



GLOBAL TUBERCULOSIS REPORT

2018



World Health
Organization



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Abbreviations

aDSM	active TB drug-safety monitoring and management	MDR/RR-TB	multidrug-resistant TB or rifampicin-resistant TB
AIDS	acquired immunodeficiency syndrome	MDR-TB	multidrug-resistant TB
AMR	antimicrobial resistance	M:F	male to female (ratio)
ART	antiretroviral therapy	MIC	minimal inhibitory concentration
BCG	bacille Calmette-Guérin	mRNA	messenger RNA
BPamZ	bedaquiline, pretomanid, moxifloxacin and pyrazinamide	NACO	National AIDS Control Organization
BRICS	Brazil, Russian Federation, India, China and South Africa	NCD	noncommunicable disease
CAD	computer-aided detection	NFC	near-field communication
CFR	case fatality ratio	NGO	nongovernmental organization
CHOICE	CHOosing Interventions that are Cost-Effective (WHO)	NHI	national health insurance
CHW	community health worker	NHIF	National Health Insurance Fund
CI	confidence interval	NHIS	National Health Insurance Scheme
CRS	creditor reporting system	NIAID	National Institute of Allergy and Infectious Diseases
CV	community volunteer	NIH	National Institutes of Health
CXR	chest X-ray	NTLD	National Tuberculosis, Leprosy and Lung Disease Programme
DALY	disability-adjusted life-year	NTP	national TB programme
DST	drug susceptibility testing	OECD	Organisation for Economic Co-operation and Development
EBA	early bactericidal activity	PanACEA	Pan-African Consortium for the Evaluation of Antituberculosis Antibiotics
EDCTP	European and Developing Countries Clinical Trial Partnership	PCR	polymerase chain reaction
EECA	Eastern Europe and Central Asia	PEPFAR	President's Emergency Plan for AIDS Relief
FIND	Foundation for Innovative New Diagnostics	PLHIV	people living with HIV
GDG	Guideline Development Group	PMDT	programmatic management of drug-resistant TB
GDP	gross domestic product	P:N	prevalence to notification (ratio)
GHCC	Global Health Cost Consortium	PPM	public-public and public-private mix
Global Fund	The Global Fund to Fight AIDS, Tuberculosis and Malaria	ReSeqTB	Relational Sequencing TB Knowledgebase
GPW	General Programme of Work (WHO)	RNA	ribonucleic acid
GRADE	Grading of Recommendations Assessment, Development and Evaluation	RNTCP	Revised National TB Control Programme
HBC	high-burden country	RR	rifampicin-resistant
HDC	Health Data Collaborative	RR-TB	rifampicin-resistant TB
HIV	human immunodeficiency virus	RT-qPCR	reverse transcriptase quantitative PCR
HLM	high-level meeting	SDG	Sustainable Development Goal
ICD-10	International classification of diseases (10th edition)	SHA	System of Health Accounts
IER	Department of Information, Evidence and Research (WHO)	SRL	supranational reference laboratory
IGRA	interferon gamma release assay	SSI	Statens Serum Institut
IHME	Institute of Health Metrics and Evaluation	TB	tuberculosis
ILO	International Labour Organization	TB Alliance	Global Alliance for TB Drug Development
LAM	lipoarabinomannan	TBTC	TB Trial Consortium
LEAP	Livelihood Empowerment Against Poverty	TNF	tumour necrosis factor
LF-LAM	lateral flow lipoarabinomannan assay	UCSR	Unit Cost Study Repository
LPA	line probe assay	UHC	universal health coverage
LTBI	latent TB infection	UN	United Nations
MAMS-TB	multi-arm, multi-stage TB	UNAIDS	Joint United Nations Programme on HIV/AIDS
MBLA	molecular bacterial load assay	US	United States
MDG	Millennium Development Goal	USA	United States of America
MDR	multidrug-resistant	USAID	US Agency for International Development
		VR	vital registration
		WHO	World Health Organization
		WRD	WHO-recommended rapid diagnostic
		XDR-TB	extensively drug-resistant TB

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The United Nations flag outside the Secretariat building of the United Nations, New York City, United States of America
Mike Segar / Reuters

Executive Summary

Context

On 26 September 2018, the United Nations (UN) will hold its first high-level meeting on tuberculosis (TB), at its headquarters in New York. The title of the meeting – United to End TB: An Urgent Global Response to a Global Epidemic – highlights the need for immediate action to accelerate progress towards the goal of ending the TB epidemic by 2030.

All Member States of WHO and the UN have committed to this goal, initially through their unanimous endorsement of WHO's End TB Strategy at the World Health Assembly in May 2014 and then their adoption of the UN Sustainable Development Goals (SDGs) in September 2015. Specific targets for 2030 set in the End TB Strategy are a 90% reduction in the absolute number of TB deaths and an 80% reduction in TB incidence (new cases per 100 000 population per year), compared with levels in 2015.¹

The UN high-level meeting follows the first WHO global ministerial conference on ending TB in the SDG era, which was held in November 2017 in the Russian Federation. The conference brought together over 1000 participants, including ministers of health and other leaders from 120 countries, and over 800 partners, including civil society. That conference resulted in the Moscow Declaration to End TB. At the World Health Assembly in May 2018, all WHO Member States committed to accelerate their actions to end TB, building on the Moscow Declaration.

In the months leading up to the UN high-level meeting, major country blocs have issued communiqués on the need for action on TB, including drug-resistant TB in the wider context of antimicrobial resistance (AMR). Examples include the G20, the G7, the BRICS group (Brazil, the Russian Federation, India, China and South Africa) and the Asia-Pacific Economic Cooperation (APEC). New commitments were made by ministers from countries in the WHO South-East Asia Region at the Delhi End TB Summit in March 2018 and by African leaders at a meeting of the African Union in July 2018.

This report

WHO has published a global TB report every year since 1997. This 2018 edition is published in the lead up to the UN high-level meeting on TB. It provides a comprehen-

sive and up-to-date assessment of the TB epidemic, and of progress in the response to the epidemic, at global, regional and country levels. The report is based primarily on data reported annually to WHO by countries, and databases maintained by other UN agencies and the World Bank.

Latest status of the TB epidemic

Worldwide, TB is one of the top 10 causes of death and the leading cause from a single infectious agent (above HIV/AIDS). Millions of people continue to fall sick with TB each year.

In 2017, TB caused an estimated 1.3 million deaths (range, 1.2–1.4 million)² among HIV-negative people and there were an additional 300 000 deaths from TB (range, 266 000–335 000) among HIV-positive people.³

Globally, the best estimate is that 10.0 million people (range, 9.0–11.1 million) developed TB disease in 2017: 5.8 million men, 3.2 million women and 1.0 million children. There were cases in all countries and age groups, but overall 90% were adults (aged ≥15 years), 9% were people living with HIV (72% in Africa) and two thirds were in eight countries: India (27%), China (9%), Indonesia (8%), the Philippines (6%), Pakistan (5%), Nigeria (4%), Bangladesh (4%) and South Africa (3%). These and 22 other countries in WHO's list of 30 high TB burden countries accounted for 87% of the world's cases.⁴ Only 6% of global cases were in the WHO European Region (3%) and WHO Region of the Americas (3%).

The severity of national epidemics varies widely among countries. In 2017, there were fewer than 10 new cases per 100 000 population in most high-income countries, 150–400 in most of the 30 high TB burden countries, and above 500 in a few countries including Mozambique, the Philippines and South Africa.

Drug-resistant TB continues to be a public health crisis. The best estimate is that, worldwide in 2017, 558 000 people (range, 483 000–639 000) developed TB that was resistant to rifampicin (RR-TB), the most effective first-line drug, and of these, 82% had multidrug-resistant TB (MDR-TB).⁵ Three countries accounted for almost half of the world's cases of MDR/RR-TB: India (24%), China (13%) and the Russian Federation (10%).

Globally, 3.6% of new TB cases and 17% of previously

treated cases had MDR/RR-TB. The highest proportions (>50% in previously treated cases) are in countries of the former Soviet Union. Among cases of MDR-TB in 2017, 8.5% (95% confidence interval, 6.2–11%) were estimated to have extensively drug-resistant TB (XDR-TB).⁶

About 1.7 billion people, 23% of the world's population, are estimated to have a latent TB infection, and are thus at risk of developing active TB disease during their lifetime.

Progress in reducing TB cases and deaths

The disease burden caused by TB is falling globally, in all WHO regions, and in most countries, but not fast enough to reach the first (2020) milestones of the End TB Strategy.

By 2020, the TB incidence rate (new cases per 100 000 population per year) needs to be falling at 4–5% per year, and the proportion of people with TB who die from the disease (the case fatality ratio, CFR) needs to fall to 10%.

In 2017, the proportion of people with TB who died from the disease was 16%, down from 23% in 2000.

Worldwide, the TB incidence rate is falling at about 2% per year.⁷ The fastest regional declines from 2013 to 2017 were in the WHO European Region (5% per year) and the WHO African Region (4% per year). In the same 5 years, particularly impressive reductions (4–8% per year) occurred in southern Africa (e.g. Eswatini, Lesotho, Namibia, South Africa, Zambia and Zimbabwe), following a peak in the HIV epidemic and the expansion of TB and HIV prevention and care; and in the Russian Federation (5% per year), following intensified efforts to reduce the burden of TB and scrutiny of progress from the highest political levels.

Globally, the absolute number of TB deaths among HIV-negative people has fallen by a best estimate of 29% since 2000, from 1.8 million in 2000 to 1.3 million in 2017, and by 5% since 2015 (the baseline year of the End TB Strategy). The number of TB deaths among HIV-positive people has fallen by 44% since 2000, from 534 000 in 2000 to 300 000 in 2017, and by 20% since 2015.

The TB mortality rate (i.e. TB deaths among HIV-negative people per 100 000 population per year) is falling at about 3% per year, and the overall reduction in the period 2000–2017 was 42%. Of the WHO regions, the fastest declines in the 5 years 2013–2017 were in the WHO European Region (11% per year) and the WHO South-East Asia Region (4% per year). High TB burden countries with rates of decline exceeding 6% per year in the 5 years 2013–2017 include the Russian Federation (13% per year), Ethiopia (12% per year), Sierra Leone (10% per year), Kenya (8% per year) and Viet Nam (8% per year).

TB diagnosis and treatment

Diagnosis and successful treatment of people with TB averts millions of deaths each year (an estimated 54 million over the period 2000–2017), but there are still large and persistent gaps in detection and treatment.

Worldwide in 2017, 6.4 million new cases of TB were officially notified to national authorities and then reported to WHO. This number has been increasing since 2013, following 4 years (2009–2012) in which 5.7–5.8 million new cases were reported annually, mainly due to increased reporting of detected cases by the private sector in India and, in 2017, an upturn in notifications in Indonesia.

The 6.4 million cases reported represented 64% of the estimated 10.0 million new cases that occurred in 2017. Ten countries accounted for 80% of the 3.6 million global gap, the top three being India (26%), Indonesia (11%) and Nigeria (9%).⁸

Gaps between the estimated number of new cases and the number actually reported are due to a mixture of underreporting of detected cases, and underdiagnosis (either because people do not access health care, or because they are not diagnosed when they do). Underestimation or overestimation of the total number of new cases is also possible. An informative example is Indonesia; in 2017, a national study found that although about 80% of new cases were detected, 41% of these cases were not reported. Actions to correct underreporting are being put in place.

There were 464 633 reported cases of TB among people living with HIV in 2017 (51% of the estimated 920 000 new cases in the same year), of whom 84% were on antiretroviral therapy. Most of the gaps in detection and treatment were in the WHO African Region, where the burden of HIV-associated TB is highest.

To support countries to close gaps in TB detection and treatment, in 2018 WHO, in collaboration with the Stop TB Partnership and the Global Fund to Fight AIDS, Tuberculosis and Malaria, launched an initiative called Find. Treat. All.⁹ The initiative includes a target of detecting and treating 40 million people with TB in the period 2018–2022.

The latest treatment outcome data for new cases show a global treatment success rate of 82% in 2016. This is a reduction from 86% in 2013 and 83% in 2015; in countries where notifications have increased, reporting of treatment outcomes has not kept pace.

Drug-resistant TB: diagnosis and treatment

Urgent action is required to improve the coverage and quality of diagnosis, treatment and care for people with drug-resistant TB.

Globally, 160 684 cases of MDR/RR-TB were detected and notified in 2017 (a small increase from 153 119

in 2016). Of these, a total of 139 114 people (87%) were enrolled on treatment with a second-line regimen, up from 129 689 in 2016 but still only 25% of the estimated 558 000 people who developed MDR/RR-TB in 2017. China and India alone accounted for 40% of the global gap; these and eight other countries¹⁰ accounted for 75%.

Treatment success remains low, at 55% globally. Examples of high burden countries in which better treatment success rates are being achieved include Bangladesh, Ethiopia, Kazakhstan, Myanmar and Viet Nam (all of which have rates above 70%).¹¹

Closing gaps in detection and treatment requires much higher coverage of drug susceptibility testing among people diagnosed with TB, reducing underdiagnosis of TB, models of care that make it easier to access and continue treatment, new diagnostics, and new medicines and treatment regimens with higher efficacy and better safety.

In July 2018, the latest evidence on treatment of drug-resistant TB was reviewed by an independent panel of experts convened by WHO. A rapid communication on key changes to recommendations for the treatment of drug-resistant TB has been issued by WHO, to be followed by the release of updated and consolidated WHO policy guidelines later in the year.

TB prevention services

The main health-care interventions to prevent new infections of *Mycobacterium tuberculosis* and their progression to TB disease are treatment of latent TB infection and vaccination of children with the bacille Calmette-Guérin (BCG) vaccine. TB preventive treatment for a latent TB infection is expanding, but most of those for whom it is strongly recommended are not yet accessing care, whereas coverage of BCG vaccination is high.

WHO has strongly recommended treatment for latent TB infection in two priority groups: people living with HIV, and children aged under 5 years who are household contacts of someone who has bacteriologically confirmed pulmonary TB.

The number of people living with HIV reported to have been started on preventive treatment was 958 559 in 2017. Of the 15 high TB/HIV burden countries that reported data, coverage ranged from 1% in Eswatini to 53% in South Africa. The number for children aged under 5 years reached 292 182 in 2017 – a threefold increase from 2015 but still only around 23% of the 1.3 million estimated to be eligible.

In countries with a high incidence of TB, WHO guidance issued in 2018 includes a new recommendation to consider testing and treatment for people aged 5 years or more who are household contacts of bacteriologically confirmed pulmonary TB cases. This substantially increases the potential number of people eligible for treatment.

WHO estimates that at least 30 million people will be eligible for TB preventive treatment between 2018 and 2022.

BCG vaccination should be provided as part of national childhood immunization programmes according to a country's TB epidemiology. In 2017, 158 countries reported providing BCG vaccination, of which 120 reported coverage of at least 90%.

Financing for TB prevention, diagnosis and treatment

Funding for the provision of TB prevention, diagnostic and treatment services has more than doubled since 2006 but continues to fall short of what is needed.

In 119 low- and middle-income countries that reported data (and accounted for 97% of reported TB cases globally), funding reached US\$ 6.9 billion in 2018. The amount available each year has been in the range US\$ 6–7 billion since 2014, after increasing from US\$ 3.3 billion in 2006. The Stop TB Partnership's Global Plan to End TB 2016–2020 estimated that US\$ 10.4 billion is required in these countries in 2018, leaving a gap of US\$ 3.5 billion. Without an increase in funding, the annual gap will widen to US\$ 5.4 billion in 2020 and to at least US\$ 6.1 billion in 2022.¹²

As in previous years, most of the funding (86%) available in 2018 is from domestic sources. However, this global aggregate figure is strongly influenced by BRICS, in which 96% (range 91–100%) of funding is from domestic sources. In India, domestic funding more than tripled between 2016 and 2018.

International donor funding (US\$ 0.9 billion in 2018, a slight decrease from 2017) accounts for 39% of funding in the 25 high TB burden countries outside BRICS and for 57% of funding in low-income countries.

Universal health coverage, social protection and social determinants

The End TB Strategy milestones for 2020 and 2025 can only be achieved if TB diagnosis, treatment and prevention services are provided within the context of progress towards universal health coverage (UHC), and if there is multisectoral action to address the social and economic factors that drive TB epidemics.

TB incidence needs to be falling at 10% per year by 2025, and the proportion of people with TB who die from the disease needs to fall to 6.5% by 2025. Such levels have only been achieved in the context of UHC, combined with social and economic development that reduces known risk factors for TB infection and disease.

UHC means that everyone – irrespective of their living standards – receives the health services they need, and that using health services does not cause financial hardship. SDG Target 3.8 is to achieve UHC by 2030.

A 2017 WHO/World Bank report on UHC found that at least half of the world's population lacks access to essential health services and almost 10% experience catastrophic expenditures on health. All of the 30 high TB burden countries need to increase service coverage and reduce levels of catastrophic expenditures to reach UHC, consistent with findings from surveys of costs faced by TB patients and their households.

WHO projections published in 2017 suggest that most middle-income countries could mobilize the funding needed to achieve UHC by 2030 from domestic resources, while this is unlikely in low-income countries.

This report features a TB-SDG monitoring framework that focuses attention on 14 indicators (from seven SDGs) that are associated with TB incidence. Monitoring of these indicators can be used to identify key influences on the TB epidemic at national level and inform the multisectoral actions required to end it.

Many new cases of TB are attributable to undernourishment, HIV infection, smoking, diabetes and alcohol use (five of the indicators featured in the TB-SDG framework). A recent modelling study shows that eliminating extreme poverty and providing social protection (both targets under SDG 1, and two other indicators in the TB-SDG framework) could substantially reduce TB incidence.

TB research and development

The SDG and End TB Strategy targets set for 2030 cannot be met without intensified research and development.

Technological breakthroughs are needed by 2025, so that the annual decline in the global TB incidence rate can be accelerated to an average of 17% per year. Priorities include a vaccine to lower the risk of infection, a vaccine or new drug treatment to cut the risk of TB disease in the 1.7 billion people already latently infected, rapid diagnostics for use at the point of care and simpler, shorter drug regimens for treating TB disease.

The development pipelines are progressing, but slowly. Few diagnostic technologies emerged in 2017. There are 20 drugs, several treatment regimens and 12 vaccine candidates in clinical trials.

Annual reports by Treatment Action Group published since 2006 show that funding for TB research and development has increased in recent years, peaking at US\$ 724 million in 2016. However, this is only 36% of the estimated requirement of US\$ 2 billion per year.

Actions needed to accelerate progress

Accelerating progress towards ending TB requires closing gaps in TB diagnosis, treatment and prevention within the context of progress towards UHC, multisectoral efforts to address the social and economic determinants and consequences of TB, intensified TB research and development, and strengthened accountability using a framework to track and review progress towards

commitments and actions needed to end TB at global, regional and national levels. These are only possible with increased and sustained funding, including from domestic sources (especially in middle-income countries), international donors and public-private partnerships.

For countries where the burden of TB is already low, the focus should be on actions needed to eliminate TB, paying particular attention to vulnerable groups with the highest risk of infection and disease.

Conclusion

TB is an old disease that was once a death sentence. Effective drug treatments first became available in the 1940s, and in combination with social and economic development they allowed countries in western Europe, North America and some other parts of the world to reduce their burden of TB disease to very low levels.¹³ For most countries, however, the "end" of TB as an epidemic and major public health problem remains an aspiration rather than a reality. The UN high-level meeting on TB on 26 September 2018, with attendance of heads of state and other eminent people, provides a platform to step up the commitments and actions needed to end the global TB epidemic, by the SDG deadline of 2030.

¹ The first milestones, for 2020, are a 35% reduction in TB deaths and a 20% reduction in TB incidence, compared with 2015. The SDG target of ending the TB epidemic is part of SDG Target 3.3, under the SDG health goal (SDG 3).

² Here and throughout the report, "range" refers to the 95% uncertainty interval.

³ When an HIV-positive person dies from TB disease, the underlying cause is coded as HIV in the International classification of diseases system.

⁴ The other 22 countries are Angola, Brazil, Cambodia, Central African Republic, Congo, the Democratic People's Republic of Korea, the Democratic Republic of the Congo, Ethiopia, Kenya, Lesotho, Liberia, Mozambique, Myanmar, Namibia, Papua New Guinea, the Russian Federation, Sierra Leone, Thailand, the United Republic of Tanzania, Viet Nam, Zambia and Zimbabwe.

⁵ Defined as resistance to rifampicin and isoniazid.

⁶ Defined as MDR-TB plus resistance to at least one drug in the following two classes of medicines used in treatment of MDR-TB: fluoroquinolones and second-line injectable agents.

⁷ The absolute number has been around 10 million per year since 2000, and has fallen slowly since 2005.

⁸ The other seven countries are shown in Fig. 4.17.

⁹ <http://www.who.int/tb/joint-initiative/en/>

¹⁰ The other eight countries are shown in Fig. 4.21.

¹¹ The countries listed are those treating at least 500 MDR/RR-TB patients annually.

¹² This figure is based on a recent extension of Global Plan projections, which indicate that at least US\$ 13 billion will be required annually by 2022.

¹³ Around 10 or fewer new TB cases per 100 000 population per year and less than one TB death per 100 000 population per year.



**UNITED TO END TUBERCULOSIS:
AN URGENT GLOBAL RESPONSE TO A GLOBAL EPIDEMIC**

BOX 1.1

Basic facts about TB

TB is an infectious disease caused by the bacillus *Mycobacterium tuberculosis*. It typically affects the lungs (pulmonary TB), but can also affect other sites (extrapulmonary TB). The disease is spread when people who are sick with pulmonary TB expel bacteria into the air, for example by coughing.

A relatively small proportion (5–10%) of the estimated 1.7 billion people infected with *M. tuberculosis* will develop TB disease during their lifetime. However, the probability of developing TB disease is much higher among people infected with HIV; it is also higher among people affected by risk factors such as undernutrition, diabetes, smoking and alcohol consumption.

Overall, about 90% of cases occur among adults, with more cases among men than women. The male:female ratio among adults is approximately 2:1.

Diagnostic tests for TB disease include:

- **Rapid molecular tests** – The only rapid test for diagnosis of TB currently recommended by WHO is the Xpert® MTB/RIF assay (Cepheid, USA). It can provide results within 2 hours, and was initially recommended (in 2010) for diagnosis of pulmonary TB in adults. Since 2013, it has also been recommended for use in children and to diagnose specific forms of extrapulmonary TB. The test has much better accuracy than sputum smear microscopy.
- **Sputum smear microscopy** – Developed more than 100 years ago, this technique requires the examination of sputum samples using a microscope to determine the presence of bacteria.
- **Culture-based methods** – These form the current reference standard; they require more developed laboratory capacity and can take up to 12 weeks to provide results.

Globally, use of rapid molecular tests is increasing, and many countries are phasing out the use of smear microscopy for diagnostic purposes (although microscopy and culture remain necessary for treatment monitoring).

There are also tests for TB that is resistant to first-line and second-line anti-TB drugs. They include Xpert MTB/RIF, which simultaneously tests for TB and resistance to rifampicin (the most effective first-line anti-TB drug); rapid line probe assays (LPAs) that test for resistance to rifampicin and isoniazid (referred to as first-line LPAs); a rapid

LPA that tests for resistance to fluoroquinolones and injectable anti-TB drugs (referred to as a second-line LPA); and sequencing technologies. First-line LPAs were first recommended by WHO in 2008; the second-line LPA was first recommended in May 2016. Culture-based methods currently remain the reference standard for drug susceptibility testing.

Without treatment, the mortality rate from TB is high. Studies of the natural history of TB disease in the absence of treatment with anti-TB drugs (conducted before drug treatments became available) found that about 70% of individuals with sputum smear-positive pulmonary TB died within 10 years of being diagnosed, as did about 20% of people with culture-positive (but smear-negative) pulmonary TB.^a

Effective drug treatments were first developed in the 1940s. The currently recommended treatment for cases of drug-susceptible TB is a 6-month regimen of four first-line drugs: isoniazid, rifampicin, ethambutol and pyrazinamide. The Global TB Drug Facility supplies a complete 6-month course for about US\$ 40 per person. Treatment success rates of at least 85% for cases of drug-susceptible TB are regularly reported to WHO by its 194 Member States. Treatment for rifampicin-resistant TB (RR-TB) and multidrug-resistant TB (MDR-TB)^b is longer, and requires more expensive (≥US\$ 1000 per person) and more toxic drugs. The latest data reported to WHO show a treatment success rate for MDR-TB of 55%, globally.

There are 20 TB drugs in clinical trials, and combination regimens that include new compounds as well as other drugs are also being tested in clinical trials. The bacille Calmette-Guérin (BCG) vaccine, which was developed almost 100 years ago and has been shown to prevent severe forms of TB in children, is still widely used. However, there is currently no vaccine that is effective in preventing TB disease in adults, either before or after exposure to TB infection. There are 12 TB vaccines in Phase I, Phase II or Phase III trials.

^a Tiemersma EW, van der Werf MJ, Borgdorff MW, Williams BG, Nagelkerke NJ. Natural history of tuberculosis: duration and fatality of untreated pulmonary tuberculosis in HIV negative patients: a systematic review. *PLoS One*. 2011;6(4):e17601 (<http://www.ncbi.nlm.nih.gov/pubmed/21483732>, accessed 3 July 2018).

^b Defined as resistance to isoniazid and rifampicin, the two most powerful anti-TB drugs.

Chapter 1. Introduction

Tuberculosis (TB) is an old disease – studies of human skeletons show that it has affected humans for thousands of years.¹ The cause remained unknown until 24 March 1882, when Dr Robert Koch announced that he had discovered the bacillus *Mycobacterium tuberculosis*, an event that is now commemorated every year as World TB Day.² The disease is spread when people who are sick with TB expel bacteria into the air, for example by coughing. Basic facts about TB are provided in **Box 1.1**.

In the late 1800s, cause-of-death data from national vital registration systems show that TB was one of the leading causes of death in some European countries. With social and economic development – such as improvements in incomes, housing and nutrition – numbers of TB cases and deaths started to decline in western Europe, North America and some other parts of the world around the turn of the 20th century, albeit slowly (1–2% per year).^{3,4} From the 1940s, the discovery, development and use of effective drug treatments substantially accelerated these trends, with national case rates (per 100 000 population) falling by up to 10% per year and mortality rates falling even faster. In countries that have experienced such reductions in disease burden, and now have only around 10 or fewer cases and less than 1 death per 100 000 population per year, TB is often regarded as a disease of the past.

For many countries, however, the “end” of TB as an epidemic and major public health problem is still a distant reality. This is despite the fact that, with a timely diagnosis and correct drug treatment, most people who develop the disease can be cured. Twenty-five years ago, in 1993, WHO declared TB a global health emergency.⁵

There has been major progress in subsequent years – more than 60 million people have been documented as treated and cured since 2000, and case and death rates have fallen steadily. Nevertheless, worldwide, around 10 million people still fall ill with the disease each year (more adults than children, and more men than women), and TB is one of the top 10 causes of death. It is also the leading cause of death from a single infectious agent, ranking above HIV/AIDS.

In 2014 and 2015, all Member States of WHO and the United Nations (UN) committed to ending the TB epidemic. They did this by unanimously endorsing WHO’s End TB Strategy at the World Health Assembly in May 2014, and by adopting the UN Sustainable Development Goals (SDGs) in September 2015. The End TB Strategy has the overall goal of ending the global TB epidemic, and it defines the targets (2030, 2035) and milestones (2020, 2025) for reductions in TB cases and deaths needed to achieve that goal. The SDGs include a target to end the TB epidemic by 2030.

In 2017 and 2018, efforts to step up political commitment to the fight against TB have intensified. The first global ministerial conference on TB was held in November 2017. The UN’s first high-level meeting (HLM) on TB, on 26 September 2018 at its headquarters in New York, includes heads of state. The title is United to End TB: An Urgent Global Response to a Global Epidemic.

WHO has published a global TB report every year since 1997. This 2018 edition is published in association with the UN HLM. It provides a comprehensive and up-to-date assessment of the TB epidemic, and of progress in the response, at global, regional and country levels. This is based primarily on data gathered by WHO from countries and territories in annual rounds of data collection, and databases maintained by other multilateral agencies.

The topics covered in the main chapters of the report are: global commitments to end TB and multisectoral accountability; estimates of TB disease burden 2000–2017; TB diagnosis and treatment; TB prevention services; financing for TB prevention, diagnosis and treatment; universal health coverage, social protection and social determinants of TB; and TB research and development.

The report’s annexes describe WHO’s online global TB database, present profiles for 30 high TB burden countries and WHO’s six regions, and contain data for key indicators for all countries, for the latest available year.

¹ Hershkovitz I, Donoghue HD, Minnikin DE, May H, Lee OY, Feldman M, et al. Tuberculosis origin: the Neolithic scenario. *Tuberculosis* (Edinb). 2015;95 Suppl 1:S122–6 (<https://www.ncbi.nlm.nih.gov/pubmed/25726364>, accessed 3 July 2018).

² Sakula A. Robert Koch: centenary of the discovery of the tubercle bacillus, 1882. *Thorax*. 1982;37(4):246–51 (<https://www.ncbi.nlm.nih.gov/pubmed/6180494>, accessed 3 July 2018).

³ Styblo K, Meijer J, Sutherland I. The transmission of tubercle bacilli: its trend in a human population. *Bull World Health Organ*. 1969;41:137–78 (<https://www.ncbi.nlm.nih.gov/pubmed/5309081>, accessed 3 July 2018).

⁴ Grange JM, Gandy M, Farmer P, Zumla A. Historical declines in tuberculosis: nature, nurture and the biosocial model. *Int J Tuberc Lung Dis*. 2001;5(3):208–12 (<https://www.ncbi.nlm.nih.gov/pubmed/11326817>, accessed 3 July 2018).

⁵ World Health Organization. TB: a global emergency, WHO report on the TB epidemic (WHO/TB/94.177). Geneva: WHO; 1994 (<http://apps.who.int/iris/handle/10665/58749>, accessed 21 June 2018).



Primary school children in a village in northern Lao People's Democratic Republic
Hadynyah / Getty Images

Chapter 2. Global commitments to end TB and multisectoral accountability

From 2000 to 2015, global and national efforts to reduce the burden of tuberculosis (TB) disease were focused on achieving targets set within the context of the Millennium Development Goals (MDGs). The MDGs were established by the United Nations (UN) in 2000, and targets were set for 2015. Target 6c of MDG 6 was to “halt and reverse” TB incidence. The Stop TB Partnership, established in 2001, adopted this target and set two additional targets: to halve TB prevalence and TB mortality rates by 2015 compared with their levels in 1990. The global TB strategy developed by WHO for the decade 2006–2015, the Stop TB Strategy, had the overall goal of reaching all three of these targets. In October 2015, WHO published its assessment of whether the 2015 global TB targets for reductions in TB incidence, prevalence and mortality had been achieved.¹

In 2016, the MDGs were succeeded by a new set of goals, known as the Sustainable Development Goals (SDGs). Adopted by the UN in September 2015 following 3 years of consultations, the SDG framework of goals, targets and indicators covers the period 2016–2030.² Similarly, in 2012 WHO initiated work on a new global TB strategy, which was completed in 2014. The End TB Strategy was unanimously endorsed by all WHO Member States at the 2014 World Health Assembly, and covers the period 2016–2035.³ The SDGs and the End TB Strategy provide the framework for national and international efforts to end the TB epidemic during the period 2016–2030.

This chapter provides an overview of both the SDGs (Section 2.1) and the End TB Strategy (Section 2.2). It then describes the Moscow Declaration from the first global ministerial conference on ending TB (Section 2.3),⁴ which was held in November 2017 with the aim of accelerating progress towards targets set in the SDGs

and End TB Strategy through a multisectoral response. The Moscow Declaration includes commitments by WHO Member States and calls to partner agencies, and has informed the first UN high-level meeting on TB at UN headquarters in New York in September 2018. In Section 2.3, specific attention is given to the development of a multisectoral accountability framework to accelerate progress towards ending TB, which was one of four topics featured in the declaration and which has been a major focus of work for WHO, in collaboration with WHO Member States and partner agencies, in 2018.

Given the multisectoral influences on the TB epidemic and the multisectoral actions needed to end it, WHO developed a TB-SDG monitoring framework in 2017.⁵ This is described and explained in Section 2.4. The framework is designed to focus attention on, and encourage analysis of, SDG targets and indicators that will influence the course of the TB epidemic, with findings then used to drive action. Analysis of the 14 indicators included in the framework is part of Chapter 7.⁶

At the 2018 World Health Assembly, Member States endorsed WHO’s General Programme of Work (GPW) for 2019–2023.⁷ The GPW is based on the foundation of SDG 3, the health goal of the SDGs, and it includes TB targets for 2023 that are consistent with those of the End TB Strategy. Section 2.5 describes the GPW’s three strategic goals and associated outcomes, and its targets for TB, highlighting how these goals, outcomes and targets link with the SDGs and the End TB Strategy.

For the first 5 years of the SDGs and End TB Strategy (2016–2020), WHO has defined three lists of high-burden countries (HBCs): for TB, TB/HIV and multidrug-resistant TB (MDR-TB). Particular attention is given to the countries in each of these lists throughout this report. For this reason, they are presented and explained in Section 2.6.

¹ World Health Organization. Global tuberculosis report 2015. Geneva: WHO; 2015 (http://apps.who.int/iris/bitstream/10665/191102/1/9789241565059_eng.pdf, accessed 21 June 2018).

² United Nations. Sustainable Development Goals (<https://sustainabledevelopment.un.org/topics/sustainabledevelopmentgoals>, accessed 21 June 2018). A short summary of the main findings is also available in Chapter 2 of the 2016 edition of the report.

³ Uplekar M, Weil D, Lönnroth K, Jaramillo E, Lienhardt C, Dias HM, et al. WHO’s new End TB Strategy. *Lancet*. 2015;385(9979):1799–1801 (<http://www.ncbi.nlm.nih.gov/pubmed/25814376>, accessed 21 June 2018).

⁴ The conference was titled “Ending TB in the Sustainable Development Era: a multisectoral response”.

⁵ World Health Organization. Global tuberculosis report 2017 (WHO/HTM/TB/2017.23). Geneva: WHO; 2017 (<http://apps.who.int/iris/bitstream/handle/10665/259366/9789241565516-eng.pdf>, accessed 21 June 2018).

⁶ In addition, Annex 2 shows the latest data and recent trends for each indicator for the 30 high TB burden countries. For other countries, the same data are available in country profiles that can be accessed online at www.who.int/tb/data.

⁷ See: http://apps.who.int/gb/ebwha/pdf_files/WHA71/A71_4-en.pdf?ua=1

2.1 The Sustainable Development Goals

The 17 SDGs are shown in **Box 2.1**.

The consolidated goal for health is SDG 3, which is defined as “Ensure healthy lives and promote well-being for all at all ages”. Thirteen targets have been set for this goal (**Box 2.2**), and one of these targets, Target 3.3, explicitly mentions TB: “By 2030, end the epidemics of AIDS, tuberculosis, malaria and neglected tropical diseases and combat hepatitis, water-borne diseases and other communicable diseases”. The language of “ending epidemics” is also now a prominent element of global health strategies developed by WHO and the Joint United Nations Programme on HIV/AIDS (UNAIDS) for the post-2015 era,¹ including the End TB Strategy (**Section 2.2**). The TB indicator for Target 3.3 is the TB incidence rate (new TB cases per 100 000 population per year).

SDG 3 also includes a target (Target 3.8) related to universal health coverage (UHC) in which TB is explicitly mentioned. The WHO/World Bank definition of UHC is that all people receive the health services they need, while at the same time ensuring that the use of these services does not expose the user to financial hardship.^{2,3} Target 3.8 includes an indicator for the coverage of essential prevention, treatment and care interventions. This is a composite indicator based on the coverage of 16 so-called “tracer interventions”,⁴ one of which is TB treatment.

The SDGs include considerable emphasis on disaggregated analysis and reporting of data (as well as reporting for an entire country). Depending on the indicator, examples include disaggregation by age, sex, location and economic status (e.g. bottom 40%, or bottom versus top income quintiles). Some indicators also give particular attention to specific subpopulations, such as pregnant women, people with disabilities, victims of work injuries, and migrants.

In support of the requirement for disaggregation for many indicators, SDG 17 includes two targets and associated indicators under the subheading of “Data, monitoring and accountability” that specifically refer to disaggregated data and the mechanisms needed to generate such data (**Table 2.1**). Emphasis is also given to the importance of death registration within national vital registration systems, to allow for accurate tracking

of causes of death (this is Part b of Indicator 17.19.2). Strengthening national vital registration systems as the basis for direct measurement of the number of TB deaths is one of the five strategic areas of work of the WHO Global Task Force on TB Impact Measurement, as discussed in **Chapter 3**.

Disaggregation is intended to inform analysis of within-country inequalities and associated assessments of equity, with findings used to identify particular areas or subpopulations where progress is lagging and greater attention is needed. Such disaggregation is also an important consideration for the TB community, given the influence of sex, age, socioeconomic status and differential access to health care on the risks for and consequences of TB infection and disease. **Chapter 3** and **Chapter 4** of this report include analyses of TB data disaggregated by age and sex.

2.2 The End TB Strategy

The End TB Strategy “at a glance” is shown in **Box 2.3**.

The overall goal is to “End the global TB epidemic”, and there are three high-level, overarching indicators and related targets (for 2030 – linked to the SDGs – and for 2035) and milestones (for 2020 and 2025). The three indicators are:

- the number of TB deaths per year;
- the TB incidence rate (new cases per 100 000 population per year); and
- the percentage of TB-affected households that experience catastrophic costs as a result of TB disease.

The 2030 targets are a 90% reduction in TB deaths and an 80% reduction in the TB incidence rate, compared with levels in 2015. The 2035 targets are a 95% reduction in TB deaths and a 90% reduction in the TB incidence rate, compared with levels in 2015. The most immediate milestones, set for 2020, are a 35% reduction in TB deaths and a 20% reduction in the TB incidence rate, compared with levels in 2015. The trajectories of TB incidence and TB deaths that are required to reach these milestones and targets are shown in **Fig. 2.1**. For the third indicator (the percentage of TB-affected households that experience catastrophic costs as a result of TB disease), the milestone for 2020 is zero, to be sustained thereafter.

The Stop TB Partnership has developed a *Global Plan to End TB, 2016–2020*,⁵ which focuses on the actions and funding needed to reach the 2020 milestones of the End TB Strategy. More details about this plan are provided in **Chapter 6**.

Progress towards UHC and actions to address health-related risk factors for TB (as well as broader social and economic determinants of TB) will be fundamental to achieving the targets and milestones for

¹ World Health Organization. Accelerating progress on HIV, tuberculosis, malaria, hepatitis and neglected tropical diseases: a new agenda for 2016–2030. Geneva: WHO; 2015 (<http://www.who.int/about/structure/organigram/html/progress-hiv-tb-malaria-ntd/en/>, accessed 21 June 2018).

² World Health Organization/World Bank Group. Tracking universal health coverage: first global monitoring report. Geneva: WHO; 2015 (http://apps.who.int/iris/bitstream/10665/174536/1/9789241564977_eng.pdf?ua=1, accessed 21 June 2018).

³ World Health Organization/World Bank Group. Tracking universal health coverage: 2017 global monitoring report. Geneva: WHO; 2017 (<http://apps.who.int/iris/bitstream/handle/10665/259817/9789241513555-eng.pdf>, accessed 21 June 2018).

⁴ There are many different prevention and treatment interventions. SDG indicator 3.8.1 is based on the coverage of 16 interventions that have been selected as “tracers” for assessment of progress towards UHC for all interventions. Further details are provided in **Chapter 7**.

⁵ The Global Plan to End TB, 2016–2020. Geneva: Stop TB Partnership; 2015 (<http://www.stoptb.org/global/plan/>, accessed 21 June 2018).

BOX 2.1

The Sustainable Development Goals

- Goal 1. End poverty in all its forms everywhere
- Goal 2. End hunger, achieve food security and improved nutrition and promote sustainable agriculture
- Goal 3. Ensure healthy lives and promote well-being for all at all ages
- Goal 4. Ensure inclusive and equitable quality education and promote lifelong learning opportunities for all
- Goal 5. Achieve gender equality and empower all women and girls
- Goal 6. Ensure availability and sustainable management of water and sanitation for all
- Goal 7. Ensure access to affordable, reliable, sustainable and modern energy for all
- Goal 8. Promote sustained, inclusive and sustainable economic growth, full and productive employment and decent work for all
- Goal 9. Build resilient infrastructure, promote inclusive and sustainable industrialization and foster innovation
- Goal 10. Reduce inequality within and among countries
- Goal 11. Make cities and human settlements inclusive, safe, resilient and sustainable
- Goal 12. Ensure sustainable consumption and production patterns
- Goal 13. Take urgent action to combat climate change and its impacts^a
- Goal 14. Conserve and sustainably use the oceans, seas and marine resources for sustainable development
- Goal 15. Protect, restore and promote sustainable use of terrestrial ecosystems, sustainably manage forests, combat desertification, and halt and reverse land degradation and halt biodiversity loss
- Goal 16. Promote peaceful and inclusive societies for sustainable development, provide access to justice for all and build effective, accountable and inclusive institutions at all levels
- Goal 17. Strengthen the means of implementation and revitalize the Global Partnership for Sustainable Development



^a Acknowledging that the United Nations Framework Convention on Climate Change is the primary international, intergovernmental forum for negotiating the global response to climate change.

BOX 2.2

Sustainable Development Goal 3 and its 13 targets

SDG 3: Ensure healthy lives and promote well-being for all at all ages

Targets

- 3.1 By 2030, reduce the global maternal mortality ratio to less than 70 per 100 000 live births
- 3.2 By 2030, end preventable deaths of newborns and children under 5 years of age, with all countries aiming to reduce neonatal mortality to at least as low as 12 per 1000 live births and under-5 mortality to at least as low as 25 per 1000 live births
- 3.3 By 2030, end the epidemics of AIDS, tuberculosis, malaria and neglected tropical diseases and combat hepatitis, water-borne diseases and other communicable diseases
- 3.4 By 2030, reduce by one third premature mortality from non-communicable diseases through prevention and treatment and promote mental health and well-being
- 3.5 Strengthen the prevention and treatment of substance abuse, including narcotic drug abuse and harmful use of alcohol
- 3.6 By 2020, halve the number of global deaths and injuries from road traffic accidents
- 3.7 By 2030, ensure universal access to sexual and reproductive health-care services, including for family planning, information and education, and the integration of reproductive health into national strategies and programmes
- 3.8 Achieve universal health coverage, including financial risk protection, access to quality essential health-care services and access to safe, effective, quality and affordable essential medicines and vaccines for all
- 3.9 By 2030, substantially reduce the number of deaths and illnesses from hazardous chemicals and air, water and soil pollution and contamination
- 3.a Strengthen the implementation of the World Health Organization Framework Convention on Tobacco Control in all countries, as appropriate
- 3.b Support the research and development of vaccines and medicines for the communicable and non-communicable diseases that primarily affect developing countries, provide access to affordable essential medicines and vaccines, in accordance with the Doha Declaration on the TRIPS Agreement and Public Health, which affirms the right of developing countries to use to the full the provisions in the Agreement on Trade-Related Aspects of Intellectual Property Rights regarding flexibilities to protect public health, and, in particular, provide access to medicines for all
- 3.c Substantially increase health financing and the recruitment, development, training and retention of the health workforce in developing countries, especially in least developed countries and small island developing States
- 3.d Strengthen the capacity of all countries, in particular developing countries, for early warning, risk reduction and management of national and global health risks

TRIPS, Trade-Related Aspects of Intellectual Property Rights

reductions in TB cases and deaths, for two reasons. First, reaching the milestones for reductions in TB cases and deaths set for 2020 and 2025 requires the annual decline in the global TB incidence rate to accelerate from 1.5% per year in 2015 to 4–5% per year by 2020, and then to 10% per year by 2025. A decline of 10% per year is equivalent to the best-ever performance to date at national level (e.g. in countries in western Europe during the 1950s and 1960s), and has only been documented in the context of UHC combined with broader social and economic development. Second, the global proportion of people with TB who die from the disease (the case fatality ratio, or CFR) needs to be reduced to 10% by 2020 and then to 6.5% by 2025. A CFR of 6.5% is similar

to the current level in many high-income countries, but is only possible if all those with TB disease can access high-quality treatment. Analysis of CFRs at global and national levels is included in [Chapter 3](#).

The percentage of TB patients and their households facing catastrophic costs is a good tracer indicator for progress towards UHC as well as social protection. If UHC and social protection are in place, then people with TB should be able to access high-quality diagnosis and treatment without incurring catastrophic costs.¹

After 2025, reaching the 2030 and 2035 targets will require an unprecedented acceleration in the rate at

¹ This indicator, including results from recent national surveys to measure it, is discussed in more detail in [Chapter 7](#).

TABLE 2.1

SDG 17, and targets and indicators related to data, monitoring and accountability

SDG 17: Strengthen the means of implementation and revitalize the global partnership for sustainable development	
TARGETS	INDICATORS
17.18 By 2020, enhance capacity-building support to developing countries, including for least developed countries and small island developing States, to increase significantly the availability of high-quality, timely and reliable data disaggregated by income, gender, age, race, ethnicity, migratory status, disability, geographic location and other characteristics relevant in national contexts	17.18.1 Proportion of sustainable development indicators produced at the national level with full disaggregation when relevant to the target, in accordance with the Fundamental Principles of Official Statistics
17.19 By 2030, build on existing initiatives to develop measurements of progress on sustainable development that complement gross domestic product, and support statistical capacity-building in developing countries	17.19.2 Proportion of countries that (a) have conducted at least one population and housing census in the last 10 years; and (b) have achieved 100 per cent birth registration and 80 per cent death registration

BOX 2.3

The End TB Strategy at a glance

VISION	A WORLD FREE OF TB — zero deaths, disease and suffering due to TB			
	END THE GLOBAL TB EPIDEMIC			
GOAL	MILESTONES		TARGETS	
	2020	2025	SDG 2030 ^a	END TB 2035
INDICATORS				
Percentage reduction in the absolute number of TB deaths (compared with 2015 baseline)	35%	75%	90%	95%
Percentage reduction in the TB incidence rate (compared with 2015 baseline)	20%	50%	80%	90%
Percentage of TB-affected households experiencing catastrophic costs due to TB (level in 2015 unknown)	0%	0%	0%	0%

PRINCIPLES

1. Government stewardship and accountability, with monitoring and evaluation
2. Strong coalition with civil society organizations and communities
3. Protection and promotion of human rights, ethics and equity
4. Adaptation of the strategy and targets at country level, with global collaboration

PILLARS AND COMPONENTS

1. INTEGRATED, PATIENT-CENTRED CARE AND PREVENTION

- A. Early diagnosis of TB including universal drug-susceptibility testing, and systematic screening of contacts and high-risk groups
- B. Treatment of all people with TB including drug-resistant TB, and patient support
- C. Collaborative TB/HIV activities, and management of comorbidities
- D. Preventive treatment of persons at high risk, and vaccination against TB

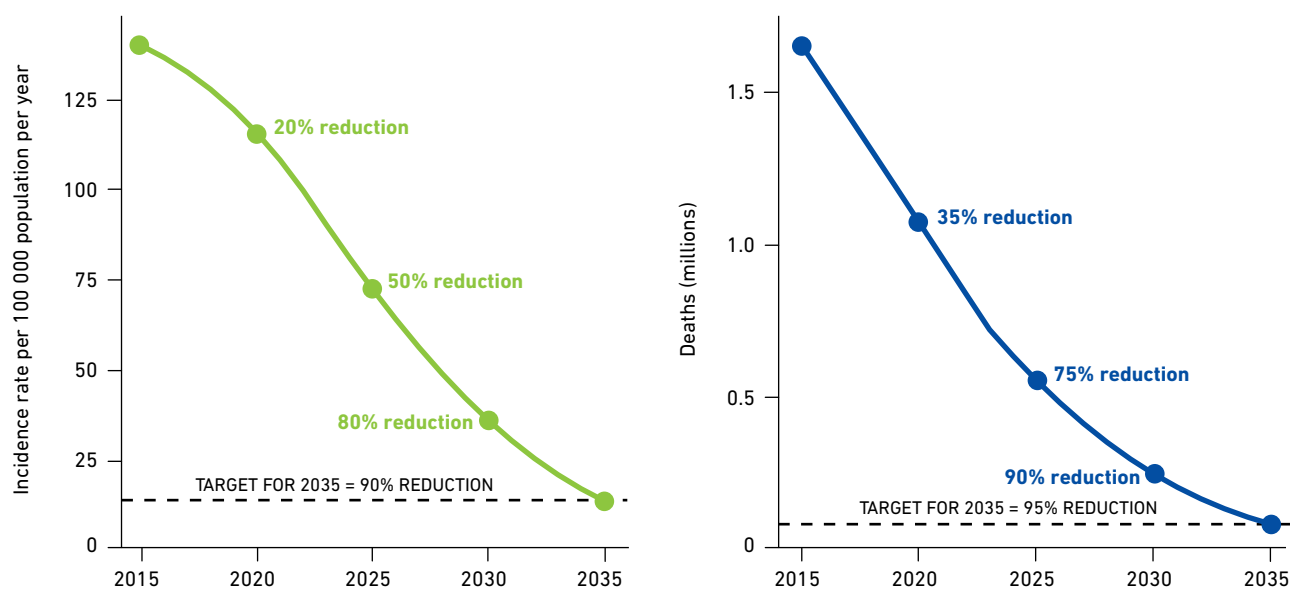
2. BOLD POLICIES AND SUPPORTIVE SYSTEMS

- A. Political commitment with adequate resources for TB care and prevention
- B. Engagement of communities, civil society organizations, and public and private care providers
- C. Universal health coverage policy, and regulatory frameworks for case notification, vital registration, quality and rational use of medicines, and infection control
- D. Social protection, poverty alleviation and actions on other determinants of TB

3. INTENSIFIED RESEARCH AND INNOVATION

- A. Discovery, development and rapid uptake of new tools, interventions and strategies
- B. Research to optimize implementation and impact, and promote innovations

^a Targets linked to the Sustainable Development Goals (SDGs).

FIG. 2.1**Projected incidence and mortality curves that are required to reach End TB Strategy targets and milestones, 2015–2035**

which TB incidence falls globally, to an average of 17% per year. Such an acceleration will depend on a technological breakthrough that can substantially reduce the risk of developing TB disease among the approximately 1.7 billion people¹ (approximately one quarter of the world's population) who are already infected with *Mycobacterium tuberculosis*. Examples include an effective post-exposure vaccine or a short, efficacious and safe treatment for latent TB infection. The latest status of the development pipelines for new TB diagnostics, drugs and vaccines is presented in [Chapter 8](#).

To achieve the targets and milestones, the End TB Strategy has four underlying principles and three pillars. The four principles are government stewardship and accountability, with monitoring and evaluation; a strong coalition with civil society organizations and communities; protection and promotion of human rights, ethics and equity; and adaptation of the strategy and targets at country level, with global collaboration. The three pillars are integrated, patient-centred TB care and prevention; bold policies and supportive systems (including UHC, social protection, and action on TB determinants); and intensified research and innovation. The 10 components of the three pillars of the End TB Strategy are shown in [Box 2.3](#).

WHO has defined 10 priority indicators for monitoring of progress in implementing the End TB Strategy. These are shown in [Table 2.2](#). The table also indicates the particular chapter of this report in which available data for each indicator can be found.

¹ Houben RM, Dodd PJ. The global burden of latent tuberculosis infection: a re-estimation using mathematical modelling. *PLoS Med*. 2016;13(10):e1002152 (<https://www.ncbi.nlm.nih.gov/pubmed/27780211>, accessed 21 June 2018).

Data for five of the 10 indicators cannot be captured routinely using the standard recording and reporting forms for paper-based systems that are included in the latest revision of WHO's framework for TB case definitions and reporting.² Collection of data on the costs faced by TB patients and their households, and assessment of whether these are catastrophic (Indicator 3 in [Table 2.2](#)) requires periodic surveys of a representative sample of TB patients; further details are provided in [Chapter 7](#). For the other four indicators (Indicators 4, 5, 6 and 8 in [Table 2.2](#)), data may already be captured routinely in countries that have electronic case-based systems for recording and reporting of data; if this is not the case, these systems can be adapted to capture the information. Alternatively, countries can undertake periodic surveys of the medical records or patient cards of a random sample of TB patients. Further guidance is provided in the WHO operational guidance on the End TB Strategy.³

2.3 The Moscow Declaration to end TB

The first global ministerial conference on ending TB was held in Moscow in November 2017. It was organized by WHO and the Ministry of Health of the Russian Federation, in recognition of the fact that investments and actions have been falling short of those needed to reach SDG and End TB Strategy targets and milestones.

² World Health Organization. Definitions and reporting framework for tuberculosis – 2013 revision (updated December 2014) (WHO/HTM/TB/2013.2). Geneva: WHO; 2013 (www.who.int/iris/bitstream/10665/79199/1/9789241505345_eng.pdf, accessed 21 June 2018).

³ World Health Organization. Implementing the End TB Strategy: the essentials. Geneva: WHO; 2016 (http://www.who.int/tb/publications/2015/The_Essentials_to_End_TB/en/, accessed 21 June 2018). See in particular Part II, Section 2.4.

TABLE 2.2

Top 10 indicators (not ranked) for monitoring implementation of the End TB Strategy at global and national levels, with recommended target levels that apply to all countries. The target level is for 2025 at the latest.

INDICATOR	RECOMMENDED TARGET LEVEL	MAIN RATIONALE FOR INCLUSION IN TOP 10	MAIN METHOD OF MEASUREMENT, AND RELEVANT CHAPTER OF THIS REPORT
1 TB treatment coverage <i>Number of new and relapse cases that were notified and treated, divided by the estimated number of incident TB cases in the same year, expressed as a percentage.</i>	≥90%	High-quality TB care is essential to prevent suffering and death from TB and to cut transmission. High coverage of appropriate treatment is a fundamental requirement for achieving the milestones and targets of the End TB Strategy.	Routinely collected notification data used in combination with estimate of TB incidence. Chapter 4
2 TB treatment success rate <i>Percentage of notified TB patients who were successfully treated. The target is for drug-susceptible and drug-resistant TB combined, although outcomes should also be reported separately.</i>	≥90%		Routinely collected data. Chapter 4
3 Percentage of TB-affected households that experience catastrophic costs due to TB^a <i>Number of people treated for TB (and their households) who incur catastrophic costs (direct and indirect combined), divided by the total number of people treated for TB.</i>	0%	One of the End TB Strategy's three high-level indicators; a key marker of financial risk protection (one of the two key elements of UHC) and social protection for TB-affected households.	National survey of notified TB patients. Chapter 7
4 Percentage of new and relapse TB patients tested using a WHO-recommended rapid diagnostic (WRD) at the time of diagnosis <i>Number of new and relapse TB patients tested using a WRD at the time of diagnosis, divided by the total number of new and relapse TB patients, expressed as a percentage.</i>	≥90%	Accurate diagnosis is a fundamental component of TB care. Rapid molecular diagnostic tests help to ensure early detection and prompt treatment.	Routinely collected data (as part of case-based surveillance), or national survey of medical records or patient cards of TB patients. Chapter 4
5 Latent TB infection (LTBI) treatment coverage <i>Number of people living with HIV newly enrolled in HIV care and the number of children aged <5 years who are household contacts of cases started on LTBI treatment, divided by the number eligible for treatment, expressed as a percentage (separately for each of the two groups).</i>	≥90%	Treatment of LTBI is the main treatment intervention available to prevent development of active TB disease in those already infected with <i>Mycobacterium tuberculosis</i> .	Routinely collected data (as part of case-based surveillance), or national survey of medical records or patient cards of people living with HIV and TB patients. Chapter 5
6 Contact investigation coverage <i>Number of contacts of people with bacteriologically confirmed TB who were evaluated for TB, divided by the number eligible, expressed as a percentage.</i>	≥90%	Contact tracing is a key component of TB prevention, especially in children.	As above for LTBI.
7 Drug-susceptibility testing (DST) coverage for TB patients <i>Number of TB patients with DST results for at least rifampicin, divided by the total number of notified (new and retreatment) cases in the same year, expressed as a percentage. DST coverage includes results from molecular (e.g. Xpert MTB/RIF) as well as conventional phenotypic DST results.</i>	100%	Testing for drug susceptibility for WHO-recommended drugs is essential to provide the right treatment for every person diagnosed with TB.	Routinely collected data (as part of case-based surveillance), or national survey of medical records or patient cards of TB patients. Chapter 4
8 Treatment coverage, new TB drugs <i>Number of TB patients treated with regimens that include new (endorsed after 2010) TB drugs, divided by the number of notified patients eligible for treatment with new TB drugs, expressed as a percentage.</i>	≥90%	An indicator that is relevant to monitoring the adoption of innovations in all countries. <i>The definition of which patients are eligible patients for treatment with new drugs may differ among countries.</i>	As above for DST.
9 Documentation of HIV status among TB patients <i>Number of new and relapse TB patients with documented HIV status, divided by the number of new and relapse TB patients notified in the same year, expressed as a percentage.</i>	100%	One of the core global indicators used to monitor collaborative TB/HIV activities. Documentation of HIV status is essential to provide the best care for HIV-positive TB patients, including antiretroviral therapy.	Routinely collected data for all TB patients. Chapter 4
10 Case fatality ratio (CFR) <i>Number of TB deaths divided by estimated number of incident cases in the same years, expressed as a percentage.</i>	≤5%	This is a key indicator for monitoring progress towards the 2020 and 2025 milestones. A CFR of 6% is required to achieve the 2025 global milestone for reductions in TB deaths and cases.	Mortality divided by incidence. In countries with a high-performance surveillance system, notifications approximate incidence. Chapter 3

^a Catastrophic costs are provisionally defined as total costs that exceed 20% of annual household income.

The conference brought together over 1000 participants, including ministers of health and other leaders from 120 countries, and over 800 partners, including civil society.

The key outcome of the conference was the Moscow Declaration to End TB,¹ which was adopted by almost 120 WHO Member States represented at the conference. The declaration was developed through consultations with partners and Member States, led by the the Russian Federation.

The Declaration includes both commitments by Member States and calls for actions by global agencies and other partners in four key areas (Fig. 2.2):²

- Advancing the TB response within the SDG agenda;
- Ensuring sufficient and sustainable financing;
- Pursuing science, research and innovation;
- Developing a multisectoral accountability framework.

At the World Health Assembly in May 2018, all Member States committed to accelerate their actions to end TB, building on the commitments of the Moscow Declaration. They also welcomed a draft version of a multisectoral accountability framework for TB, and supported its further development, adaptation and use (Box 2.4).



The topics of the Moscow Declaration are prominently featured in the UN high-level meeting on TB on 26 September 2018,³ which will seek further commitments from Heads of State. The title of the meeting was United to End TB: An Urgent Global Response to a Global Epidemic.

In the months leading up to the UN high-level meeting, ministers and heads of state of major country blocs have issued communiqués on the need for action on TB, including drug-resistant TB in the wider context of antimicrobial resistance (AMR). Examples include the G20; the G7; Brazil, the Russian Federation, India, China and South Africa (BRICS); and the Asia-Pacific Economic Cooperation (APEC). New commitments have also been made by ministers from countries in the WHO South-East Asia Region at the Delhi End TB Summit in March 2018 and by African leaders at a summit of the African Union in July 2018.

¹ Moscow Declaration to End TB; First WHO Global Ministerial Conference on Ending TB in the Sustainable Development Era: A Multisectoral Response, November 2017. WHO and the Ministry of Health of the Russian Federation. http://www.who.int/tb/features_archive/Moscow_Declaration_to_End_TB_final_ENGLISH.pdf?ua=1, accessed 21 June 2018).

² The SDG agenda and a multisectoral accountability framework for TB are discussed in this chapter. The topic of financing is covered in Chapter 6 and Chapter 7; research and development is the subject of Chapter 8.

³ United Nations General Assembly. Resolution adopted by the General Assembly on 4 April 2018. A/RES/72/268. Scope, modalities, format and organization of the high level meeting on the fight against tuberculosis (http://www.un.org/en/ga/search/view_doc.asp?symbol=A/RES/72/268, accessed 21 June 2018).

FIG. 2.2

The four outcome areas of the Moscow Declaration



2.4 A TB-SDG monitoring framework

Monitoring of TB-specific indicators is well established at global and national levels. For example, standardized monitoring of notifications of TB cases and their treatment outcomes at global and national levels has been in place since 1995, and WHO has been publishing annual estimates of TB incidence and mortality for more than a decade. In the era of the End TB Strategy and SDGs, such monitoring needs to be sustained, alongside continued efforts to strengthen notification and vital registration systems so that they can provide reliable data for direct measurement of TB incidence and TB deaths (see also Chapter 3), and monitoring of the newer priority indicators (five of those listed in Table 2.2 were introduced in the context of the End TB Strategy).

As explained in Section 2.2, achieving the End TB Strategy targets and milestones requires progress in reducing health-related risk factors for TB infection and disease, as well as broader social and economic determinants of TB infection and disease. As explained in Section 2.1, the SDG framework includes targets and indicators related to these risk factors and determinants, and the global ministerial conference on ending TB and the associated Moscow Declaration emphasized the need for a multisectoral approach and a multisectoral accountability framework (Section 2.3). In this context, TB monitoring needs to include analysis of selected SDG indicators that will influence the course of the TB epidemic, with findings used to inform the multisectoral actions needed to end the TB epidemic.

WHO developed a TB-SDG monitoring framework in 2017, based on previously published work that identified clear linkages between various SDG indicators and TB

BOX 2.4

Developing a draft multisectoral accountability framework to accelerate progress to end TB

Background

The first WHO global ministerial conference on TB, titled “Ending TB in the Sustainable Development Era: a multisectoral response”, was held in Moscow in November 2017. The aim was to accelerate implementation of the End TB Strategy, in recognition of the fact that investments and actions have, to date, fallen short of those needed to reach SDG and End TB Strategy targets, and to inform the UN high-level meeting on TB in September 2018.

The Moscow Declaration to End TB includes both commitments by Member States and calls for actions by global agencies and other partners.^a It addresses four key areas for action, one of which is multisectoral accountability.^b Member States committed to “supporting the development of a multisectoral accountability framework” in advance of the UN HLM on TB, and called on WHO to develop such a framework, working in close cooperation with Member States and partners, for consideration by WHO’s governing bodies (i.e. the Executive Board and World Health Assembly). The rationale for such a framework is that strengthened accountability for the response to TB at national and global levels should contribute to faster progress towards the targets and milestones of the SDGs and End TB Strategy.

The Executive Board reiterated this request in its January 2018 meeting, in the form of a decision point.^c This included a specific request, addressed to the WHO Director-General, to develop a draft framework for consideration by Member States at the World Health Assembly in May 2018, and for presentation at the UN HLM on ending TB in September 2018.

Definitions and concepts

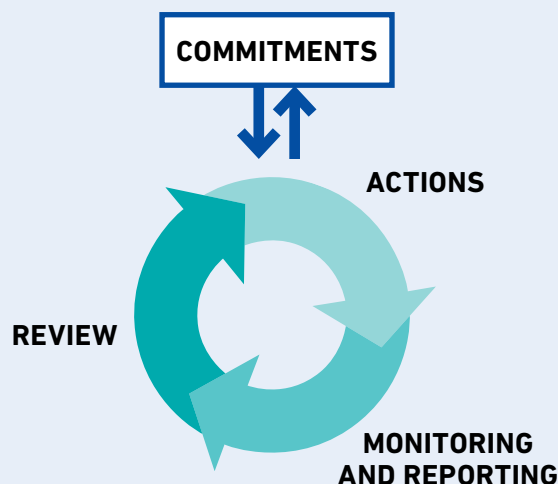
Accountability means being responsible (or answerable) for commitments made or actions taken.

An *accountability framework* defines who is accountable (e.g. an individual, organization or national government, or the global community), what commitments and actions they are accountable for, and how they will be held to account. Broadly, mechanisms for how specific entities are held to account fall into two major categories: monitoring and reporting, and review. A generic illustration of an accountability framework, represented as a cycle of components, is shown in [Fig. B2.4.1](#).^d

Conceptually, commitments should be followed by the actions needed to keep or achieve them. Monitoring and reporting are then used to track progress related to commitments and actions. Review is used to assess the results from monitoring that are documented in reports and associated products, and to make recommendations for future actions. The cycle of action, monitoring and reporting, and review can be repeated many times. The results from monitoring and reporting, and the recommendations from reviews based on those results, should drive the next cycle of actions. Periodically, new commitments or reinforcement of commitments may be required, based on reviews of progress.

FIG. B2.4.1

Generic accountability framework



Accountability can be strengthened by reinforcing one or more of the four components of the framework. Examples include adding new actions or improving existing ones; increasing the quality and coverage of data and reports available to inform reviews of progress; elevating reviews to a higher level; improving review processes (e.g. by making them more independent, more transparent and with wider participation); and ensuring that the results of reviews have meaningful consequences for action. If each of the “actions – monitoring and reporting – review” components of the cycle is strong, progress towards commitments should be faster than if one or more of those components is weak.

Development process to date

WHO prepared a background document on the development of a multisectoral accountability framework in February 2018.^e The background document was used as the basis for consultations with Member States and other partners, including in a 2-day consultation held in March 2018.^f Consultations informed the development of a “draft multisectoral accountability framework to accelerate progress to end TB”, which was posted for public review in mid-April 2018. Input from this review, from Member States and partners, was used to produce an updated draft.

The draft multisectoral accountability framework for TB that was submitted for the consideration of the 2018 World Health Assembly is available online;^g a brief outline is provided here.

The framework has two major parts: global and regional levels; and national (including local) level. The four components of each part of the framework are the same as those in the generic framework shown in [Fig. B2.4.1](#) – namely, commitments, actions, monitoring and reporting, and review.

The content of the framework is based on the following principles:

- The SDGs and the End TB Strategy, and associated political declarations form the foundation of the framework, as do existing monitoring and reporting systems, and associated best practices.
- Civil society, TB-affected communities and patient groups have a fundamental role to play in all aspects of accountability.
- It is not possible to be exhaustive in listing all elements of relevance under each of the four major components, especially at national level; hence, major examples are provided, using generic language.
- The national component of the framework requires adaptation. For example, there are differences between low and high TB burden countries, differences between countries that fund their TB responses entirely from domestic resources and those that rely on external resources for a large share of their total funding, and differences that relate to national administrative structures and legislative frameworks.

The *global and regional* part of the framework defines commitments, actions, monitoring and reporting processes, and review mechanisms that apply to all countries collectively or at regional level. The national part of the framework defines commitments, actions, monitoring and reporting processes, and review mechanisms that apply to individual countries, at national and local levels.

Discussions at the 2018 World Health Assembly and next steps

At the 2018 World Health Assembly, a resolution (EB142.R3) that included text on the UN HLM for TB and the multisectoral accountability framework was presented for consideration by Peru and the Russian Federation.^c With respect to the multisectoral accountability framework specifically, the resolution:

- welcomed the draft framework;
- requested the Secretariat to continue to develop it, working with Member States and partners;
- requested the Director-General of WHO to present the draft framework at the UN HLM on TB in September 2018; and
- requested that the Secretariat report on progress at the next World Health Assembly in 2019.

The resolution was unanimously endorsed by all Member States.

Next steps for WHO include the following:

1. Presentation of the draft framework by the WHO Director-General at the UN HLM on TB.
2. Further work on the framework, in consultation with Member States and partners. The first updates to the framework will be based on the content of the Political Declaration from the UN HLM on TB. There may be a need for additional consultations on the “review” component of the draft framework.
3. Submission of an updated version of the framework for consideration by the World Health Assembly in 2019.
4. Provision of support to Member States that express interest in beginning to adapt and use the draft multisectoral accountability framework.

^a Moscow Declaration to End TB. Geneva: WHO; 2017 (http://www.who.int/tb/Moscow_Declaration_MinisterialConference_TB/en/, accessed 21 June 2018).

^b The others areas for action were advancing the response within the 2030 Agenda for Sustainable Development; ensuring sufficient and sustainable financing; and pursuing science, research and innovation.

^c See resolution EB142.R3, operative paragraph 1 (http://apps.who.int/gb/ebwha/pdf_files/EB142/B142_R3-en.pdf, accessed 21 June 2018). Also see WHA documents A71/15 and A71/16 (http://apps.who.int/gb/ebwha/pdf_files/WHA71/A71_15-en.pdf and http://apps.who.int/gb/ebwha/pdf_files/WHA71/A71_16-en.pdf, accessed 21 June 2018).

^d This figure is derived from the unified accountability framework for women’s, children’s and adolescents’ health. That framework depicts the action–monitoring–review cycle in a circle, as here, for the global and country levels separately. The accountability framework for TB adds a component for “commitments” and highlights “monitoring and reporting” in its third component.

^e Developing a draft TB multisectoral accountability framework. Background document. Stakeholder consultation convened by Global TB Programme, World Health Organization, Chateau de Penthes, Geneva, 1–2 March 2018. Geneva: World Health Organization; 2018 (http://www.who.int/tb/TBAccountabilityFramework_Consultation1_2March_BackgroundDocument_20180228.pdf?ua=1, accessed 21 June 2018).

^f Developing a draft TB multisectoral accountability framework. Stakeholder consultation convened by the Global TB Programme, World Health Organization, Geneva, 1–2 March 2018. Meeting report (http://www.who.int/tb/TB_MAF_1_2Marchconsultation_meetingreport_20180322.pdf?ua=1, accessed 21 June 2018).

^g Developing a multisectoral accountability framework to accelerate progress to end TB. Document submitted to the 2018 World Health Assembly (A71/16.Add.1.)

incidence,^{1,2,3,4} and further analysis of the relationship between SDG indicators and TB incidence.⁵ The framework, which comprises 14 indicators under seven SDGs, is shown in **Table 2.3**.

For SDG 3, the seven indicators selected for monitoring are:

- coverage of essential health services;
- percentage of health expenditures that are out-of-pocket;
- health expenditure per capita;
- HIV prevalence;
- prevalence of smoking;
- prevalence of diabetes; and
- prevalence of alcohol use disorder.

For SDGs 1, 2, 7, 8, 10 and 11, the seven indicators selected for monitoring are:

- proportion of the population living below the international poverty line;
- proportion of the population covered by social protection floors or systems;
- prevalence of undernourishment;
- proportion of the population with primary reliance on clean fuels and technology;
- gross domestic product (GDP) per capita;
- Gini index for income inequality;⁶ and
- proportion of the urban population living in slums.

The rationale for the selection of these 14 indicators and data sources is provided in **Table 2.3**, with comments on whether it is relevant to collect data for TB patients specifically.

The framework includes only indicators for which a relationship with TB incidence could be established. It excludes:

- indicators that are sub-indicators of indicators that have already been included (e.g. sub-indicators related to UHC, under SDG 3); and

- indicators that are only remotely associated with TB risks, and that operate mainly through other SDGs (e.g. education under SDG 4, gender equality under SDG 5 and resilient infrastructure under SDG 9).

Importantly, collection and reporting of data for the 14 indicators shown in **Table 2.3** does not require any additional data collection and reporting efforts by national TB programmes. Nor does it require data collection and reporting efforts that go beyond those to which countries have already committed in the context of the SDGs.

At the global level, the UN has established a monitoring system for SDG indicators, and countries are expected to report data on an annual basis via the appropriate UN agencies (including WHO). Therefore, analysis of the status of, and trends in, the 14 indicators related to TB will be based primarily on accessing the data held in the UN's SDG database, as shown in **Table 2.3**.⁷ In some cases, the SDG indicator is not considered the best metric, and a better (but closely related) alternative has been identified and justified (five indicators under SDG 3, one under SDG 8 and one under SDG 10). In such cases, the data sources are either WHO, the Organisation for Economic Co-operation and Development (OECD), UNAIDS or the World Bank (also shown in **Table 2.3**).

The data for the indicators shown in **Table 2.3** that will be available in the WHO, OECD, UN and World Bank databases will be for national populations. These data will not be available for TB patients specifically, with the exception of data on HIV prevalence, given that HIV status among TB patients has been routinely monitored as part of national TB surveillance for more than a decade. This is not a problem for monitoring of TB risk factors and determinants, since it is the population-level prevalence that influences population-level TB risks.

For a few of the indicators included in **Table 2.3**, collection of data for TB patients specifically could be considered. However, there is a clear risk of making routine TB surveillance far too complex, and this risk needs to be avoided. If the indicator is considered important enough to monitor among TB patients at country level, periodic surveys should be considered as an alternative to routine surveillance (in which data would be collected for all TB patients).

Analysis of the indicators in the TB-SDG monitoring framework for high TB burden countries is included in **Chapter 7**. The latest status and recent trends in each indicator are also shown for these countries on the second page of the country profiles in **Annex 2** (this information is shown for other countries in profiles that are available online).⁸

¹ Lönnroth K, Jaramillo E, Williams B, Dye C, Raviglione M. Tuberculosis: the role of risk factors and social determinants. In: Blas E, Kurup A (editors), *Equity, social determinants and public health programmes*. Geneva: WHO; 2010 (http://apps.who.int/iris/bitstream/10665/44289/1/9789241563970_eng.pdf, accessed 2 August 2017).

² Lönnroth K, Castro KG, Chakaya JM, Chauhan LS, Floyd K, Glaziou P, et al. Tuberculosis control and elimination 2010–50: cure, care, and social development. *Lancet*. 2010;375(9728):1814–29 (<https://www.ncbi.nlm.nih.gov/pubmed/20488524>, accessed 2 August 2017).

³ Lienhardt C, Glaziou P, Uplekar M, Lönnroth K, Getahun H, Raviglione M. Global tuberculosis control: lessons learnt and future prospects. *Nat Rev Microbiol*. 2012;10(6):407–16 (<https://www.ncbi.nlm.nih.gov/pubmed/22580364>, accessed 2 August 2017).

⁴ Lönnroth K, Jaramillo E, Williams BG, Dye C, Raviglione M. Drivers of tuberculosis epidemics: the role of risk factors and social determinants. *Soc Sci Med*. 2009;68(12):2240–6 (<https://www.ncbi.nlm.nih.gov/pubmed/19394122>, accessed 2 August 2017).

⁵ Monitoring and evaluation of TB in the context of the Sustainable Development Goals: Background Paper for WHO Ministerial Conference on “TB in the context of the Sustainable Development Goals”. Available on request.

⁶ The index can take values between 0 and 1, with 0 representing perfect equality and 1 representing perfect inequality.

⁷ Further details are provided in **Annex 1**.

⁸ <http://www.who.int/tb/data/en/>

TABLE 2.3A

TB-SDG monitoring framework: indicators to monitor within SDG 3

SDG 3: Ensure healthy lives and promote well-being for all at all ages					
SDG TARGETS FOR 2030	SDG INDICATORS	ALTERNATIVE INDICATORS TO MONITOR	RATIONALE	DATA SOURCE	COLLECT DATA FOR TB PATIENTS SPECIFICALLY?
3.3 End the epidemics of AIDS, TB, malaria and neglected tropical diseases and combat hepatitis, water-borne diseases and other communicable diseases	3.3.1 Number of new HIV infections per 1000 uninfected population 3.3.2 TB incidence per 100 000 population	HIV prevalence	HIV is a strong risk factor for development of TB disease and is associated with poorer treatment outcomes. HIV prevalence (rather than incidence) will be monitored because it is directly measured and those newly infected with HIV are at lower risk of developing TB compared with those who have been infected for more than 1 year.	UNAIDS	Yes, already routinely collected.
				WHO	NA
3.4 Reduce premature mortality by one third from non-communicable diseases and promote mental health and well-being	3.4.1 Mortality rate attributed to cardiovascular disease, cancer, diabetes or chronic respiratory disease	Prevalence of diabetes	Diabetes is a strong risk factor for development of TB disease, although a link with TB incidence at the national (as opposed to individual) level has been difficult to establish due to confounding. Diabetes prevalence is more relevant than mortality for TB since it directly influences the risk of developing TB.	WHO	Could be considered at country level, to inform planning of care for comorbidities.
3.5 Strengthen prevention and treatment of substance abuse, including narcotic drug abuse and harmful use of alcohol	3.5.2 Alcohol consumption per capita per year (in litres of pure alcohol) among those aged ≥15 years (harmful level defined nationally)	Prevalence of alcohol use disorder	Alcohol use is a strong risk factor for TB disease and poorer treatment outcomes at the individual level, although a link with TB incidence at the national (as opposed to individual) level has been hard to establish due to confounding. The prevalence of alcohol use disorder is the most relevant indicator in the context of TB.	WHO	Could be considered at country level, to inform planning of care for comorbidities.
3.8 Achieve UHC, including financial risk protection, access to quality essential health-care services and access to safe, effective, quality and affordable essential medicines and vaccines for all	3.8.1 Coverage of essential health services (defined as the average coverage of essential services based on tracer interventions that include reproductive, maternal, newborn and child health, infectious diseases, non-communicable diseases and service capacity and access, among the general and the most disadvantaged population). 3.8.2 Proportion of population with large household expenditures on health as a share of total household expenditure or income	NA	Achieving UHC is required to achieve the three high-level targets of the End TB Strategy for reductions in the TB incidence rate, the number of TB deaths and eliminating catastrophic costs for TB patients and their households. TB treatment coverage has been monitored for years and is one of the 16 tracer indicators that have been selected to measure SDG indicator 3.8.1. Health expenditure per capita is correlated with TB incidence.	WHO	To assess progress in elimination of catastrophic costs for TB patients and their households, periodic surveys of TB patients are recommended.
		Percentage of health expenditures that are out-of-pocket Health expenditure per capita			
3.a Strengthen implementation of the WHO Framework Convention on Tobacco Control	3.a.1 Age-standardized prevalence of current tobacco use among those aged ≥15 years	Prevalence of smoking among those aged ≥15 years (%)	Smoking is a strong risk factor for TB disease at the individual level, although a link with TB incidence at the national (as opposed to individual) level has been difficult to establish due to confounding.	WHO	Could be considered (e.g. to inform access to smoking cessation interventions).

TABLE 2.3B

TB-SDG monitoring framework: indicators to monitor beyond SDG 3

SDG 1: End poverty in all its forms everywhere					
SDG TARGETS FOR 2030	SDG INDICATORS	ALTERNATIVE INDICATORS TO MONITOR	RATIONALE	DATA SOURCE	COLLECT DATA FOR TB PATIENTS SPECIFICALLY?
1.1 Eradicate extreme poverty for all people everywhere	1.1.1 Proportion of population living below the international poverty line	NA	Poverty is a strong risk factor for TB, operating through several pathways. Reducing poverty should also facilitate prompt health-care seeking.	UN SDG database, World Bank	No
1.3 Nationally appropriate social protection systems and measures for all, including floors	1.3.1 Proportion of population covered by social protection floors/systems	NA	Countries with higher levels of social protection have lower TB burden. Progress on both indicators will help to achieve the End TB Strategy target to eliminate catastrophic costs for TB patients and their households.		Could be considered (e.g. to facilitate access to social protection).
SDG 2: End hunger, achieve food security and improved nutrition and promote sustainable agriculture					
2.1 End hunger and ensure access by all people to safe, nutritious and sufficient food year-round	2.1.1 Prevalence of undernourishment	NA	Under-nutrition weakens the body's defence against infections and is a strong risk factor for TB at the national and individual level.	UN SDG database	Could be considered (e.g. to plan food support).
SDG 7: Ensure access to affordable, reliable, sustainable, and modern energy for all					
7.1 Ensure universal access to affordable, reliable and modern energy services	7.1.2 Proportion of population with primary reliance on clean fuels and technology	NA	Indoor air pollution is a risk factor for TB disease at the individual level. There has been limited study of ambient air pollution but it is plausible that it is linked to TB incidence.	WHO	No
SDG 8: Promote sustained, inclusive and sustainable economic growth, full and productive employment and decent work for all					
8.1 Sustain per capita growth with at least 7% growth in GDP per year in the least developed countries	8.1.1 Annual growth rate of real GDP per capita	GDP per capita	Historic trends in TB incidence are closely correlated with changes in the absolute level of GDP per capita (but not with the growth rate).	World Bank	No
SDG 10: Reduce inequality within and among countries					
10.1 Achieve and sustain income growth of the bottom 40% of the population at a rate higher than the national average	10.1.1 Growth rates of household expenditure or income per capita, overall and for the bottom 40% of the population	Gini index for income inequality	TB is a disease of poverty, and decreasing income inequalities combined with economic growth should have an effect on the TB epidemic.	World Bank OECD	No
SDG 11: Make cities and human settlements inclusive, safe, resilient and sustainable					
11.1 Ensure access for all to adequate, safe and affordable housing and basic services and upgrade slums	11.1.1 Proportion of urban population living in slums, informal settlements or inadequate housing	NA	Living in a slum is a risk factor for TB transmission due to its link with overcrowding. It is also a risk factor for developing TB disease, due to links with air pollution and under-nutrition.	UN SDG database	No

2.5 WHO's General Programme of Work 2019–2023

WHO's GPW for 2019–2023 was drafted in 2017; it was then revised following review by WHO's Executive Board in January 2018, and a final version adopted by the World Health Assembly in May 2018.¹ The thirteenth in a series of GPWs since WHO was founded in 1948, the current GPW sets out the organization's strategic direction for the next 5 years.

The GPW for 2019–2023 is based on the foundation of the SDGs and is relevant to all countries: low-, middle- and high-income. In the context of SDG 3 in particular (Box 2.2), GPW 13 provides a vision that is rooted in Article 1 of WHO's Constitution: "A world in which all people attain the highest possible standard of health and well-being". GPW 13 also recognizes the influence of other SDGs on health, and the need for multisectoral approaches to address the social, economic and environmental determinants of health.

The GPW is structured around three strategic priorities and associated goals (Fig. 2.3). The three strategic priorities are UHC, addressing health emergencies and promoting healthier populations. The associated goals for 2023 are the so-called "triple billion goals": that 1 billion more people are benefiting from UHC, 1 billion more people are better protected from health emergencies, and 1 billion more people are enjoying better health and well-being. WHO and the World Bank have estimated that only about half the world's population had access to essential health services in 2015.² Achieving all three goals depends on joint efforts by Member States, WHO and other partners.

The GPW goal related to UHC is directly linked to SDG Target 3.8 (Box 2.2). As explained in Section 2.2, progress towards this goal will be crucial to reaching End TB Strategy milestones and targets. Similarly, progress in TB prevention and care will contribute to the UHC target, with TB treatment being one of the tracer indicators for SDG Target 3.8 (Section 2.1, Table 2.3), and to more people enjoying better health and well-being. Protecting people from the public health threat posed by TB contributes to the health emergencies goal. At the same time, broader progress towards the goals related to health emergencies and better health and well-being contribute to progress towards End TB Strategy targets and milestones by positively influencing the determinants of TB included in the TB-SDG monitoring framework shown in Table 2.3.

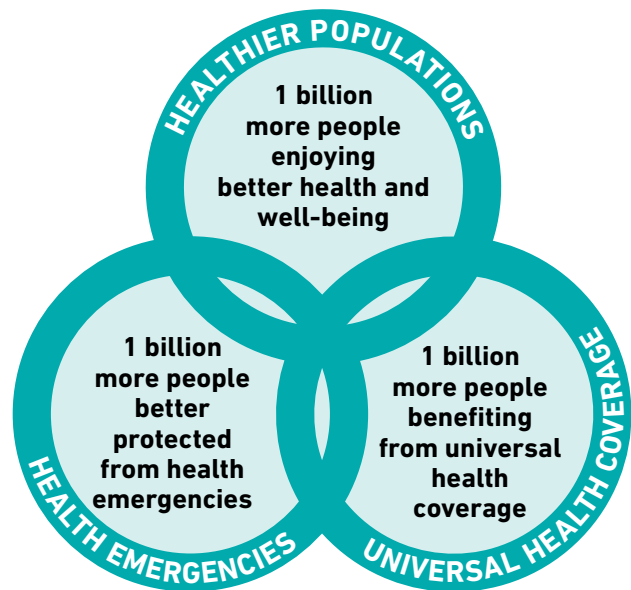
GPW 13 includes 10 outcomes, one of which is based

¹ World Health Organization. Report by the Director General. A71/4. Draft thirteenth general programme of work 2019–2023 (http://apps.who.int/gb/ebwha/pdf_files/WHA71/A71_4-en.pdf?ua=1, accessed 21 June 2018).

² World Health Organization/World Bank Group. Tracking universal health coverage: 2017 global monitoring report. Geneva: WHO; 2017 (<http://apps.who.int/iris/bitstream/handle/10665/259817/9789241513555-eng.pdf>, accessed 21 June 2018).

FIG. 2.3

WHO's 13th General Programme of Work: a set of interconnected strategic priorities and goals to ensure healthy lives and promote well-being for all at all ages



on the SDG 3 target related to ending epidemics. Outcome 5 is defined as "Accelerated elimination and eradication of high-impact communicable diseases". Under this outcome, there are two targets for TB: that the number of TB deaths is reduced by at least 50% between 2018 and 2023, and that by 2023 there is at least 80% treatment coverage for people with drug-resistant TB.

Linked to the GPW, WHO's Global TB Programme has also defined two other targets:

- 40 million people with TB are reached with care during the period 2018–2022, including 3.5 million children and 1.5 million people with drug-resistant TB;
- At least 30 million people are reached with TB prevention services during the period 2018–2022.

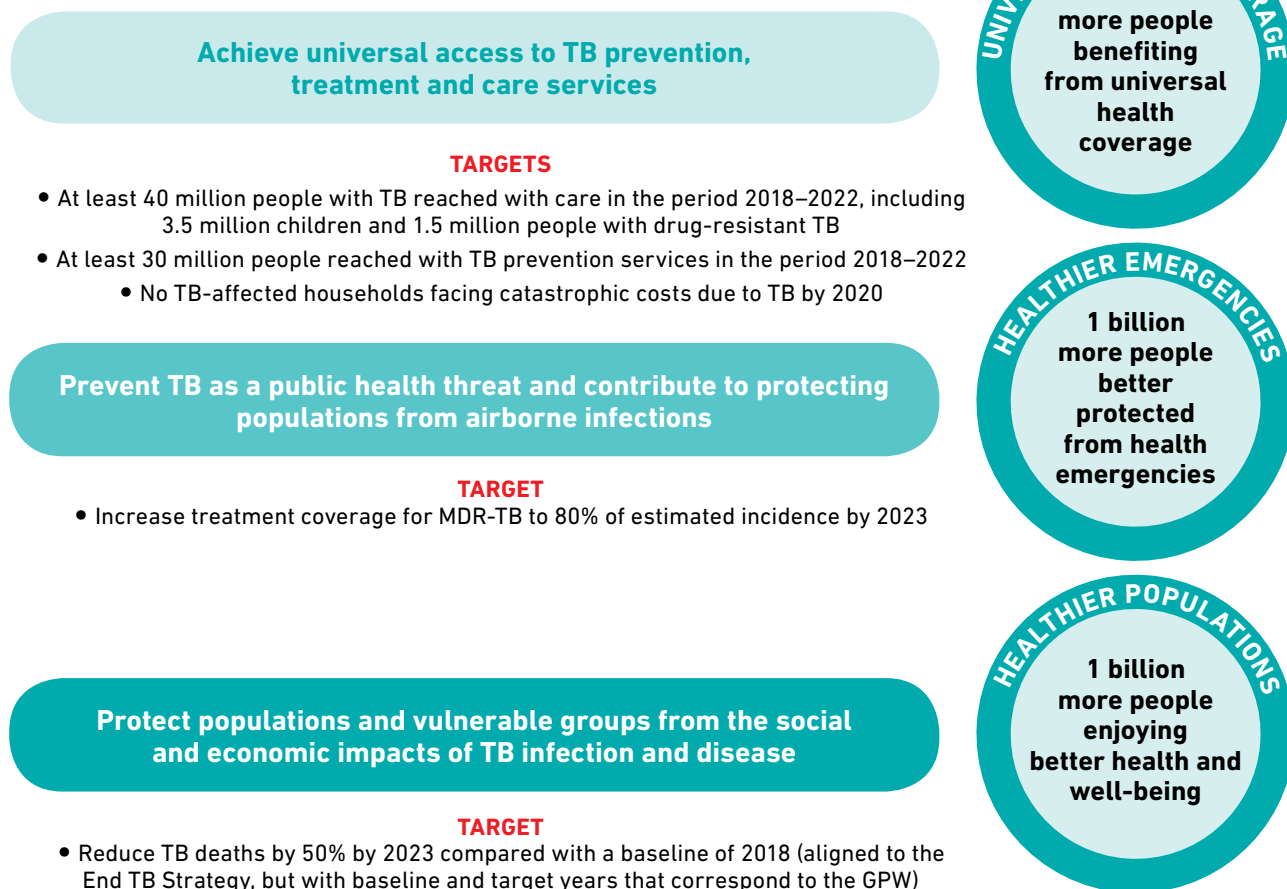
TB targets that have been aligned with the GPW are illustrated in Fig. 2.4. To catalyze global efforts to support the achievement of these targets, WHO, the Stop TB Partnership and the Global Fund to Fight AIDS, Tuberculosis and Malaria have launched a joint initiative, called "Find.Treat.All" (Box 2.5).

2.6 Lists of high-burden countries being used by WHO during the period 2016–2020

During the period 1998–2015, the concept of an HBC became familiar and widely used in the context of TB. In 2015, three HBC lists – for TB, TB/HIV and MDR-TB – were in use. The HBC list for TB (22 countries) had remained unchanged since 2002; also, the HBC lists for TB/HIV (41 countries) had not been updated since 2009, and those for MDR-TB (27 countries) had not been updated since 2008.

FIG. 2.4

Strategic priorities and targets for TB aligned with WHO's General Programme of Work



BOX 2.5

FIND.TREAT.ALL: a joint initiative to scale up the global response towards universal access to TB prevention and care, 2018–2022

Following the first global ministerial conference on TB in 2017 and in the advance of the UN high-level meeting on TB in 2018, WHO, the Stop TB Partnership, and The Global Fund to Fight AIDS, Tuberculosis and Malaria have launched a joint initiative to scale up the End TB response towards universal access to TB prevention and care. The initiative is for the five-year period 2018–2022.

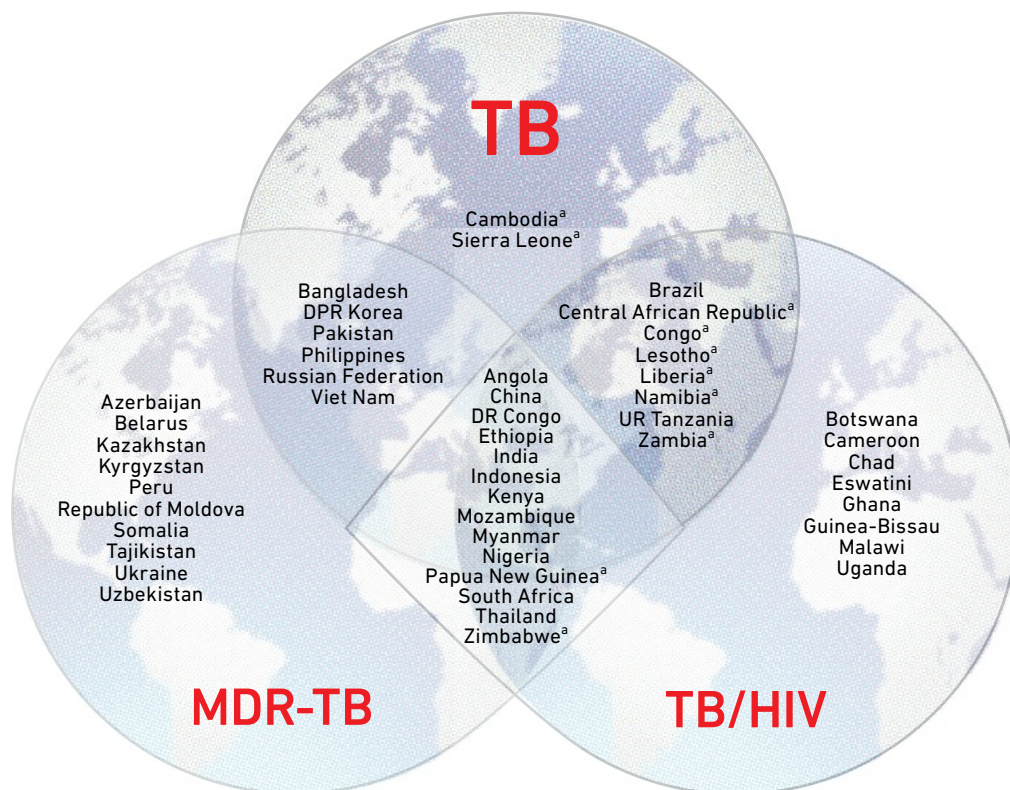
The initiative includes a target to diagnose, treat and report 40 million people with TB, including 3.5 million children and 1.5 million people with drug-resistant-TB, between 2018 and 2022.

Achieving the targets will require the efforts of many stakeholders, including country leaders, government ministries, civil society and TB-affected communities, the private sector and global agencies. Global, regional and national responses will need to be transformed in terms of commitments, actions to accelerate access to prevention and care, and measurement of progress.

The initiative encompasses all countries, with priority given to the 30 high TB burden countries. It has become a flagship initiative at WHO.

FIG. 2.5

Countries in the three high-burden country lists for TB, TB/HIV and MDR-TB being used by WHO during the period 2016–2020, and their areas of overlap



^a Indicates countries that are included in the list of 30 high TB burden countries on the basis of the severity of their TB burden (i.e. TB incident cases per 100 000 population per year), as opposed to the top 20, which are included on the basis of their absolute number of incident cases per year. Also see [Table 2.4](#).

With 2015 marking the end of the MDGs and a new era of SDGs, as well as the last year of the Stop TB Strategy before its replacement with the End TB Strategy, it was an ideal time to revisit these three HBC lists.

Following a wide consultation process,¹ in 2015 WHO defined three HBC lists for the period 2016–2020: one for TB, one for MDR-TB and one for TB/HIV ([Fig. 2.5](#), [Table 2.4](#)).² Each list contains 30 countries ([Table 2.4](#)). These are defined as the top 20 countries in terms of the absolute number of estimated incident cases, plus the additional 10 countries with the most severe burden in terms of incidence rates per capita that do not already appear in the top 20 and that meet a minimum threshold in terms of their absolute numbers of incident cases (10 000 per year for TB, and 1000 per year for TB/HIV and MDR-TB). The lists were defined using the latest estimates of TB disease burden available in October 2015. Each list accounts for about 90% of the global burden,

with most of this accounted for by the top 20 countries in each list.

There is overlap among the three lists, but 48 countries appear in at least one list. The 14 countries that are in all three lists (shown in the central diamond in [Fig. 2.5](#)) are Angola, China, the Democratic Republic of the Congo, Ethiopia, India, Indonesia, Kenya, Mozambique, Myanmar, Nigeria, Papua New Guinea, South Africa, Thailand and Zimbabwe.³

The 30 high TB burden countries are given particular attention in the main body of this report. Where estimates of disease burden and assessment of progress in the response are for TB/HIV and MDR-TB specifically, the countries in the TB/HIV and MDR-TB lists, respectively, are given particular attention. [Annex 2](#) contains a two-page profile for each of the 30 high TB burden countries; the annex has a clear demarcation between the 20 countries included on the basis of absolute numbers of incident cases and the 10 additional countries included on the basis of the incidence rate per capita. Country profiles for all countries (with the same content as those presented in [Annex 2](#)) are also available online.⁴

¹ World Health Organization Strategic and Technical Advisory Group for TB. Use of high-burden country lists for TB by WHO in the post-2015 era (discussion paper). Geneva: WHO; 2015 (http://www.who.int/tb/publications/global_report/high_tb_burdencountrylists2016-2020.pdf, accessed 21 June 2018).

² As explained in the last row of [Table 2.4](#), the three lists have a lifetime of 5 years, and the countries included in each list and the criteria used to define each list will be reviewed in June 2020.

³ These 14 countries accounted for 64% of the estimated global number of incident TB cases in 2017.

⁴ See: www.who.int/tb/data

TABLE 2.4

The three high-burden country lists for TB, TB/HIV and MDR-TB being used by WHO during the period 2016–2020

LIST	THE 30 HIGH TB BURDEN COUNTRIES		THE 30 HIGH TB/HIV BURDEN COUNTRIES		THE 30 HIGH MDR-TB BURDEN COUNTRIES	
Purpose and target audience	To provide a focus for global action on TB in the countries where progress is most needed to achieve End TB Strategy and SDG targets and milestones, to help build and sustain national political commitment and funding in the countries with the highest burden in terms of absolute numbers or severity, and to promote global monitoring of progress in a well-defined set of countries.		To provide a focus for global action on HIV-associated TB in the countries where progress is most needed to achieve End TB Strategy, UNAIDS and SDG targets and milestones, to help build and sustain national political commitment and funding in the countries with the highest burden in terms of absolute numbers or severity, and to promote global monitoring of progress in a well-defined set of countries.		To provide a focus for global action on the MDR-TB crisis in the countries where progress is most needed to achieve End TB Strategy targets and milestones, to help build and sustain national political commitment and funding in the countries with the highest burden in terms of absolute numbers or severity, and to promote global monitoring of progress in a well-defined set of countries.	
Definition	The 20 countries with the highest estimated numbers of incident TB cases, plus the top 10 countries with the highest estimated TB incidence rate that are not in the top 20 by absolute number (threshold, >10 000 estimated incident TB cases per year).		The 20 countries with the highest estimated numbers of incident TB cases among people living with HIV, plus the top 10 countries with the highest estimated TB/HIV incidence rate that are not in the top 20 by absolute number (threshold, >1000 estimated incident TB/HIV cases per year).		The 20 countries with the highest estimated numbers of incident MDR-TB cases, plus the top 10 countries with the highest estimated MDR-TB incidence rate that are not in the top 20 by absolute number (threshold, >1000 estimated incident MDR-TB cases per year).	
Countries in the list	<i>The top 20 by estimated absolute number (in alphabetical order):</i> Angola Bangladesh Brazil China DPR Korea DR Congo Ethiopia India Indonesia Kenya Mozambique Myanmar Nigeria Pakistan Philippines Russian Federation South Africa Thailand UR Tanzania Viet Nam	<i>The additional 10 by estimated incidence rate per 100 000 population and with a minimum number of 10 000 cases per year (in alphabetical order):</i> Cambodia Central African Republic Congo Lesotho Liberia Namibia Papua New Guinea Sierra Leone Zambia Zimbabwe	<i>The top 20 by estimated absolute number (in alphabetical order):</i> Angola Brazil Cameroon China DR Congo Ethiopia India Indonesia Kenya Lesotho Malawi Mozambique Myanmar Nigeria South Africa Thailand Uganda UR Tanzania Zambia Zimbabwe	<i>The additional 10 by estimated incidence rate per 100 000 population and with a minimum number of 1000 cases per year (in alphabetical order):</i> Botswana Central African Republic Chad Congo Eswatini Ghana Guinea-Bissau Liberia Namibia Papua New Guinea	<i>The top 20 by estimated absolute number (in alphabetical order):</i> Bangladesh China DPR Korea DR Congo Ethiopia India Indonesia Kazakhstan Kenya Mozambique Myanmar Nigeria Pakistan Philippines Russian Federation South Africa Thailand Ukraine Uzbekistan Viet Nam	<i>The additional 10 by estimated incidence rate per 100 000 population and with a minimum number of 1000 cases per year (in alphabetical order):</i> Angola Azerbaijan Belarus Kyrgyzstan Papua New Guinea Peru Republic of Moldova Somalia Tajikistan Zimbabwe
Share of global incidence in 2017 (%)	84%	2.9%	83%	4.7%	87%	4.4%
Lifetime of list	5 years (review criteria and included countries in June 2020).		5 years (review criteria and included countries in June 2020).		5 years (review criteria and included countries in June 2020).	



Transporting chest X-ray equipment during the 2016 national TB prevalence survey of the Philippines

Raldy Benavente / FACE Inc (Philippines)

Chapter 3. TB disease burden

KEY FACTS AND MESSAGES

Worldwide, tuberculosis (TB) is one of the top 10 causes of death, and the leading cause from a single infectious agent (above HIV/AIDS); millions of people continue to fall sick with the disease each year.

In 2017, TB caused an estimated 1.3 million deaths (range, 1.2–1.4 million) among HIV-negative people, and there were an additional 300 000 deaths from TB (range, 266 000–335 000) among HIV-positive people.^a There were an estimated 10.0 million new cases of TB (range, 9.0–11.1 million), equivalent to 133 cases (range, 120–148) per 100 000 population.

TB affects all countries and all age groups, but overall the best estimates for 2017 were that 90% of cases were adults (aged ≥15 years), 64% were male, 9% were people living with HIV (72% of them in Africa) and two thirds were in eight countries: India (27%), China (9%), Indonesia (8%), the Philippines (6%), Pakistan (5%), Nigeria (4%), Bangladesh (4%) and South Africa (3%). Only 6% of cases were in the WHO European Region and the WHO Region of the Americas, each of which had 3% of cases.

The severity of national epidemics varies widely. In 2017, there were under 10 new cases per 100 000 population in most high-income countries, 150–400 in most of the 30 high TB burden countries, and above 500 in a few countries including Mozambique, the Philippines and South Africa.

Globally in 2017, there were an estimated 558 000 new cases (range, 483 000–639 000) of rifampicin-resistant TB (RR-TB), of which almost half were in three countries: India (24%), China (13%) and the Russian Federation (10%). Among RR-TB cases, an estimated 82% had multidrug-resistant TB (MDR-TB).

Globally, 3.6% of new TB cases and 17% of previously treated cases had MDR/RR-TB, with the highest proportions (>50% in previously treated cases) in countries of the former Soviet Union.

Globally, the absolute number of deaths from TB among HIV-negative people has been estimated to have fallen by 29% since 2000, from 1.8 million in 2000 to 1.3 million in 2017, and by 5% since 2015 (the baseline year for targets set in the End TB Strategy). The number of TB deaths among HIV-positive people^a

has fallen by 44% since 2000, from 534 000 in 2000 to 300 000 in 2017, and by 20% since 2015.

The TB mortality rate (TB deaths among HIV-negative people per 100 000 population per year) is falling at about 3% per year, and the best estimate for the overall reduction during 2000–2017 is 42%. The fastest regional declines in mortality rates in the 5-year period 2013–2017 were in the WHO European Region and WHO South-East Asia Region (11% and 4% per year, respectively).

Worldwide, TB incidence (new cases per 100 000 population per year) is falling at about 2% per year. The fastest regional declines from 2013 to 2017 were in the WHO European Region (5% per year) and WHO African Region (4% per year). In the same 5 years, particularly impressive reductions (4–8% per year) occurred in southern Africa (e.g. Eswatini [formerly Swaziland], Lesotho, Namibia, South Africa, Zambia, Zimbabwe) following a peak in the HIV epidemic, and the expansion of TB and HIV prevention and care, and in the Russian Federation (5% per year) following intensified efforts to reduce the burden of TB.

In 2017, the best estimate of the proportion of people with TB who died from the disease (the case fatality ratio, CFR) was 16%, down from 23% in 2000. The CFR needs to fall to 10% by 2020 to reach the first milestones of the End TB Strategy. There is considerable country variation in the CFR, from under 5% in a few countries to more than 20% in most countries in the WHO African Region, demonstrating large inequalities among countries in access to TB diagnosis and treatment.

National notification and vital registration systems need to be strengthened towards the goal of direct measurement of TB incidence and mortality in all countries. National TB prevalence surveys provide an interim approach to directly measuring the burden of TB disease in an important subset of high TB burden countries; 25 surveys were completed between 2007 and the end of 2017.

^a When an HIV-positive person dies from TB disease, the underlying cause is classified as HIV in the international classification of diseases system (ICD-10).

The burden of tuberculosis (TB) disease can be measured in terms of:

- *incidence* – the number of new and relapse cases of TB arising in a given time period, usually 1 year;
- *prevalence* – the number of cases of TB at a given point in time; and
- *mortality* – the number of deaths caused by TB in a given time period, usually 1 year.

Global targets and milestones for reductions in the burden of TB disease have been set as part of the Sustainable Development Goals (SDGs) and WHO's End TB Strategy (Chapter 2).¹ SDG 3 includes a target to end the global TB epidemic by 2030, with TB incidence (new and relapse cases per 100 000 population per year) defined as the indicator for measurement of progress. The 2030 targets set in the End TB Strategy are a 90% reduction in TB deaths and an 80% reduction in TB incidence, compared with levels in 2015. Targets for 2035 and milestones for 2020 and 2025 have also been defined (Table 3.1).

TABLE 3.1
Targets for percentage reductions in TB disease burden set in WHO's End TB Strategy

INDICATORS	MILESTONES		TARGETS	
	2020	2025	2030	2035
Percentage reduction in the absolute number of TB deaths per year (compared with 2015 baseline)	35	75	90	95
Percentage reduction in the TB incidence rate (new and relapse cases per 100 000 population per year) (compared with 2015 baseline)	20	50	80	90

This chapter has five major sections. Section 3.1 and Section 3.2 present the latest WHO estimates of TB incidence and mortality between 2000 and 2017, and highlight sources of data and actions needed to improve measurement of TB incidence and mortality. Section 3.3 focuses on the burden of drug-resistant TB, including progress in global surveillance of resistance to anti-TB drugs, and estimates of the incidence of multidrug-resistant TB (MDR-TB) and rifampicin-resistant TB (RR-TB). Section 3.4 discusses national TB prevalence surveys. TB prevalence is not an indicator for which a global target has been set during the period 2016–2035.² Nevertheless, in many countries, national TB prevalence

¹ World Health Organization. WHO End TB Strategy: global strategy and targets for tuberculosis prevention, care and control after 2015. Geneva: WHO; 2015 (http://www.who.int/tb/post2015_strategy/en/, accessed 30 August 2018).

² This is in contrast to the period covered by the Stop TB Strategy (2006–2015), when the target was to halve prevalence by 2015 compared with a baseline of 1990.

surveys still provide the best method for estimating the burden of TB disease (including by age and sex) and for planning actions needed to reduce that burden. In addition, results from national TB prevalence surveys can inform estimates of TB incidence and mortality, and thus contribute to monitoring of progress towards SDG and End TB Strategy targets. Finally, Section 3.5 covers estimates of TB incidence and mortality, disaggregated by age and sex. This is in line with the increasing emphasis on the importance of within-country disaggregation of key indicators in the SDGs and the End TB Strategy (Chapter 2).

WHO updates its estimates of the burden of TB disease annually, using the latest available data and analytical methods.^{3,4} Since 2006, concerted efforts have been made to improve the available data and methods used, under the umbrella of the WHO Global Task Force on TB Impact Measurement (Box 3.1). A summary of the main updates to available data and methods since the 2017 global TB report is provided in Box 3.2. To put these updates in a broader context, comparisons of the annual updates made by WHO for TB are compared with those for HIV and malaria, as well as with estimates for TB that are updated annually by the Institute of Health Metrics and Evaluation (IHME) at the University of Washington, United States of America (USA), in Box 3.3.

3.1 TB incidence

3.1.1 Methods to estimate TB incidence

TB incidence has never been measured at national level because this would require long-term studies among large cohorts (hundreds of thousands) of people, which would involve high costs and challenging logistics. However, notifications of TB cases provide a good proxy indication of TB incidence in countries that have high-performance surveillance systems (e.g. with little underreporting of diagnosed cases), and in which the quality of and access to health care means that few cases are not diagnosed.

The ultimate goal is to directly measure TB incidence from TB notifications in all countries. This requires a combination of strengthened surveillance, better quantification of underreporting (i.e. the number of cases that are missed by surveillance systems)⁵ and universal health coverage. A TB surveillance checklist developed by the WHO Global Task Force on TB Impact Measurement (Box 3.1) defines the standards that need to be met for

³ The online technical appendix is available at www.who.int/tb/data.

⁴ The updates can affect the entire time series back to 2000. Therefore, estimates presented in this chapter for 2000–2016 supersede those of previous reports, and direct comparisons (e.g. between the 2016 estimates in this report and the 2016 estimates in the previous report) are not appropriate.

⁵ Inventory studies can be used to measure the number of cases that are diagnosed but not reported. For a guide to inventory studies, see World Health Organization. Assessing tuberculosis underreporting through inventory studies. Geneva: WHO; 2012 (http://www.who.int/tb/publications/inventory_studies/en/, accessed 15 August 2018).

BOX 3.1

The WHO Global Task Force on TB Impact Measurement

Establishment and progress made, 2006–2015

The WHO Global Task Force on TB Impact Measurement (hereafter referred to as the Task Force) was established in 2006 and is convened by the TB Monitoring and Evaluation unit of WHO's Global TB Programme. Its original aim was to ensure that WHO's assessment of whether 2015 targets set in the context of the Millennium Development Goals (MDGs) were achieved at global, regional and country levels was as rigorous, robust and consensus-based as possible. Three strategic areas of work were pursued:

- strengthening routine surveillance of TB cases (via national notification systems) and deaths (via national VR systems) in all countries;
- undertaking national TB prevalence surveys in 22 global focus countries; and
- periodically reviewing methods used to produce TB disease burden estimates.

Work on strengthened surveillance included the following:

- Development of a TB surveillance checklist of standards and benchmarks (with 10 core and three supplementary standards).³ This checklist can be used to systematically assess the extent to which a surveillance system meets the standards required for notification and VR data to provide a direct measurement of TB incidence and mortality, respectively. By the end of 2015, 38 countries including 16 high TB burden countries had used the checklist. The global status of progress in using the checklist by August 2018 is shown in [Fig. 3.1](#).
- Electronic recording and reporting. Case-based electronic databases are the reference standard for recording and reporting TB surveillance data. A guide was produced in 2011,^b and efforts to introduce such systems were supported. The global status of progress in case-based electronic surveillance for TB by August 2018 is highlighted in [Chapter 4](#).
- Development of a guide on inventory studies to measure underreporting of detected TB cases,^c and support such studies in priority countries. An inventory study can be used to quantify the number of cases that are detected but not reported to national surveillance systems, and can serve as a basis for improving estimates of TB incidence and addressing gaps in reporting. The global status of progress in implementation of inventory studies by August 2018 is shown in [Fig. 3.1](#). An excellent recent example of a national inventory study, in Indonesia, is profiled in [Chapter 4](#).
- Expanded use of data from VR systems and mortality surveys to produce estimates of the number of TB deaths, and contributions to wider efforts to promote VR systems. By 2015, VR data were used to produce estimates of TB mortality in 127 countries, up from three in 2008.

There was substantial success in the implementation of national TB prevalence surveys in the period 2007–2015, which has continued. Between 2007 and the end of 2015, a total of 23 countries completed a survey and a further two had done so by the end of 2017; this included 18 of the 22

global focus countries. A Task Force subgroup undertook a major review and update of methods between June 2008 and October 2009. A second thorough and comprehensive review was undertaken in 2015, with consensus reached on methods to be used for the 2015 targets assessment published in WHO's 2015 global TB report.^d

Updated strategic areas of work, 2016–2020

In the context of a new era of SDGs and WHO's End TB Strategy, the Task Force met in April 2016 to review and reshape its mandate and strategic areas of work for the post-2015 era. An updated mandate and five strategic areas of work for the period 2016–2020 were agreed.^e

The mandate was defined as follows:

- To ensure that assessments of progress towards End TB Strategy and SDG targets and milestones at global, regional and country levels are as rigorous, robust and consensus-based as possible.
- To guide, promote and support the analysis and use of TB data for policy, planning and programmatic action.

The five strategic areas of work are as follows:

1. Strengthening national notification systems for direct measurement of TB cases, including drug-resistant TB and HIV-associated TB specifically.
2. Strengthening national VR systems for direct measurement of TB deaths.
3. Priority studies to periodically measure TB disease burden, including:
 - a. national TB prevalence surveys
 - b. drug-resistance surveys
 - c. mortality surveys
 - d. surveys of costs faced by TB patients and their households.
4. Periodic review of methods used by WHO to estimate the burden of TB disease and latent TB infection.
5. Analysis and use of TB data at country level, including:
 - a. disaggregated analyses (e.g. by age, sex and location) to assess inequalities and equity;
 - b. projections of disease burden; and
 - c. guidance, tools and capacity-building.

The SDG and End TB Strategy targets and milestones referred to in the mandate are the targets (2030, 2035) and milestones (2020, 2025) set for the three high-level indicators; that is, TB incidence, the number of TB deaths and the percentage of TB-affected households that face catastrophic costs as a result of TB disease ([Chapter 2](#)).

Strategic areas of work 1–3 are focused on direct measurement of TB disease burden (epidemiological and, in the case of cost surveys, economic). The underlying principle for the Task Force's work since 2006 has been that estimates of the level of and trends in disease burden should be based on direct measurements from routine surveillance and surveys as much as possible (as opposed to indirect estimates based on modelling and expert opinion). However, strategic area of work 4 remains necessary because indirect estimates

will be required until all countries have the surveillance systems or the periodic studies required to provide direct measurements. Strategic area of work 5 recognizes the importance of analysing and using TB data at country level (as well as generating data, as in strategic areas of work 1–3), including the disaggregated analyses that are now given much greater attention in the SDGs and End TB Strategy.

In the years up to 2020, the top priorities for the Task Force are strengthening of national notification and VR systems as the basis for direct measurement of TB incidence and TB mortality.

Further details about the work of the Task Force are available online;^f an up-to-date summary is provided in the latest brochure about its work.^g

^a World Health Organization. Standards and benchmarks for tuberculosis surveillance and vital registration systems: checklist and user guide. Geneva: WHO; 2014 (<http://www.who.int/tb/publications/standardsandbenchmarks/en/>, accessed 25 July 2018).

^b World Health Organization. Electronic recording and reporting for tuberculosis care and control. Geneva: WHO; 2012 (http://www.who.int/tb/publications/electronic_recording_reporting/en/, accessed 25 July 2018).

^c World Health Organization. Assessing tuberculosis underreporting through inventory studies. Geneva: WHO; 2012 (http://www.who.int/tb/publications/inventory_studies/en/, accessed 15 August 2018).

^d World Health Organization Global Task Force on TB Impact Measurement. Third meeting of the TB estimates subgroup: methods to use for WHO's definitive assessment of whether 2015 global TB targets are met. Geneva: WHO; 2015 (http://www.who.int/tb/advisory_bodies/impact_measurement_taskforce/meetings/consultation_april_2015_tb_estimates_subgroup/en/, accessed 8 August 2018).

^e World Health Organization Global Task Force on TB Impact Measurement. Report of the sixth meeting of the full Task Force; 19–21 April 2016, Glion-sur-Montreux, Switzerland. Geneva: WHO; 2016 (http://www.who.int/tb/advisory_bodies/impact_measurement_taskforce/meetings/tf6_report.pdf?ua=1, accessed 8 August 2018).

^f http://www.who.int/tb/areas-of-work/monitoring-evaluation/impact_measurement_taskforce/en/

^g http://www.who.int/tb/publications/factsheet_tb_impactmeasurement.pdf?ua=1

notification data to provide a direct measure of TB incidence.¹ By August 2018, a total of 68 countries, including 26 of the 30 high TB burden countries (listed in [Table 3.1](#)) had completed the checklist, often in association with a national TB epidemiological review ([Fig. 3.1](#)).

Methods currently used by WHO to estimate TB incidence can be grouped into four major categories, as follows ([Fig. 3.2](#)):

- **Results from TB prevalence surveys.** Incidence is estimated using prevalence survey results and estimates of the duration of disease, with the latter derived from a model that accounts for the impact of HIV coinfection and antiretroviral therapy (ART) on the distribution of disease duration.² This method is used for 22 countries, of which 22 have national survey data and one – India – has a survey in one state. The 23 countries accounted for 60% of the estimated global number of incident cases in 2017.
- **Notifications adjusted by a standard factor to account for underreporting, over-diagnosis and under-diagnosis.** This method is used for 144 countries that are all high-income countries except the Netherlands and the United Kingdom, plus selected middle-income countries with low levels of underreporting, including Brazil, China and the Russian Federation. For three countries (France, Republic of Korea and Turkey) the

adjustment was country specific, based on results from studies of underreporting. These 144 countries accounted for 15% of the estimated global number of incident cases in 2017.

- **Results from national inventory studies that measured the level of underreporting of detected TB cases combined with capture–recapture analysis.** This method is used for six countries: Egypt, Indonesia (yielding best estimates that are lower but statistically consistent with those previously derived from the 2013–2014 national TB prevalence survey),³ Iraq, the Netherlands, the United Kingdom and Yemen. These countries accounted for 9% of the estimated global number of incident cases in 2017.⁴
- **Case notification data combined with expert opinion about case-detection gaps.** Expert opinion, elicited through regional workshops or country missions, is used to estimate levels of underreporting, over-diagnosis and under-diagnosis. Trends are estimated through mortality data, surveys of the annual risk of infection or exponential interpolation using estimates of case-detection gaps for 3 years. In this report, this

¹ World Health Organization. Standards and benchmarks for tuberculosis surveillance and vital registration systems: checklist and user guide. Geneva: WHO; 2014 (<http://www.who.int/tb/publications/standardsandbenchmarks/en/>, accessed 25 July 2018). One of the standards is that levels of underreporting of detected TB cases should be minimal.

² Estimation of prevalence from incidence is not straightforward. For example, it requires assumptions about the duration of disease for different categories of case, and since prevalence surveys focus on bacteriologically confirmed TB in adults, adjustments to include children and extrapulmonary TB are needed.

³ The national inventory study implemented in Indonesia in 2017 is the largest of its kind to date. Further details are provided in [Box 3.2](#) and in [Chapter 4, Box 4.4](#).

⁴ Capture–recapture modelling is possible if six assumptions are met: (i) all cases should be observable; (ii) the proportion of mismatches and matching failures in record-linkage is low, which typically requires a large sampling fraction; (iii) a closed population during the study period (typically 3–6 months); (iv) if S represents the number of case lists or data sources available, then at least three data sources should be available ($S \geq 3$) and their dependencies must be accounted for in the model design, while the full S-way interaction between sources is assumed null; (v) homogeneity of within-source observation probabilities across subpopulation groups, such as those defined by socioeconomic and demographic characteristics; (vi) consistent case definitions across data sources. Few high TB burden countries are expected to be able to implement inventory studies that will meet these 6 assumptions to a sufficient degree.

BOX 3.2

Updates to estimates of TB disease burden in this report and anticipated updates

Updates in this report

Estimates of TB incidence and mortality in this report cover the period 2000–2017. Estimates of incidence and mortality for drug-resistant TB are for 2017. The main country-specific updates in this report are for estimates of TB incidence in Indonesia and South Africa.

1. Indonesia

Indonesia conducted a national inventory study to measure the underreporting of detected TB cases in the first quarter of 2017. Capture–recapture modelling based on three separate lists was then used to update incidence estimates, using methods set out in WHO guidance on inventory studies.^a The updated incidence estimates are consistent with those previously derived from the results of the national TB prevalence survey implemented in 2013–2014, but the best estimates are lower (about 15%) and also more precise (i.e. with narrower uncertainty intervals). Details are provided in **Box 4.4** in **Chapter 4**.

Estimates of TB mortality in Indonesia are based on those published by the Institute of Health Metrics and Evaluation (IHME), University of Washington, USA (see below).

2. South Africa

Trends in TB incidence in South Africa have been revised for two major reasons. First, there is now a clear, consistent and sustained downward trend in TB case notifications, which is also evident in other countries of southern Africa and which can be explained by the high coverage of ART among people living with HIV (for further details, see **Box 3.4**). Second, the values of the ratio of notified to estimated incident cases based on previously published trends, following further declines in notifications, are now implausible. For recent years, the revised estimates of TB incidence are lower than those previously published (e.g. the estimate for 2016 in this report is 21% lower than the estimate for 2016 published last year).

TB incidence will be reassessed when the results from the national TB prevalence survey 2018–2019 become available.

Estimates of TB mortality in South Africa are based on those published by IHME (see next section).

3. Newly reported data and updated estimates from other agencies

New VR data were reported to WHO between mid-2017 and mid-2018. Several countries reported historical data that were previously missing, or made corrections to previously reported data. In total, 1949 country-year data points were retained for analysis. Updated estimates of HIV prevalence and mortality were obtained from the Joint United Nations Programme on HIV/AIDS (UNAIDS) in July 2018.^b (In most instances, any resulting changes to TB burden estimates were well within the uncertainty intervals of previously published estimates, and trends were generally consistent.

In 19 countries (shown in **Fig. 3.10**), estimates of TB mortality (HIV-negative) were based on estimates published by IHME.^c These estimates use data from national and sample VR systems, and verbal autopsy surveys. Estimates of TB mortality in South Africa are adjusted by IHME for miscoding of deaths caused by HIV and TB. IHME estimates used in this report were slightly adjusted from those published by IHME to fit WHO estimates of the total number of deaths (mortality envelope). The median country-year envelope ratio (WHO/IHME) was 1.01 (interquartile range: 0.92–1.09) among 319 data points.

4. Findings from national TB epidemiological reviews

Small adjustments to incidence trajectories were made for Belarus, Malawi, Thailand and Turkmenistan, based on findings from recent national TB epidemiological reviews.

5. Drug-resistant TB

The estimated incidence of rifampicin-resistant TB in 2017 is based on the following formula:

$$I_{rr} = I[(1 - f)p_n((1 - r) + rp) + fp_r]$$

where I is overall TB incidence, I_{rr} is the incidence of rifampicin resistance, f is the cumulative risk for incident cases to receive a non-relapse retreatment (following failure or return after default), r is the proportion of relapses, p is the relative risk ratio in relapses compared with first episodes of TB, and p_n and p_r denote the proportion of rifampicin-resistant cases among previously untreated and previously treated patients, respectively.

Routine surveillance data for 2017 on levels of drug resistance were reported by 91 countries with continuous surveillance systems (i.e. routine diagnostic testing for drug resistance). New data from national surveys became available from seven countries since the publication of the 2017 global TB report: Eritrea, Eswatini, Indonesia, Lao People's Democratic Republic, Sri Lanka, Togo and the United Republic of Tanzania.

Updates anticipated in the near future

Updates to estimates of disease burden are expected towards the end of 2018 or in 2019 for Eswatini, Mozambique, Myanmar, Namibia, Nepal, South Africa and Viet Nam, following the completion of national TB prevalence surveys. Estimates of TB burden in India will be updated once results from a national TB prevalence survey planned for 2018–2019 become available.

^a World Health Organization. Assessing tuberculosis under-reporting through inventory studies. WHO, 2012. <http://apps.who.int/iris/handle/10665/78073>, accessed 15 August 2018).

^b <http://aidsinfo.unaids.org>

^c Downloaded from <http://ghdx.healthdata.org/gbd-results-tool>, June 2018.

BOX 3.3

WHO estimates for TB disease burden in the context of other estimates

The latest global estimates published by WHO and IHME are similar. For example, the best estimate for TB incidence in 2016 in this report is 10.1 million (range, 9.0–11.3 million), whereas the most recent best estimate from IHME is 10.4 million.^a The best estimate of the number of TB deaths among HIV-negative people in 2016 in this report is 1.3 million (range, 1.2–1.4 million), whereas the best estimate from IHME is 1.2 million. There is general consistency in mortality estimates in countries with VR systems and standard coding of causes of deaths of good quality, and in incidence estimates in countries with strong health-care and notification systems. Discrepancies are most apparent for countries where the underlying data are weaker, due to differences in the indirect estimation methods that are used.

When annual updates for TB are published by both WHO and IHME, entire time series (starting in 2000 for WHO, and 1990 for IHME) are updated. New information or refinements to methods used to produce estimates can result in changes to the estimates published for earlier years in previous publications. This is a normal part of disease burden estimation, and also occurs with disease burden estimates published for other diseases, such as HIV and malaria. For example, global estimates for 2015 for HIV, malaria and TB published by WHO, UNAIDS

and IHME in consecutive years (2015 and 2016) by the same agency have been within about 2–8% of each other. Global estimates of TB disease burden in 2015 published by WHO in this and the previous two global TB reports are within 4–5% of each other.

Country-specific estimates of TB disease burden published by WHO are generally consistent from year to year. In WHO reports published in 2014–2017, updates that have been apparent at global level have been due to updates for three countries: Nigeria (2014 report, following results from the country's first national TB prevalence survey in 2012); Indonesia (2015 report, following completion of a national TB prevalence survey in 2013–2014); and India (2016 report, following accumulating evidence from both survey and surveillance data).

As the availability and quality of data continue to improve, variability for the same year in consecutive reports will decrease and estimates published by WHO should converge with those published by other agencies. Ideally, estimates of TB incidence and mortality are based on national notification and vital registration systems that meet quality and coverage standards.

^a Downloaded from <http://ghdx.healthdata.org/gbd-results-tool>, July 2018.

method is used for 43 countries that accounted for 16% of the estimated global number of incident cases in 2017.

Of the four methods, the last one is the least preferred and it is relied upon only if one of the other three methods cannot be used. As explained in **Box 3.1**, the underlying principle for the WHO Global Task Force on TB Impact Measurement since its establishment in 2006 has been that estimates of the level of and trends in TB disease burden should be based on direct measurements from routine surveillance and surveys as much as possible, as opposed to indirect estimates that rely on modelling and expert opinion.

Further details about these methods are provided in the **online technical appendix**.¹

¹ The **online technical appendix** is available at www.who.int/tb/data.

3.1.2 Estimates of TB incidence in 2017

Globally in 2017, there were an estimated 10.0 million incident cases of TB (range, 9.0–11.1 million),² equivalent to 133 cases (range, 120–148) per 100 000 population. Estimates of absolute numbers are shown in **Table 3.2** and estimates of rates per capita are shown in **Table 3.3**.

Most of the estimated number of cases in 2017 occurred in the WHO South-East Asia Region (44%), the WHO African Region (25%) and the WHO Western Pacific Region (18%); smaller proportions of cases occurred in the WHO Eastern Mediterranean Region (7.7%), the WHO Region of the Americas (2.8%) and the WHO European Region (2.7%).

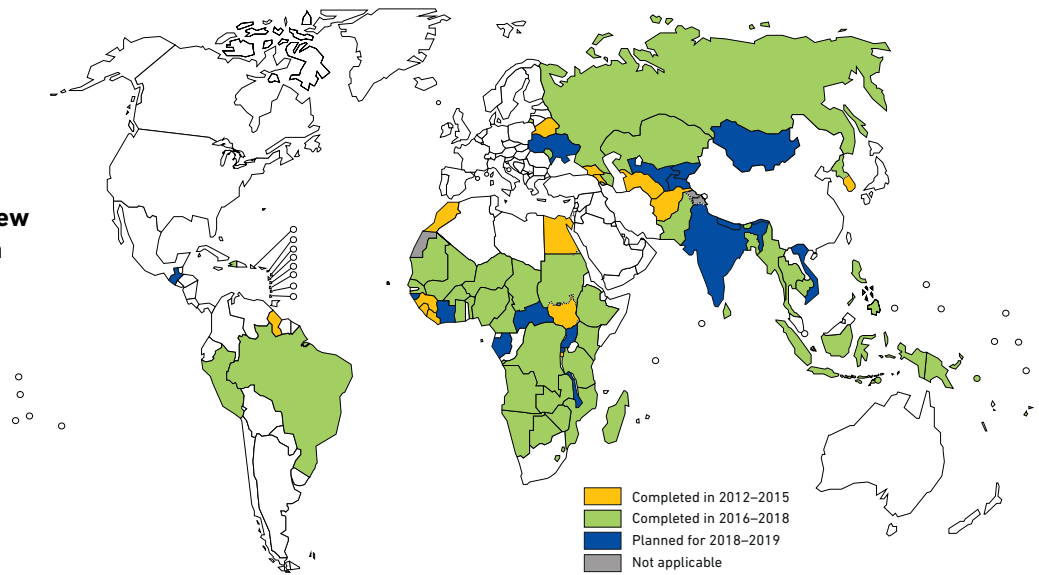
The 30 high TB burden countries³ accounted for 87% of all estimated incident cases worldwide, and eight of these countries accounted for two thirds of the global total: India (27%), China (9%), Indonesia (8%), the

² Here and elsewhere in the report, "range" refers to the 95% uncertainty interval.

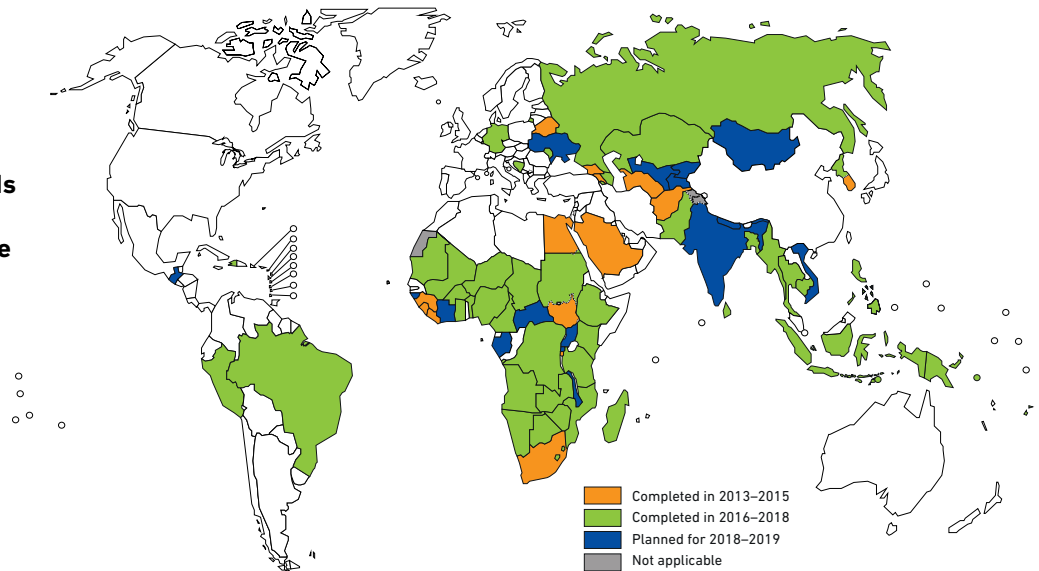
³ These countries are listed in **Table 3.2** and **Table 3.3**. For an explanation of how the list of 30 high TB burden countries was defined, see **Chapter 2**.

FIG. 3.1
Strengthening national TB surveillance (status in August 2018)

Countries in which a national TB epidemiological review has been undertaken since July 2012



Countries in which a checklist of standards and benchmarks has been completed since January 2013



Countries in which national inventory studies of the underreporting of detected TB cases have been implemented since 2000

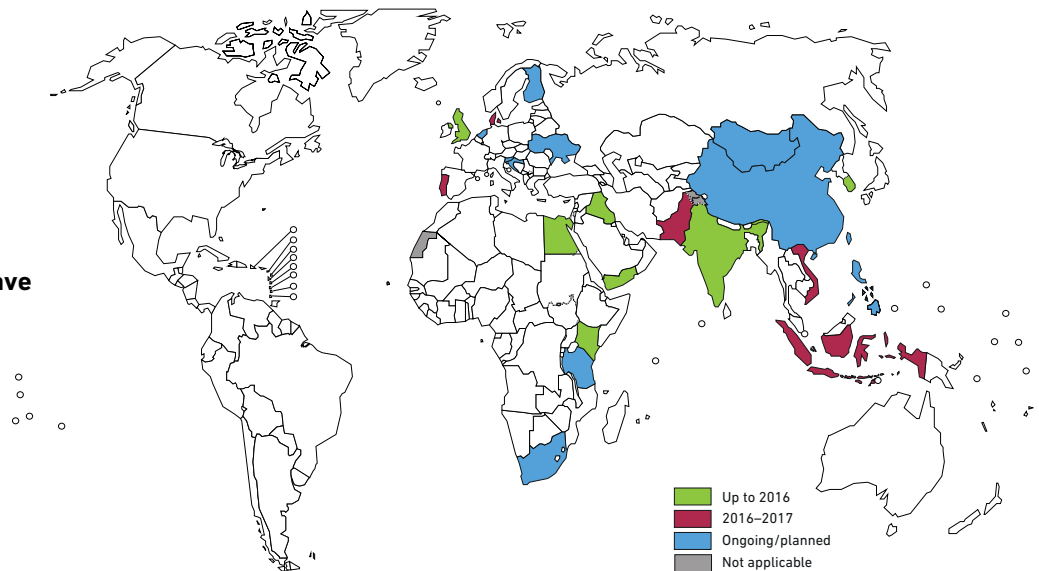


TABLE 3.2

Estimated epidemiological burden of TB in 2017 for 30 high TB burden countries, WHO regions and globally. Number in thousands.^a

	POPULATION	HIV-NEGATIVE TB MORTALITY		HIV-POSITIVE TB MORTALITY ^b		TOTAL TB INCIDENCE		HIV-POSITIVE TB INCIDENCE	
		BEST ESTIMATE	UNCERTAINTY INTERVAL	BEST ESTIMATE	UNCERTAINTY INTERVAL	BEST ESTIMATE	UNCERTAINTY INTERVAL	BEST ESTIMATE	UNCERTAINTY INTERVAL
Angola	30 000	20	12–31	7.8	3.9–13	107	69–153	18	9.1–30
Bangladesh	165 000	59	38–85	0.17	0.09–0.29	364	265–479	0.55	0.27–0.92
Brazil	209 000	5.1	4.8–5.3	1.9	1.4–2.5	91	78–105	11	9.3–13
Cambodia	16 000	3.1	2.0–4.3	0.41	0.27–0.57	52	36–72	1.3	0.89–1.8
Central African Republic	5 000	3.2	1.8–4.9	2.7	1.4–4.4	20	13–28	6.2	3.3–10
China	1 410 000	37	33–41	1.8	0.84–3.1	889	761–1 030	12	6.3–18
Congo	5 000	3.3	1.9–5.2	2.3	1.2–3.7	20	13–29	5.3	2.7–8.6
DPR Korea	25 000	16	11–22	0.04	0.02–0.07	131	114–149	0.17	0.09–0.28
DR Congo	81 000	49	29–74	7.5	3.5–13	262	169–374	20	13–29
Ethiopia	105 000	25	16–37	3.6	2.5–5.0	172	121–232	12	8.6–17
India ^c	1 340 000	410	381–441	11	6.5–16	2 740	1 870–3 770	86	57–120
Indonesia	264 000	107	100–114	9.4	5.0–15	842	767–919	36	20–57
Kenya	50 000	25	14–39	18	11–27	158	97–235	45	27–68
Lesotho	2 000	1.0	0.55–1.7	4.6	2.9–6.7	15	9.6–21	11	6.7–15
Liberia	5 000	2.7	1.6–4.1	0.91	0.57–1.3	15	9.4–21	2.2	1.4–3.2
Mozambique ^d	30 000	22	13–33	27	17–39	163	106–233	66	42–95
Myanmar ^d	53 000	27	18–39	4.9	3.5–6.6	191	141–249	17	12–22
Namibia ^d	3 000	0.75	0.48–1.1	0.80	0.55–1.1	11	8.2–14	3.9	2.5–5.5
Nigeria	191 000	120	70–183	35	21–52	418	273–594	58	37–85
Pakistan	197 000	54	42–67	2.2	1.1–3.8	525	373–704	7.3	3.6–12
Papua New Guinea	8 000	4.3	2.9–6.0	0.93	0.51–1.5	36	29–43	3.5	2.0–5.5
Philippines	105 000	26	23–31	0.38	<0.01–3.3	581	326–909	7.1	2.9–13
Russian Federation	144 000	10	9.4–12	1.7	0.85–2.8	86	56–123	18	12–26
Sierra Leone	8 000	3.0	1.8–4.5	0.78	0.49–1.1	23	15–33	2.8	1.8–4.0
South Africa ^d	57 000	22	20–24	56	39–77	322	230–428	193	137–258
Thailand	69 000	9.3	7.0–12	2.9	2.1–3.8	108	82–138	11	8.5–15
UR Tanzania	57 000	27	12–48	22	10–38	154	73–266	48	31–69
Viet Nam ^d	96 000	12	7.5–17	0.84	0.61–1.1	124	101–148	4.5	3.7–5.4
Zambia	17 000	5.0	2.9–7.7	13	8.2–19	62	40–88	36	23–52
Zimbabwe	17 000	2.0	1.3–2.9	6.3	4.5–8.5	37	27–47	23	15–33
High TB burden countries	4 760 000	1 110	1 030–1 190	247	214–282	8 720	7 680–9 810	766	680–857
Africa	1 050 000	413	348–485	252	219–287	2 480	2 210–2 760	663	585–747
The Americas	1 010 000	18	17–19	6.0	5.3–6.7	282	262–302	30	28–33
Eastern Mediterranean	682 000	89	75–104	3.0	1.8–4.5	771	611–950	9.8	6.0–15
Europe	920 000	24	23–25	5.0	3.8–6.4	273	236–313	33	26–42
South-East Asia	1 970 000	638	598–679	28	22–36	4 440	3 530–5 450	152	117–191
Western Pacific	1 900 000	92	85–100	5.0	3.8–6.4	1 800	1 490–2 130	31	24–40
GLOBAL	7 520 000	1 270	1 190–1 360	300	266–335	10 000	9 000–11 100	920	832–1 010

^a Numbers shown to two significant figures if under 100 and to three significant figures otherwise.

^b Deaths among HIV-positive TB cases are classified as HIV deaths according to ICD-10.

^c Estimates of TB incidence and mortality for India are interim, pending results from the national TB prevalence survey planned for 2019/2020.

^d Estimates of TB incidence and mortality for Mozambique, Myanmar, Namibia, South Africa and Viet Nam will be reviewed after final results from their respective national TB prevalence surveys are available in 2019.

TABLE 3.3

Estimated epidemiological burden of TB in 2017 for 30 high TB burden countries, WHO regions and globally. Rates per 100 000 population.

	HIV-NEGATIVE TB MORTALITY		HIV-POSITIVE TB MORTALITY ^a		TOTAL TB INCIDENCE		HIV PREVALENCE IN INCIDENT TB (%)	
	BEST ESTIMATE	UNCERTAINTY INTERVAL	BEST ESTIMATE	UNCERTAINTY INTERVAL	BEST ESTIMATE	UNCERTAINTY INTERVAL	BEST ESTIMATE	UNCERTAINTY INTERVAL
Angola	67	39–103	26	13–44	359	232–512	17	10–25
Bangladesh	36	23–52	0.11	0.05–0.18	221	161–291	0.15	0.08–0.24
Brazil	2.4	2.3–2.5	0.91	0.67–1.2	44	37–50	12	11–13
Cambodia	19	13–27	2.6	1.7–3.6	326	224–447	2.5	2.3–2.7
Central African Republic	68	38–105	58	31–94	423	274–604	32	21–43
China	2.6	2.4–2.9	0.13	0.06–0.22	63	54–73	1.3	0.73–2.0
Congo	63	36–98	43	22–71	376	239–545	27	17–37
DPR Korea	63	43–86	0.17	0.09–0.28	513	446–584	0.13	0.07–0.21
DR Congo	60	35–90	9.2	4.3–16	322	208–460	7.6	4.4–12
Ethiopia	24	15–35	3.5	2.4–4.8	164	115–221	7.2	6.6–7.8
India ^b	31	28–33	0.79	0.48–1.2	204	140–281	3.1	2.8–3.5
Indonesia	40	38–43	3.6	1.9–5.8	319	291–348	4.3	2.4–6.7
Kenya	50	28–78	37	22–55	319	195–472	29	26–32
Lesotho	46	25–75	206	128–302	665	430–949	71	63–78
Liberia	57	34–86	19	12–28	308	199–440	15	13–17
Mozambique ^c	73	43–111	90	56–131	551	356–787	40	36–44
Myanmar ^c	51	33–73	9.2	6.6–12	358	263–466	8.7	8.0–9.4
Namibia ^c	30	19–43	31	22–43	423	324–535	36	35–38
Nigeria	63	36–96	18	11–27	219	143–311	14	12–16
Pakistan	27	21–34	1.1	0.56–1.9	267	189–357	1.4	0.78–2.2
Papua New Guinea	53	36–73	11	6.2–18	432	352–521	9.9	5.9–15
Philippines	25	22–29	0.36	<0.01–3.1	554	311–866	1.2	0.69–1.9
Russian Federation	7.3	6.6–8.0	1.2	0.59–1.9	60	39–85	21	20–23
Sierra Leone	39	23–59	10	6.5–15	301	193–431	12	12–13
South Africa ^c	39	35–43	99	68–135	567	406–754	60	55–64
Thailand	13	10–17	4.2	3.1–5.6	156	119–199	11	9.7–11
UR Tanzania	47	21–83	39	18–67	269	127–464	31	28–35
Viet Nam ^c	12	7.8–17	0.88	0.64–1.2	129	106–155	3.6	3.4–3.9
Zambia	30	17–45	76	48–110	361	234–514	58	53–64
Zimbabwe	12	7.7–17	38	27–51	221	164–287	63	59–67
High TB burden countries	23	22–25	5.2	4.5–5.9	183	161–206	8.8	7.4–10
Africa	39	33–46	24	21–27	237	211–263	27	23–31
The Americas	1.8	1.7–1.9	0.60	0.53–0.67	28	26–30	11	9.6–12
Eastern Mediterranean	13	11–15	0.43	0.26–0.66	113	90–139	1.3	0.73–2.0
Europe	2.6	2.5–2.7	0.54	0.41–0.69	30	26–34	12	9.1–16
South-East Asia	32	30–35	1.4	1.1–1.8	226	179–277	3.5	2.4–4.7
Western Pacific	4.9	4.5–5.3	0.26	0.20–0.34	94	78–112	1.8	1.2–2.4
GLOBAL	17	16–18	4.0	3.5–4.5	133	120–148	9.2	7.9–11

^a Deaths among HIV-positive TB cases are classified as HIV deaths according to ICD-10.

^b Estimates of TB incidence and mortality for India are interim, pending results from the national TB prevalence survey planned for 2019/2020.

^c Estimates of TB incidence and mortality for Mozambique, Myanmar, Namibia, South Africa and Viet Nam will be reviewed after final results from their respective national TB prevalence surveys are available in 2019.

FIG. 3.2
Main methods used to estimate TB incidence

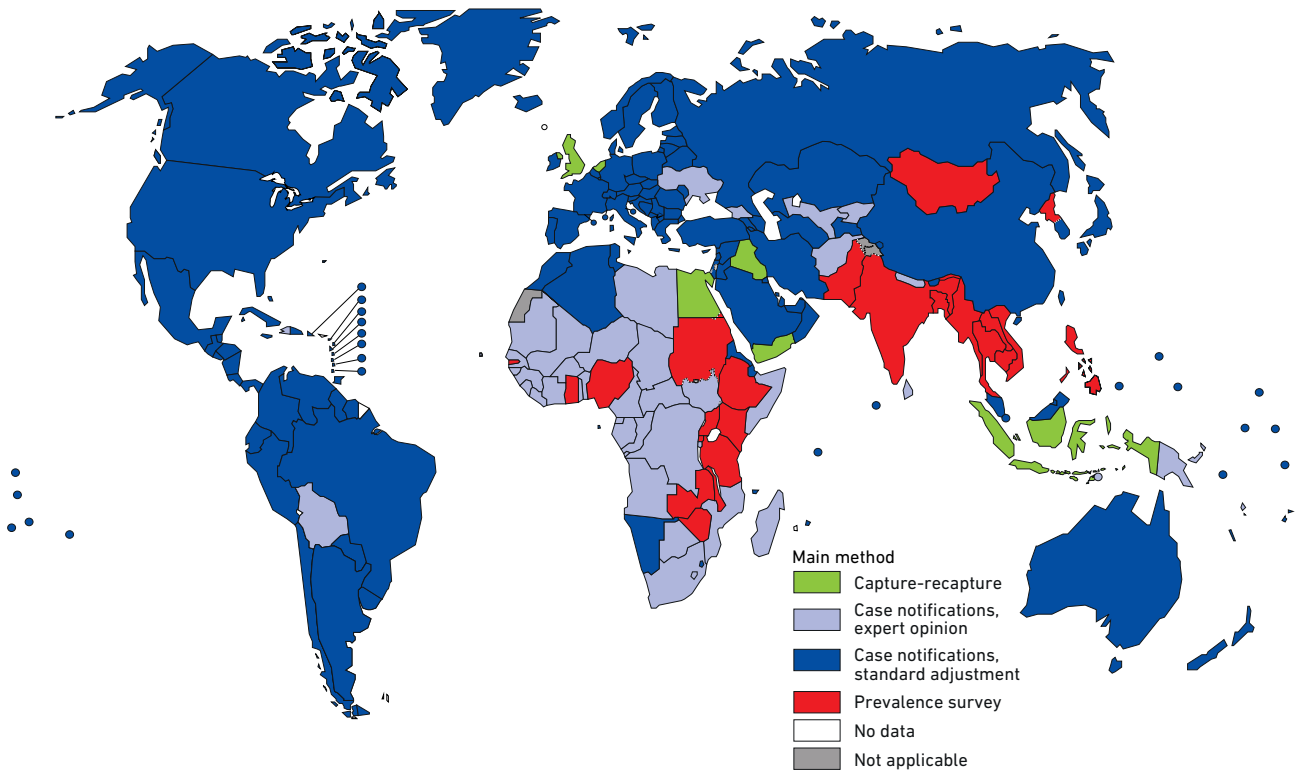


FIG. 3.3
Estimated TB incidence in 2017, for countries with at least 100 000 incident cases

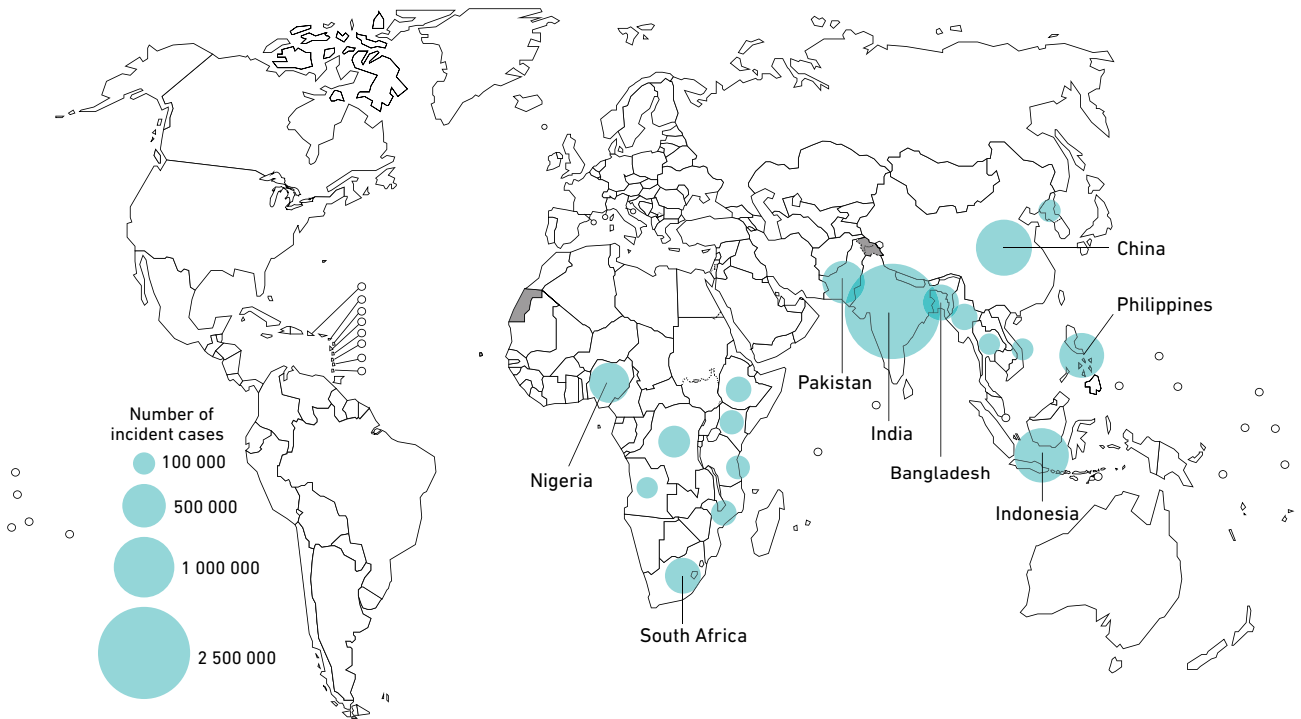
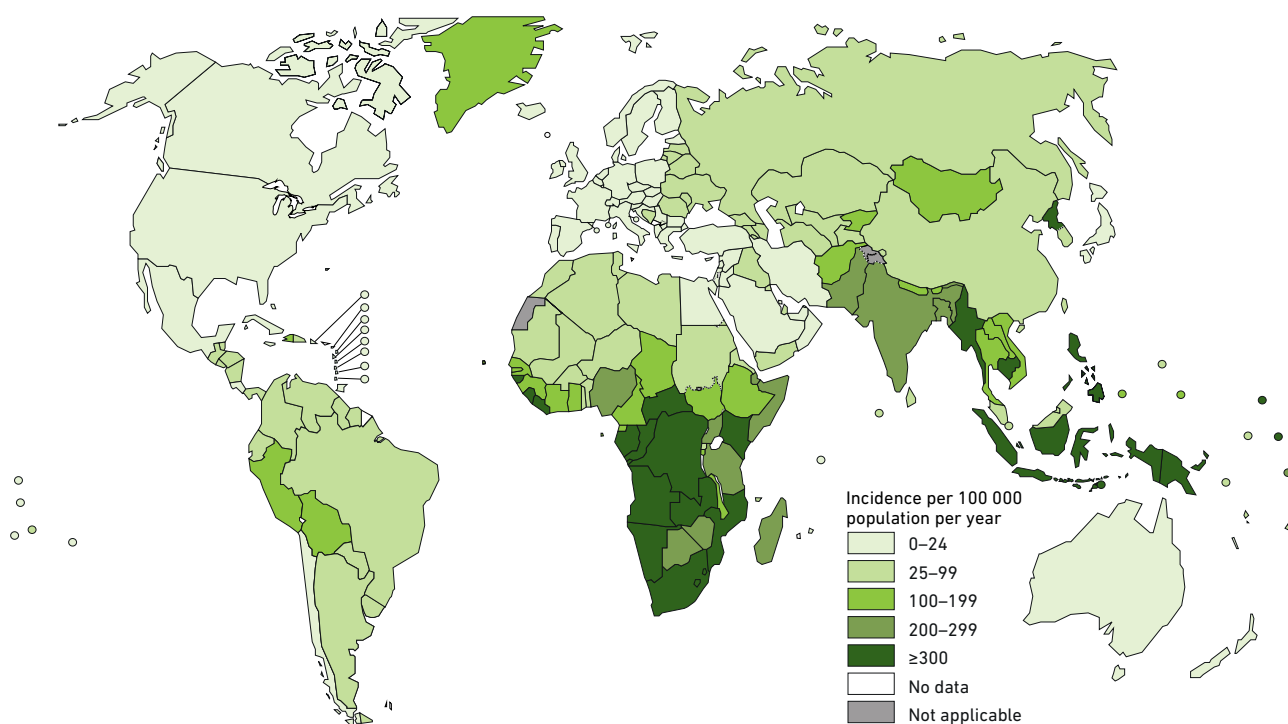


FIG. 3.4
Estimated TB incidence rates, 2017



Philippines (6%), Pakistan (5%), Nigeria (4%), Bangladesh (4%) and South Africa (3%) (Fig. 3.3).

The severity of national TB epidemics in terms of the annual number of incident TB cases relative to population size (the incidence rate) varied widely among countries in 2017. There were under 10 incident cases per 100 000 population in most high-income countries, 150–400 in most of the 30 high TB burden countries (Fig. 3.4), and above 500 in a few countries including the Democratic People’s Republic of Korea, Lesotho, Mozambique, the Philippines and South Africa (Table 3.3).

An estimated 9% (range, 7.9–11%) of the incident TB cases in 2017 were among people living with HIV (Table 3.2, Table 3.3). The proportion of TB cases coinfecting with HIV was highest in countries in the WHO African Region, exceeding 50% in parts of southern Africa (Fig. 3.5). The risk of developing TB in the 37 million people living with HIV was 20 times higher than the risk in the rest of the world population (range, 17–23), increasing with a decreasing prevalence of HIV in the general population.

Estimates of the incidence of zoonotic TB are shown in Table 3.4.

3.1.3 Estimated trends in TB incidence, 2000–2017

Consistent with previous global TB reports, the number of incident cases is falling slowly, in both absolute terms and per capita (Fig. 3.6, Fig. 3.7). Globally, the average rate of decline in the TB incidence rate was 1.5% per year

in the period 2000–2017, and 1.8% between 2016 and 2017. This needs to accelerate to 4–5% per year by 2020 and to 10% per year by 2025, to achieve the milestones for reductions in cases and deaths set in the End TB Strategy (Chapter 2).

Trends are shown for the six WHO regions in Fig. 3.8 and for the 30 high TB burden countries in Fig. 3.9.¹ The fastest declines in the 5-year period 2013–2017 were in the WHO European Region (on average, 5% per year). In the same period, particularly impressive reductions (4–8% per year) occurred in southern Africa (e.g. Eswatini [formerly Swaziland], Lesotho, Namibia, South Africa, Zambia and Zimbabwe) following a peak in the HIV epidemic and the expansion of TB and HIV prevention and care (Box 3.4), and in the Russian Federation (5% per year) following intensified efforts to reduce the burden of TB and scrutiny of progress from the highest political levels.

3.2 TB mortality

Deaths from TB among HIV-negative people are classified as TB deaths in the most recent version of the International classification of diseases (ICD-10).² When

¹ Time series of estimates for all countries are available online. Annex 1 explains how to access and download them.

² World Health Organization. International statistical classification of diseases and health related problems, ICD-10. Geneva: WHO; 2016 (<http://apps.who.int/classifications/icd10/browse/2016/en>, accessed 31 August 2018).

FIG. 3.5
Estimated HIV prevalence in new and relapse TB cases, 2017

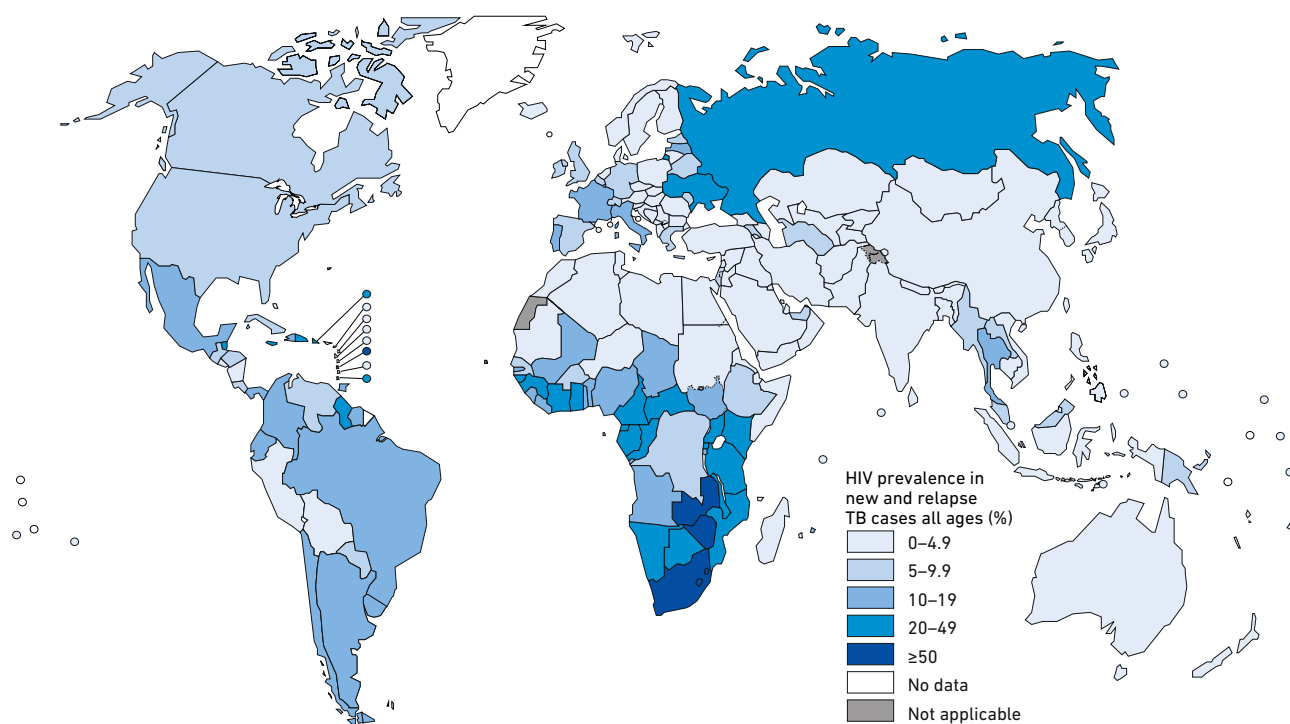


TABLE 3.4

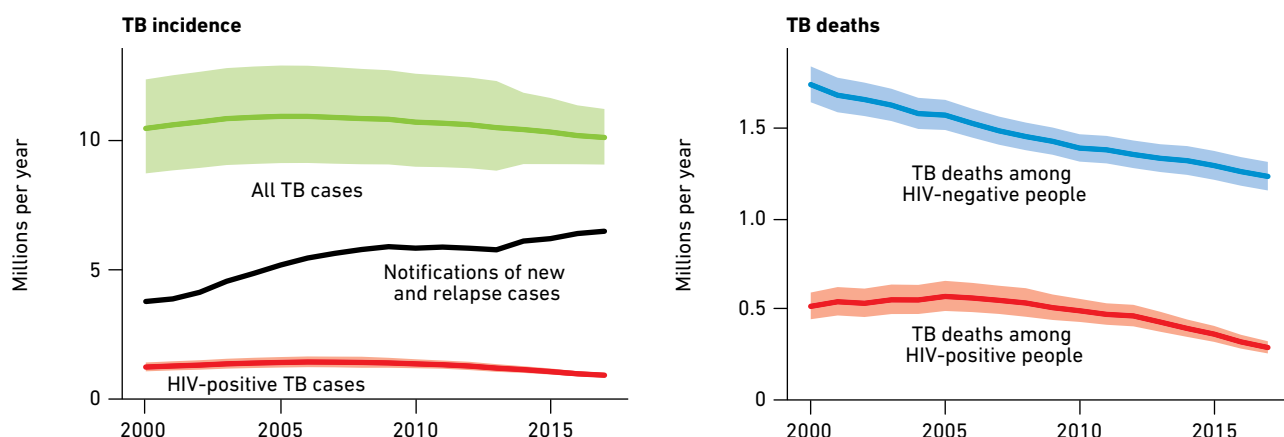
Estimated incidence and mortality due to *M. bovis* TB. Best estimates (absolute numbers) are followed by the lower and upper bounds of the 95% uncertainty interval.^a

WHO REGION	INCIDENT CASES		DEATHS	
	BEST ESTIMATE	UNCERTAINTY INTERVAL	BEST ESTIMATE	UNCERTAINTY INTERVAL
Africa	70 000	18 800–154 000	9 270	2 450–20 500
The Americas	821	222–1 800	45	12–98
Eastern Mediterranean	7 660	1 980–17 100	733	194–1 620
Europe	1 150	308–2 550	87	24–191
South-East Asia	44 900	11 500–100 000	2 090	568–4 590
Western Pacific	18 000	4 740–40 000	309	84–678
GLOBAL	142 000	70 600–239 000	12 500	4 910–23 700

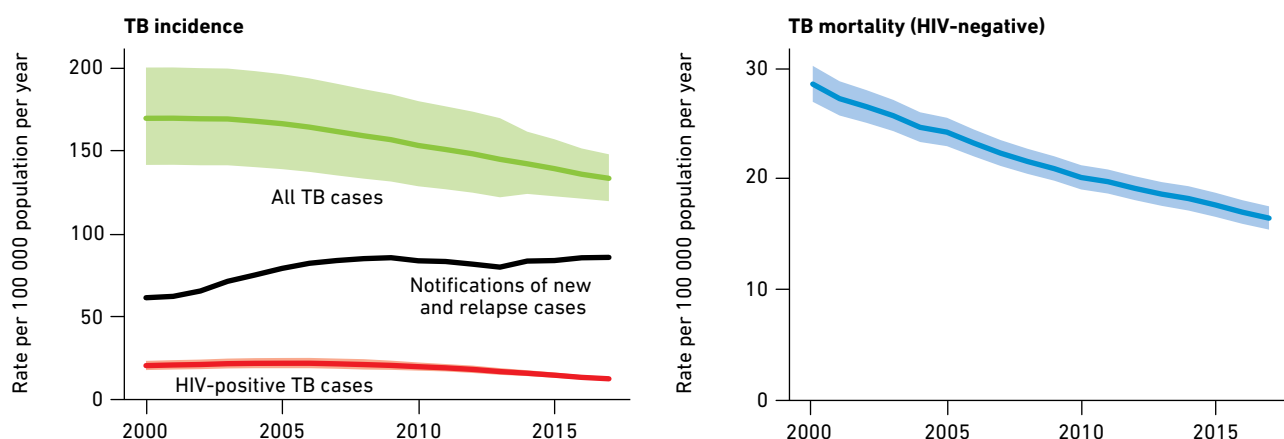
^a Numbers shown to two significant figures if under 100 and to three significant figures otherwise.

FIG. 3.6

Global trends in the estimated number of incident TB cases and the number of TB deaths (in millions), 2000–2017. Shaded areas represent uncertainty intervals.

**FIG. 3.7**

Global trends in estimated TB incidence and mortality rates, 2000–2017. Shaded areas represent uncertainty intervals.



an HIV-positive person dies from TB, the underlying cause is classified as HIV. For consistency with these international classifications, this section makes a clear distinction between TB deaths in HIV-negative people and TB deaths in HIV-positive people.

3.2.1 Methods to estimate TB mortality

TB mortality among HIV-negative people can be measured directly using data from national vital registration (VR) systems, provided that these systems have high coverage and that causes of death are accurately determined and coded according to ICD-10. Sample VR systems covering representative areas of the country (the approach used, for example, in China) provide an interim solution. Mortality surveys can also be used to estimate deaths caused by TB. In 2017, most countries with a high burden of TB lacked national or sample VR systems, and few had conducted mortality surveys. In the absence of VR systems or mortality surveys, TB mortality can be

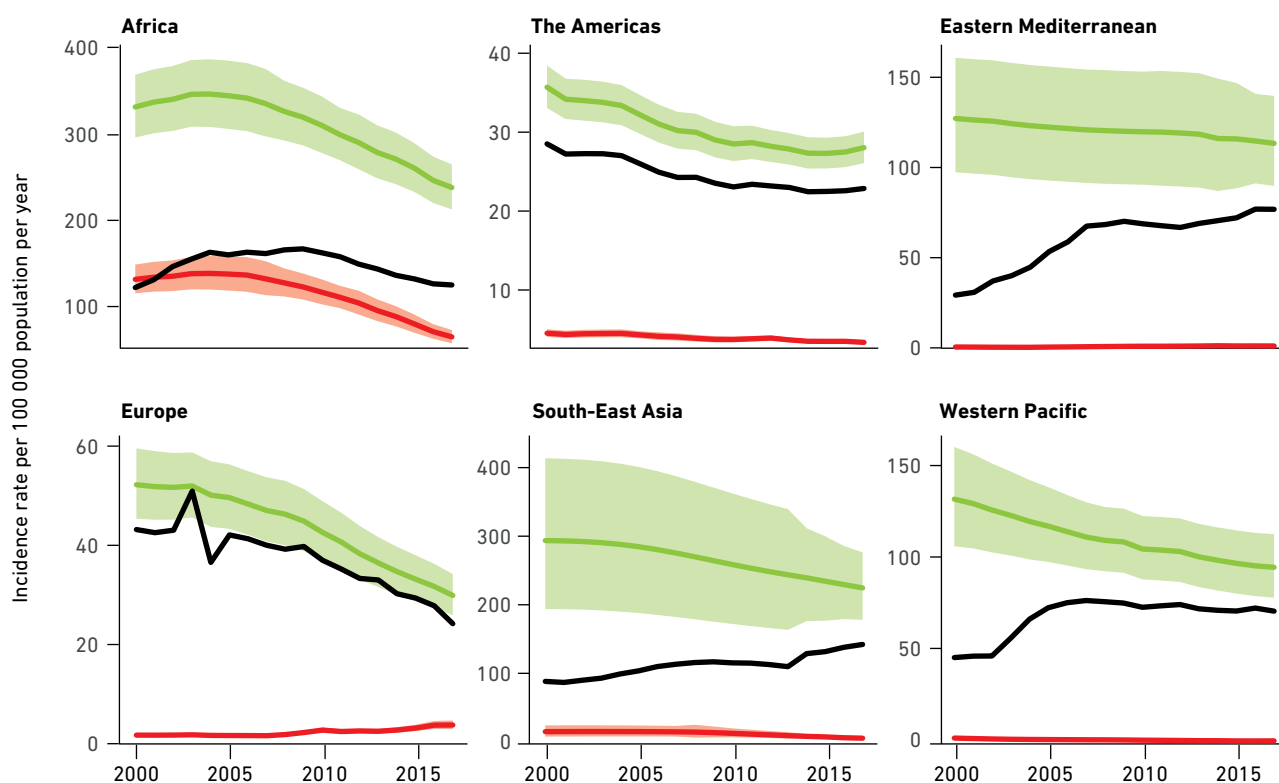
estimated as the product of TB incidence and the case fatality ratio (CFR), or through ecological modelling using mortality data from countries with VR systems.

TB mortality among HIV-positive people is hard to measure, even when VR systems are in place, because deaths among HIV-positive people are coded as HIV deaths, and contributory causes (e.g. TB) are often not reliably assessed and recorded. TB deaths among HIV-positive people were estimated as the product of TB incidence and the CFR, with the latter accounting for the protective effect of ART.

Until 2008, WHO estimates of TB mortality used VR or mortality survey data for only three countries. This was substantially improved to 89 countries in 2009, although most of the data were from countries in the WHO European Region and the WHO Region of the Americas, which accounted for less than 10% of the world's TB cases. For the current report, VR or mortality survey data were used for 127 countries (Fig. 3.10), which collectively

FIG. 3.8

Regional trends in estimated TB incidence rates by WHO region, 2000–2017. Total TB incidence rates are shown in green and incidence rates of HIV-positive TB are shown in red. Shaded areas represent uncertainty intervals. The black lines show notifications of new and relapse cases for comparison with estimates of the total incidence rate.



accounted for 57% of the estimated number of TB deaths (among HIV-negative people) globally in 2017. For 19 countries, analyses of VR data and resulting estimates of TB deaths published by the Institute of Health Metrics and Evaluation (IHME) at the University of Washington, USA, were used.¹ The WHO African Region has the greatest need to introduce or strengthen VR systems in which causes of death are classified according to ICD-10.

Details about the methods used to produce estimates of TB mortality are provided in the [online technical appendix](#).²

3.2.2 Estimates of TB mortality in 2017

Estimates of the absolute number of deaths caused by TB are shown globally, for the six WHO regions and for the 30 high TB burden countries, in [Table 3.2](#). There were an estimated 1.3 million (range, 1.2–1.4 million) deaths from TB among HIV-negative people in 2017 and an additional 300 000 (range, 266 000–335 000) deaths from TB among HIV-positive people.

TB is the tenth leading cause of death worldwide, and since 2011 it has been the leading cause of death from a single infectious agent, ranking above HIV/AIDS

([Fig. 3.11](#), [Fig. 3.12](#) and [Fig. 3.13](#)).³ Most of these deaths could be prevented with early diagnosis and appropriate treatment ([Chapter 1](#)). For example, among people whose TB was detected, reported and treated in 2016, the treatment success rate was 82% globally ([Chapter 4](#)); and in high-income countries with universal health coverage, the proportion of people who die from TB can be under 5% ([Section 3.2.4](#)).

About 82% of TB deaths among HIV-negative people occurred in the WHO African Region and the WHO South-East Asia Region in 2017; these regions accounted for 85% of the combined total of TB deaths in HIV-negative and HIV-positive people. India accounted for 32% of global TB deaths among HIV-negative people, and for 27% of the combined total TB deaths in HIV-negative and HIV-positive people.

Estimates of TB mortality rates (per 100 000 population) are shown globally, for the six WHO regions and for the 30 high TB burden countries, in [Table 3.3](#). Globally, the number of TB deaths among HIV-negative people per 100 000 population was 17 (range, 16–18) in 2017, and 21 (range, 20–22) when TB deaths among HIV-positive people were included. There was considerable variation

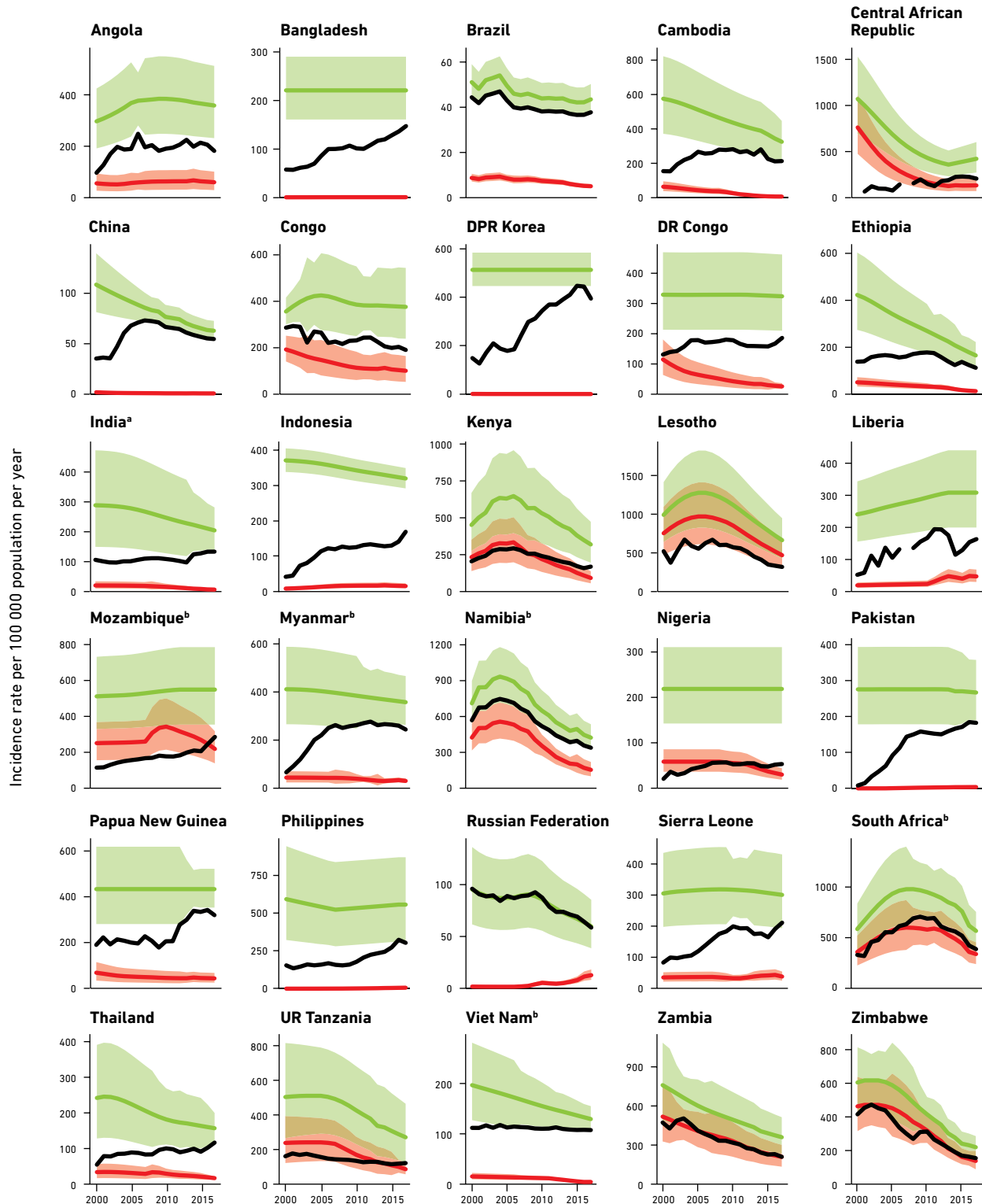
¹ Downloaded from <http://ghdx.healthdata.org/gbd-results-tool>, July 2018.

² The [online technical appendix](#) is available at www.who.int/tb/data.

³ Global Health Estimates 2016: Deaths by cause, age, sex, by country and by region, 2000–2016. Geneva, World Health Organization; 2018.

FIG. 3.9

Trends in estimated TB incidence in the 30 high TB burden countries, 2000–2017. TB incidence rates are shown in **green** and incidence rates of HIV-positive TB are shown in **red**. Shaded areas represent uncertainty intervals. The **black** lines show notifications of new and relapse cases for comparison with estimates of the total incidence rate.



^a Estimates of TB incidence for India are interim, pending results from the national TB prevalence survey planned for 2019/2020.

^b Estimates of TB incidence for Mozambique, Myanmar, Namibia, South Africa and Viet Nam will be reviewed after final results from their respective national TB prevalence surveys are available in 2019.

BOX 3.4

Rapid decline in TB incidence in six countries in southern Africa

In six countries in southern Africa, TB incidence is estimated to have fallen rapidly in the period 2010–2017, with average annual rates of decline of 18% in Eswatini, 10% in Zimbabwe, 8% in Botswana, 7% in Lesotho and South Africa, and 6% in Namibia. Such national rates of decline in TB incidence are among the fastest of recent decades. TB case notifications in all six countries have also declined in this period, at nearly the same average rates.^a

Since the 1990s, TB incidence in southern Africa has been strongly driven by the HIV epidemic. In 2017, the estimated proportion of incident TB cases coinfecting with HIV ranged from 36% in Namibia to 71% in Lesotho. The prevalence of HIV in the general population has remained relatively stable in all six countries since 2010 (ranging from 8% in Namibia to 17–19% in Eswatini). What has changed is that the coverage of HIV care in the general population and among new TB cases in people living with HIV has improved substantially.

In the general population, ART coverage among people living with HIV rapidly increased in the period 2010–2017, with improvements ranging from 24% to 61% in South Africa and from 30% to 84% in Zimbabwe (Fig. B3.4.1). In four of the six countries, ART coverage was above 80% of estimated prevalent HIV cases in 2017, a remarkable achievement. Trends in ART coverage among new TB cases coinfecting with HIV followed similar trends and reached similar levels of coverage by 2017, with the exception of Botswana (34%).

The HIV pandemic has had a major impact on the health of people in southern Africa. HIV drove the annual national incidence of TB in the six featured countries to some of the highest rates ever globally (e.g. to over 1% of the population per year in Eswatini and Lesotho, and 0.9% of the population per year in South Africa in 2010). In 2010, the estimated TB incidence rate ratio (TB incidence among HIV-positive individuals divided by TB incidence in HIV-negative individuals) ranged from 10 in Botswana to 32 in Zimbabwe. Those relative risks nearly halved within 8 years, from 4.7 in Botswana to 18 in Zimbabwe in 2017.

The rapid increase in ART coverage among people living with HIV in all six countries led to a tremendous reduction in TB incidence rates over the past 8 years, confirming predictions made in 2010.^b Further reductions will require addressing other important determinants of TB (Chapter 7) and improving the coverage of preventive treatment (Chapter 5).

^a Further details on trends in case notifications are provided in Chapter 4 and in country profiles.

^b Williams BG, Granich R, De Cock KM, Glaziou P, Sharma A, Dye C. Antiretroviral therapy for tuberculosis control in nine African countries. *PNAS* 2010; 107(45):19485–9 (<https://www.ncbi.nlm.nih.gov/pubmed/20974976>, accessed 31 August 2018).

FIG. B3.4.1

Trends in TB incidence rates expressed as a percentage of the best estimate for 2010 (in red), overall ART coverage among people living with HIV (solid black line) and ART coverage among new TB cases coinfecting with HIV (dashed line), in 6 selected countries in southern Africa, 2010–2017.

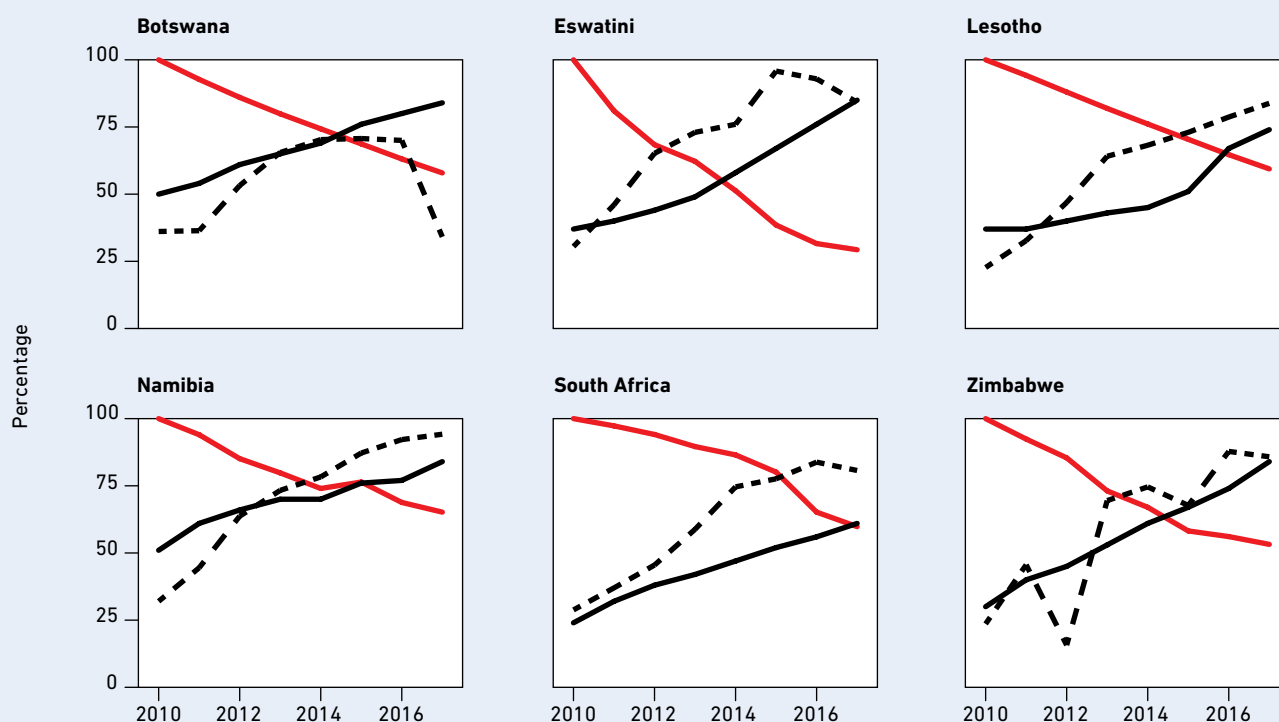
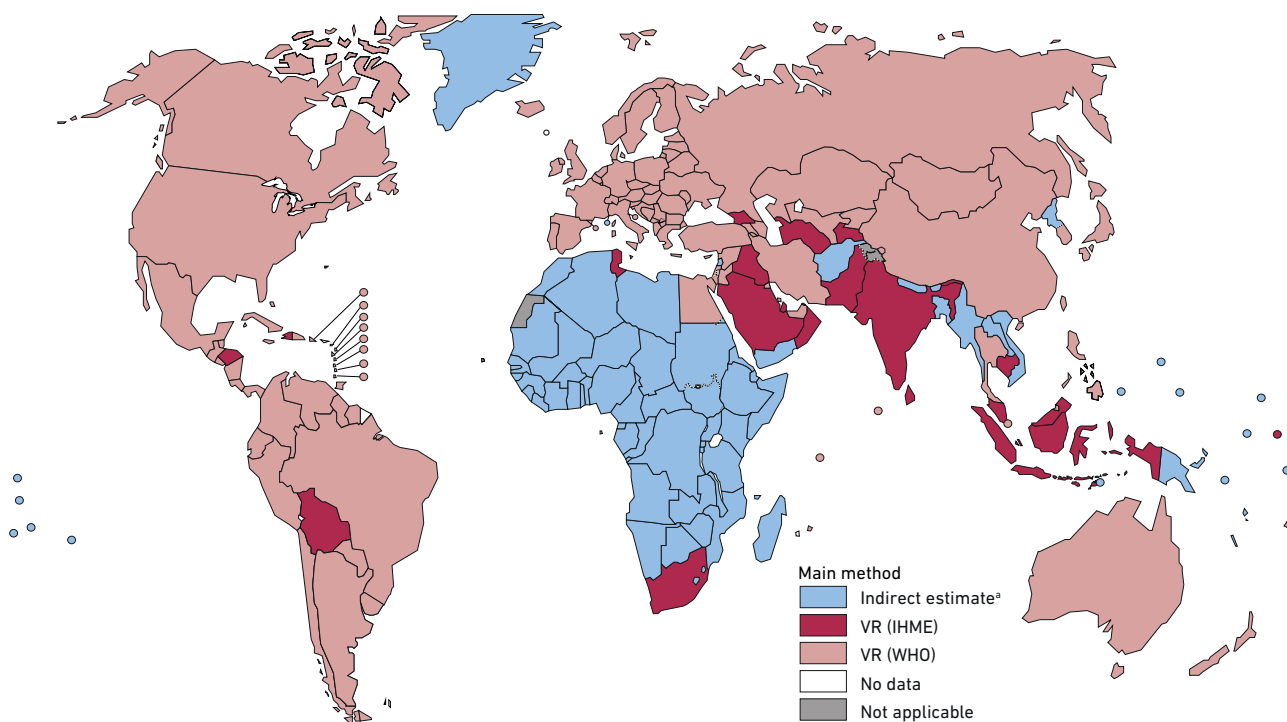


FIG. 3.10

Main methods used to estimate TB mortality in HIV-negative people

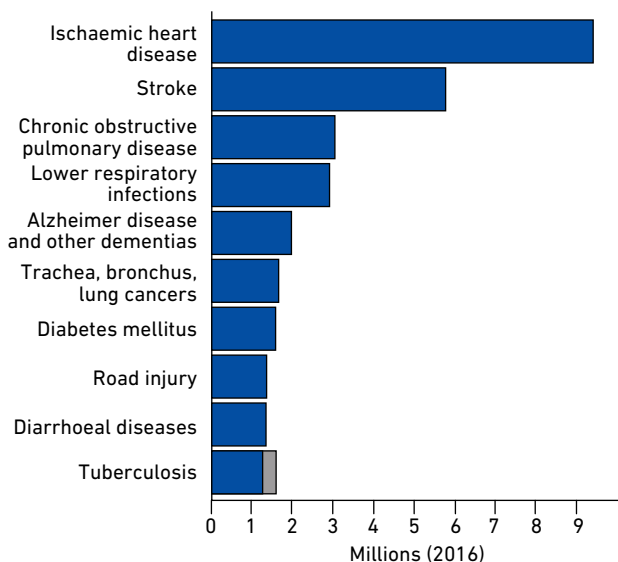


^a Mortality is estimated as the product of TB incidence and the TB case fatality ratio. Further details are provided in the [online technical appendix](#).

FIG. 3.11

Top causes of death worldwide in 2016.^{a,b}

Deaths from TB among HIV-positive people are shown in grey.



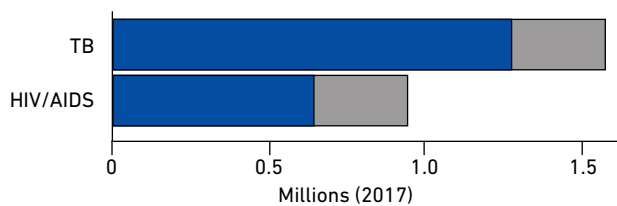
^a This is the latest year for which estimates for all causes are currently available. See WHO estimates, available at <http://apps.who.int/gho/portal/uhc-fp-cabinet-wrapper-v2.jsp?id=1020201> (accessed 15 August 2018).

^b Deaths from TB among HIV-positive people are officially classified as deaths caused by HIV/AIDS in the International classification of diseases.

FIG. 3.12

Estimated number of deaths from HIV/AIDS and TB in 2017.

Deaths from TB among HIV-positive people are shown in grey.^{a,b}

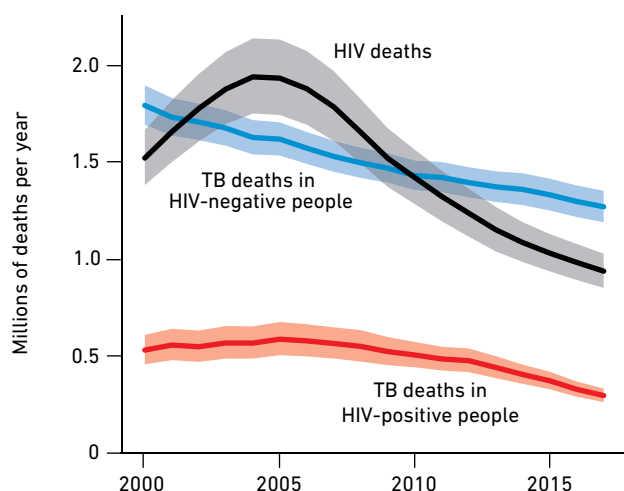


^a For HIV/AIDS, the latest estimates of the number of deaths in 2017 that have been published by UNAIDS are available at <http://www.unaids.org/en/resources/publications/all> (accessed 15 August 2018). For TB, the estimates for 2017 are those published in this report.

^b Deaths from TB among HIV-positive people are officially classified as deaths caused by HIV/AIDS in the International classification of diseases.

FIG. 3.13**Global trends in the estimated number of deaths caused by TB and HIV (in millions), 2000–2017.^{a,b}**

Shaded areas represent uncertainty intervals.



^a For HIV/AIDS, the latest estimates of the number of deaths in 2017 that have been published by UNAIDS are available at <http://www.unaids.org/en/resources/publications/all> (accessed 15 August 2018). For TB, the estimates for 2017 are those published in this report.

^b Deaths from TB among HIV-positive people are officially classified as deaths caused by HIV/AIDS in the International classification of diseases.

among countries (Fig. 3.14), ranging from less than one TB death per 100 000 population in many high-income countries, to 40 or more deaths per 100 000 population in much of the WHO African Region and in four high TB burden countries in Asia (the Democratic People's Republic of Korea, Indonesia, Myanmar and Papua New Guinea).

Estimates of the number of deaths caused by zoonotic TB are shown in Table 3.4.

3.2.3 Estimated trends in TB mortality, 2000–2017

Globally, the absolute number of deaths caused by TB among HIV-negative people has been falling since 2000, from a best estimate of 1.8 million in 2000 to 1.3 million in 2017 (Fig. 3.6), a reduction of 29%. The reduction since 2015 (the baseline year for milestones and targets set in the End TB Strategy, as shown in Table 3.1) was 5%. The TB mortality rate (TB deaths among HIV-negative people per 100 000 population per year) fell by 42% globally between 2000 and 2017 (Fig. 3.7), and by 3.2% between 2016 and 2017.

Mortality rates among HIV-negative people have been falling in all six WHO regions since 2000, but the rate of decline varies (Fig. 3.15). For example, in the 5-year period 2013–2017, the fastest average rates of decline were in the WHO European Region (11% per year) and the WHO South-East Asia Region (4.3% per year), and the slowest rate was in the WHO African Region (1.7% per year).

Trends also vary markedly among the 30 high TB burden countries (Fig. 3.16), ranging from substantial reductions (more than 50%) since 2000 (e.g. in Cambodia,

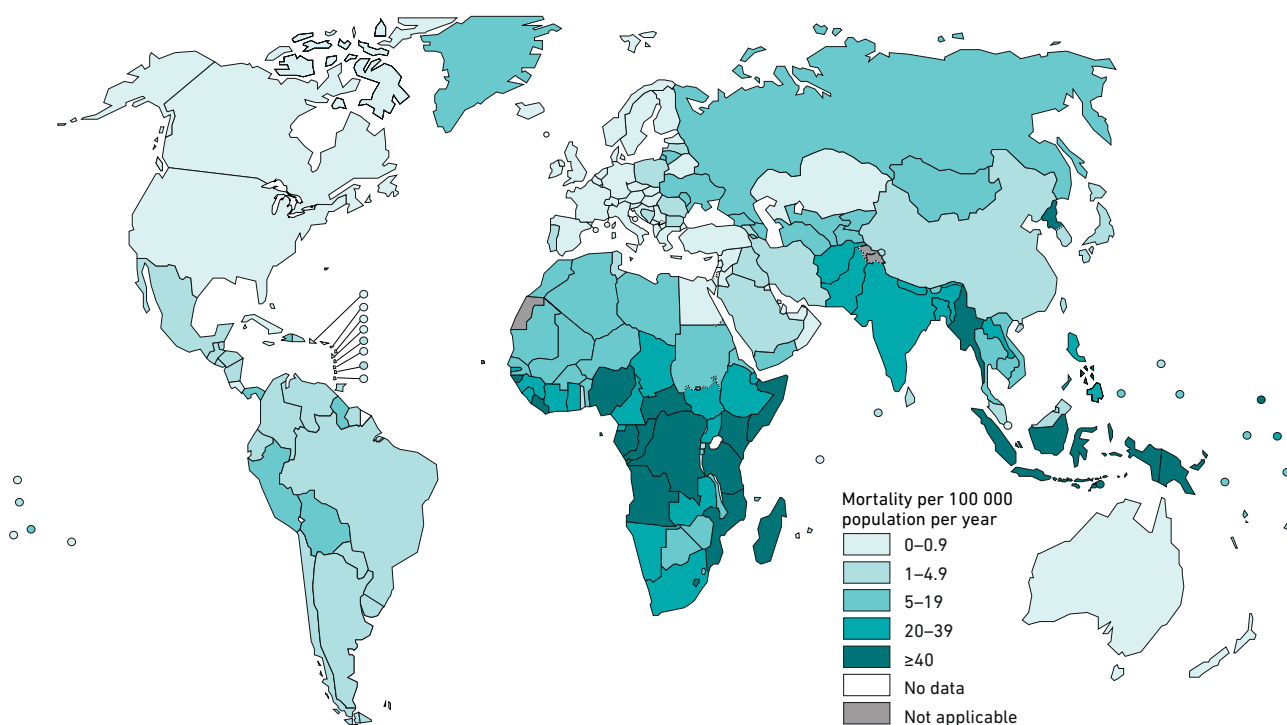
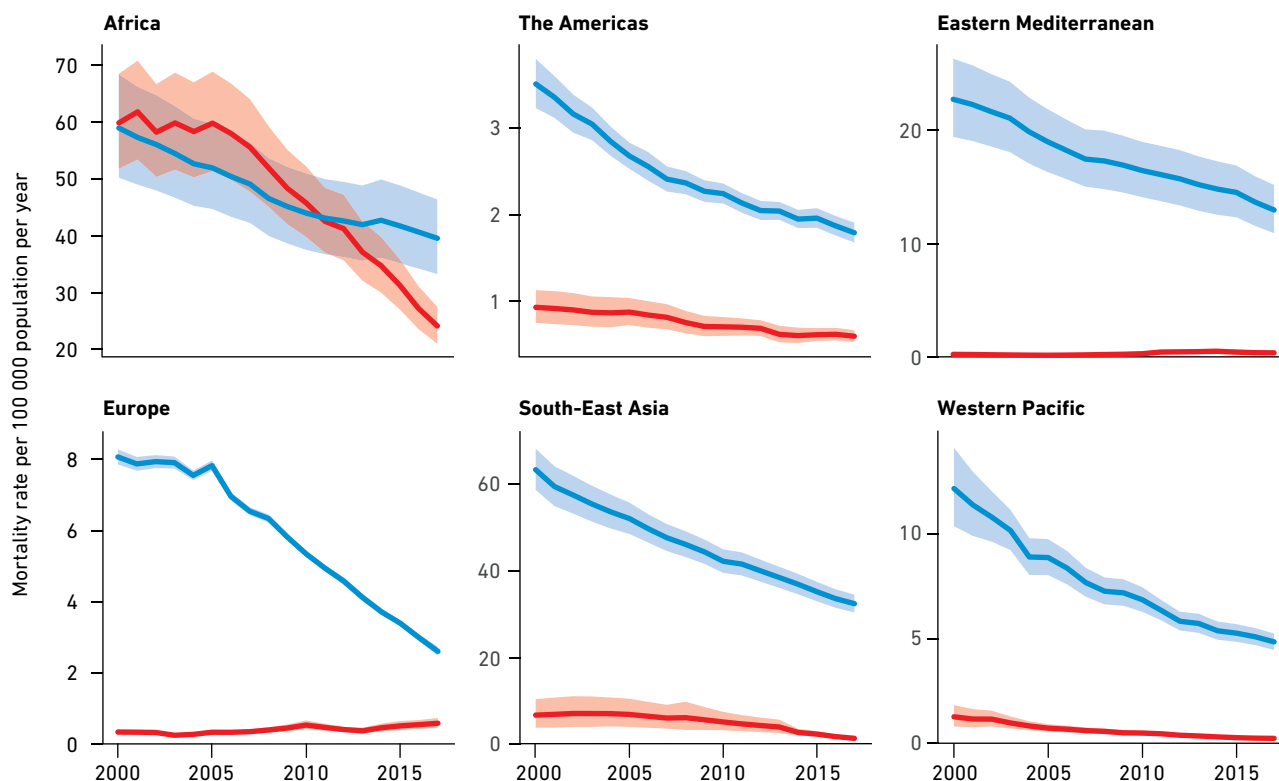
FIG. 3.14**Estimated TB mortality rates excluding TB deaths among HIV-positive people, 2017**

FIG. 3.15

Regional trends in estimated TB mortality rates by WHO region, 2000–2017. Estimated TB mortality rates excluding TB deaths among HIV-positive people are shown in **blue** and estimated mortality rates of HIV-positive TB are shown in **red**. Shaded areas represent uncertainty intervals.



China, Ethiopia, Myanmar, the Russian Federation and Viet Nam) to limited changes (e.g. in Angola and Congo).¹ High TB burden countries with rates of decline in TB deaths among HIV-negative people that exceeded 6% per year in the 5-year period 2013–2017 included the Russian Federation (13% per year), Ethiopia (12% per year), Sierra Leone (10% per year), Kenya (8% per year) and Viet Nam (8% per year).

Globally, the number of TB deaths among HIV-positive people has fallen by 44% since 2000, from 534 000 (range, 460 000–613 000) in 2000 to 300 000 (range, 266 000–335 000) in 2017, and by 20% since 2015. Most of this reduction was in the WHO African Region (Fig. 3.15). In several high TB burden countries, the number of deaths caused by TB among HIV-positive people has fallen substantially in recent years; for example, in Cambodia, Kenya, Namibia, South Africa, the United Republic of Tanzania and Zimbabwe (Fig. 3.16).

3.2.4 The case fatality ratio and across-country equity

The CFR is the proportion of people with TB who die from the disease; it can be approximated as the number of TB

deaths divided by the number of new cases in the same year. The CFR allows assessment of variation in equity in terms of access to TB diagnosis and treatment among countries (because if everyone with TB had access to timely diagnosis and high-quality treatment, the CFR would be low in all countries). To achieve the milestones for reductions in TB deaths set for 2020 and 2025 in the End TB Strategy, the global CFR needs to fall to 10% by 2020 and to 6.5% by 2025 (Chapter 2).

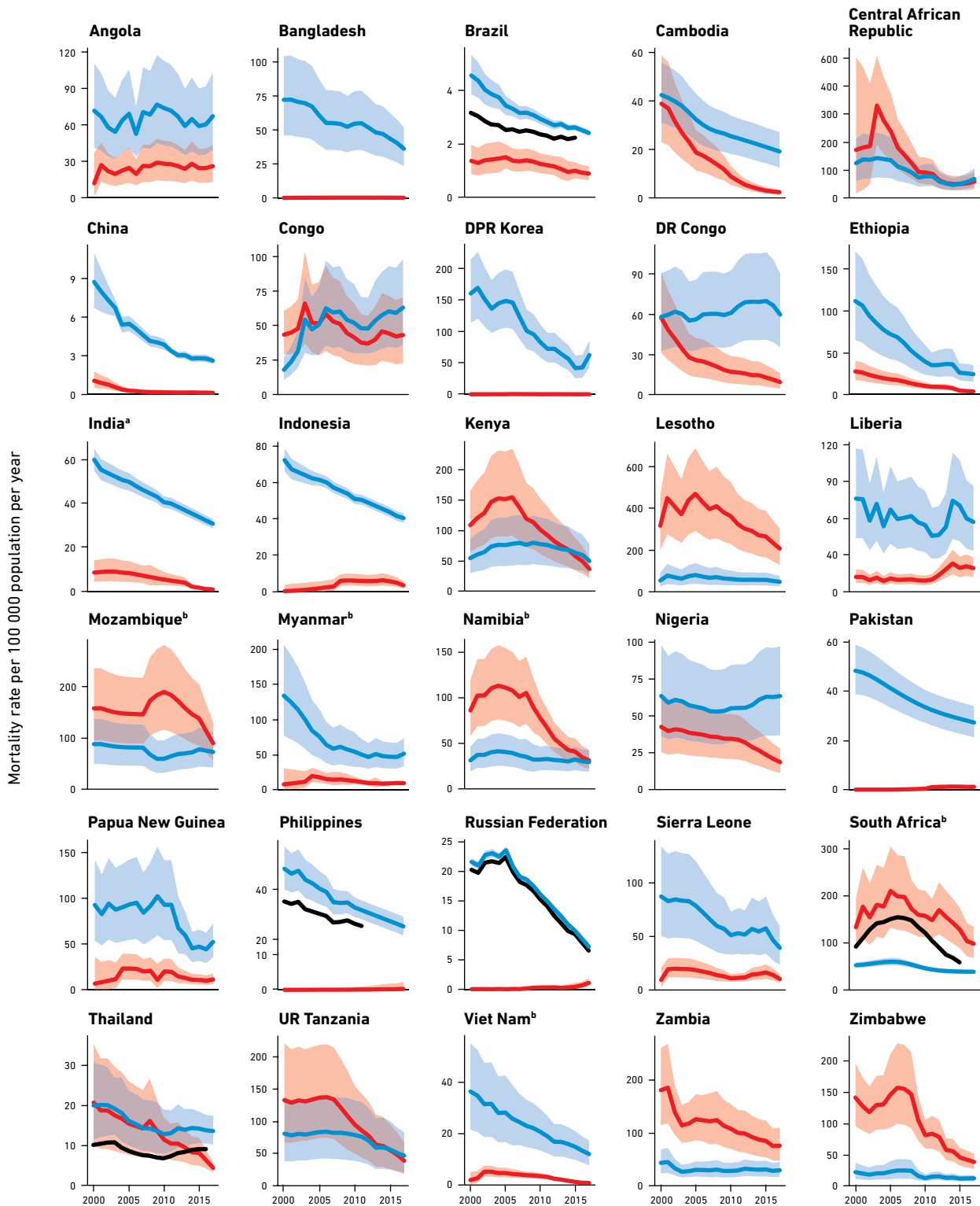
In 2017, the global CFR (calculated as the combined number of TB deaths in HIV-negative people and HIV-positive people, divided by the total number of incident cases in both HIV-negative and HIV-positive people)² was 16%, down from 23% in 2000. It varied widely among countries (Fig. 3.17), from under 5% in a few countries to more than 20% in most countries in the WHO African Region. Intensified efforts are required to reduce the CFR to 10% globally by 2020.

¹ Time series of estimates for all countries are available online. Annex 1 explains how to access and download them.

² The CFR was calculated based on the combined total of deaths in HIV-negative and HIV-positive people for the purpose of cross-country comparisons; in particular, to illustrate the high CFRs in African countries, which could be reduced by effective detection and care programmes. CFRs restricted to HIV-negative TB deaths and cases can also be calculated but are not shown here. At the subnational level, CFRs can also be restricted to HIV-negative TB deaths, depending on the country and its HIV burden.

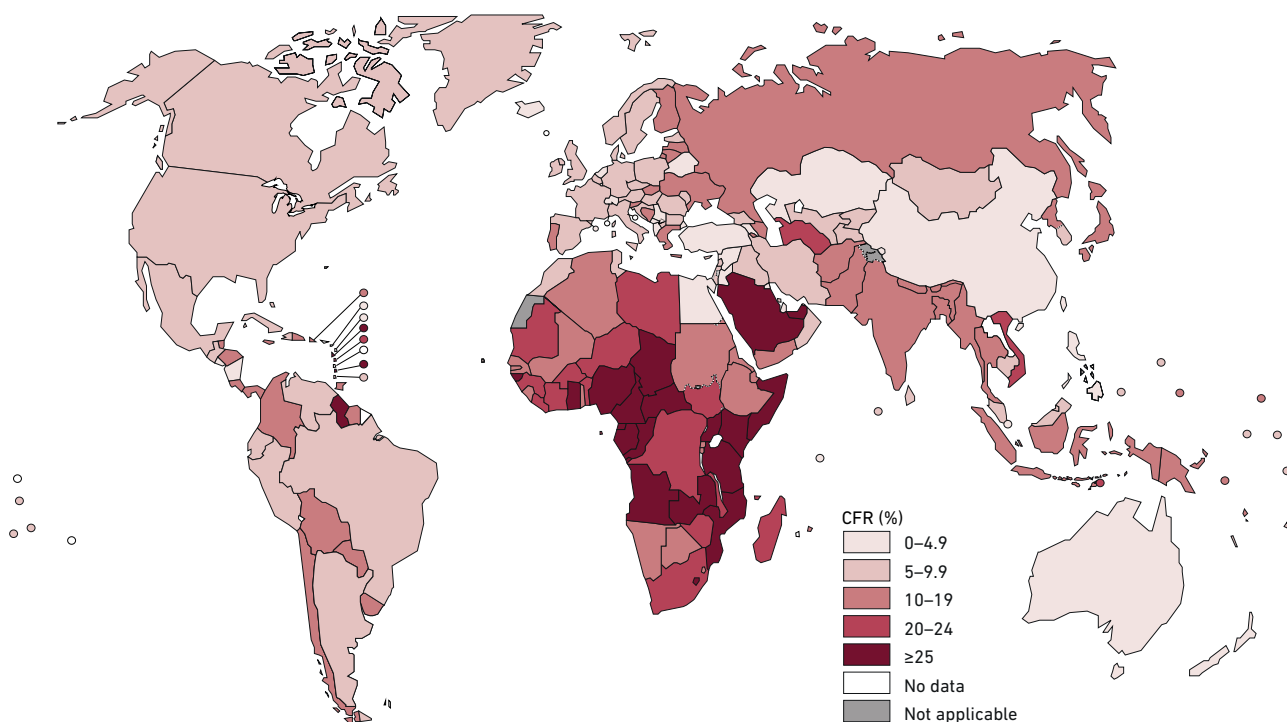
FIG. 3.16

Trends in estimated TB mortality rates in the 30 high TB burden countries, 2000–2017. TB mortality rates in HIV-negative people are shown in blue and mortality rates of HIV-positive TB are shown in red. The black lines show observations from vital registration systems. Shaded areas represent uncertainty intervals.



^a Estimates of TB mortality for India are interim, pending results from the national TB prevalence survey planned for 2019/2020.

^b Estimates of TB mortality for Mozambique, Myanmar, Namibia, South Africa and Viet Nam will be reviewed after final results from their respective national TB prevalence surveys are available in 2019.

FIG. 3.17**Estimates of the case fatality ratio (CFR), including HIV-negative and HIV-positive people, 2017**

3.2.5 Estimated number of deaths averted by TB treatment, 2000–2017

The actual numbers of TB deaths (presented above) can be compared with the number of TB deaths that would have occurred in the absence of TB treatment, to estimate the number of deaths averted by TB interventions. The number of deaths that would have occurred each year in the absence of TB treatment (and without ART provided alongside TB treatment for HIV-positive cases) can be conservatively estimated as the number of estimated incident cases (Section 3.1) multiplied by the relevant estimated CFR for untreated TB.¹ Estimates are conservative because they do not account for the impact of TB services or the availability of ART on the level of TB incidence, or for the indirect, downstream impact of these interventions on future levels of infections, cases and deaths.

Between 2000 and 2017, TB treatment alone averted an estimated 45 million deaths among HIV-negative people (Table 3.5). Among HIV-positive people, TB treatment supported by ART averted an additional 9 million deaths.

3.3 Drug-resistant TB

Drug-resistant TB remains a major public health concern in many countries. Three major categories are used for

¹ Further details about methods used to estimate lives saved, including CFRs for different categories of TB case, are provided in the [online technical appendix](#), available at www.who.int/tb/data.

global surveillance and treatment. MDR-TB is TB that is resistant to both rifampicin and isoniazid, the two most powerful anti-TB drugs; it requires treatment with a second-line regimen. RR-TB also requires treatment with second-line drugs. With increasing use of Xpert® MTB/RIF for simultaneous detection of TB and resistance to rifampicin, a growing number of RR-TB cases (without further testing for isoniazid resistance) are being detected and notified (Chapter 4). Extensively drug-resistant TB (XDR-TB) is defined as MDR-TB plus resistance to at least one drug in both of the two most important classes of medicines in an MDR-TB regimen: fluoroquinolones and second-line injectable agents (amikacin, capreomycin or kanamycin). Estimates of the disease burden caused by drug-resistant TB presented in this chapter focus on MDR/RR-TB.

3.3.1 Global surveillance of anti-TB drug resistance

Since the launch of the Global Project on Anti-tuberculosis Drug Resistance Surveillance in 1994, data on drug resistance have been systematically collected and analysed from 160 countries worldwide (82% of the 194 WHO Member States), which collectively have more than 97% of the world's population and TB cases. This includes 91 countries that have continuous surveillance systems based on routine diagnostic drug susceptibility testing (DST) of *Mycobacterium tuberculosis* isolates

TABLE 3.5

Cumulative number of deaths averted by TB and TB/HIV interventions 2000–2017 (in millions), globally and by WHO region^a

WHO REGION	HIV-NEGATIVE PEOPLE		HIV-POSITIVE PEOPLE		TOTAL	
	BEST ESTIMATE	UNCERTAINTY INTERVAL	BEST ESTIMATE	UNCERTAINTY INTERVAL	BEST ESTIMATE	UNCERTAINTY INTERVAL
Africa	5.5	4.5–6.4	6.0	5.3–6.8	12	10–13
The Americas	1.5	1.4–1.6	0.34	0.32–0.37	1.8	1.7–2.0
Eastern Mediterranean	3.5	3.0–4.0	0.06	0.04–0.07	3.6	3.1–4.1
Europe	1.9	1.7–2.1	0.21	0.20–0.23	2.1	1.9–2.3
South-East Asia	20	16–24	1.9	1.5–2.4	22	18–25
Western Pacific	13	11–14	0.35	0.31–0.39	13	12–14
GLOBAL	45	39–50	8.9	7.9–9.9	54	48–59

^a Numbers shown to two significant figures.

FIG. 3.18

Data sources available to estimate levels of TB drug resistance

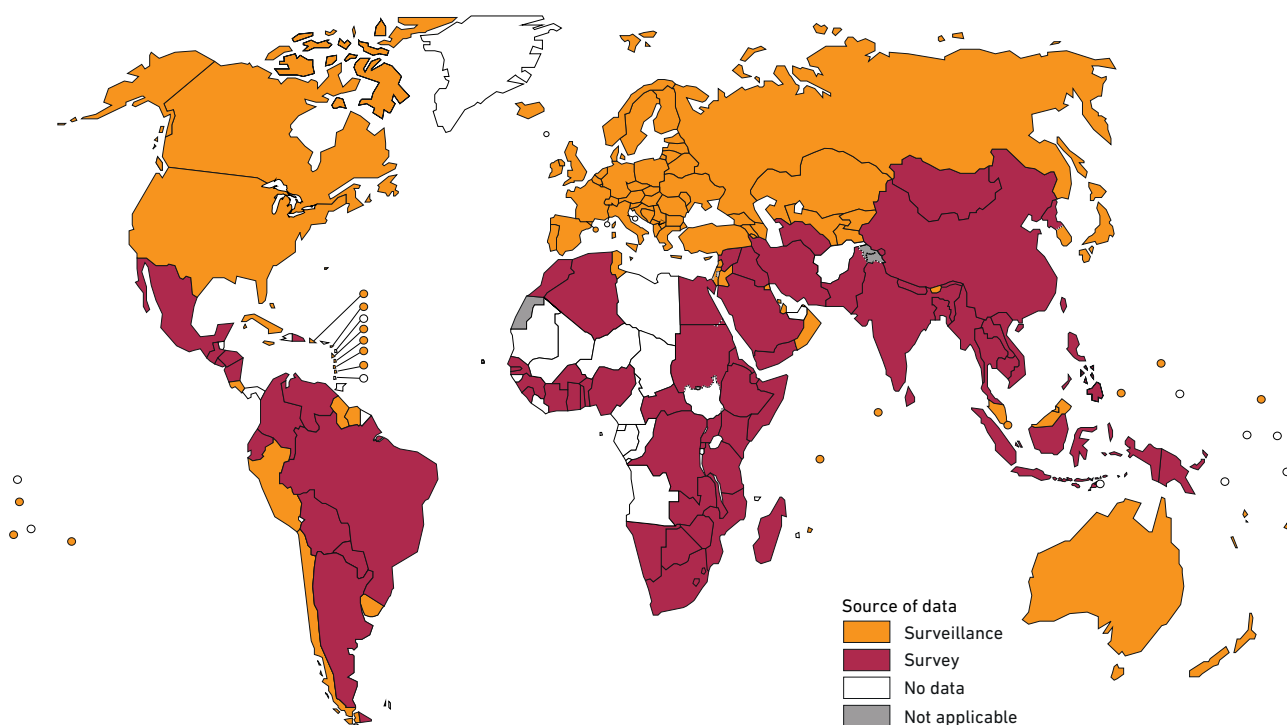
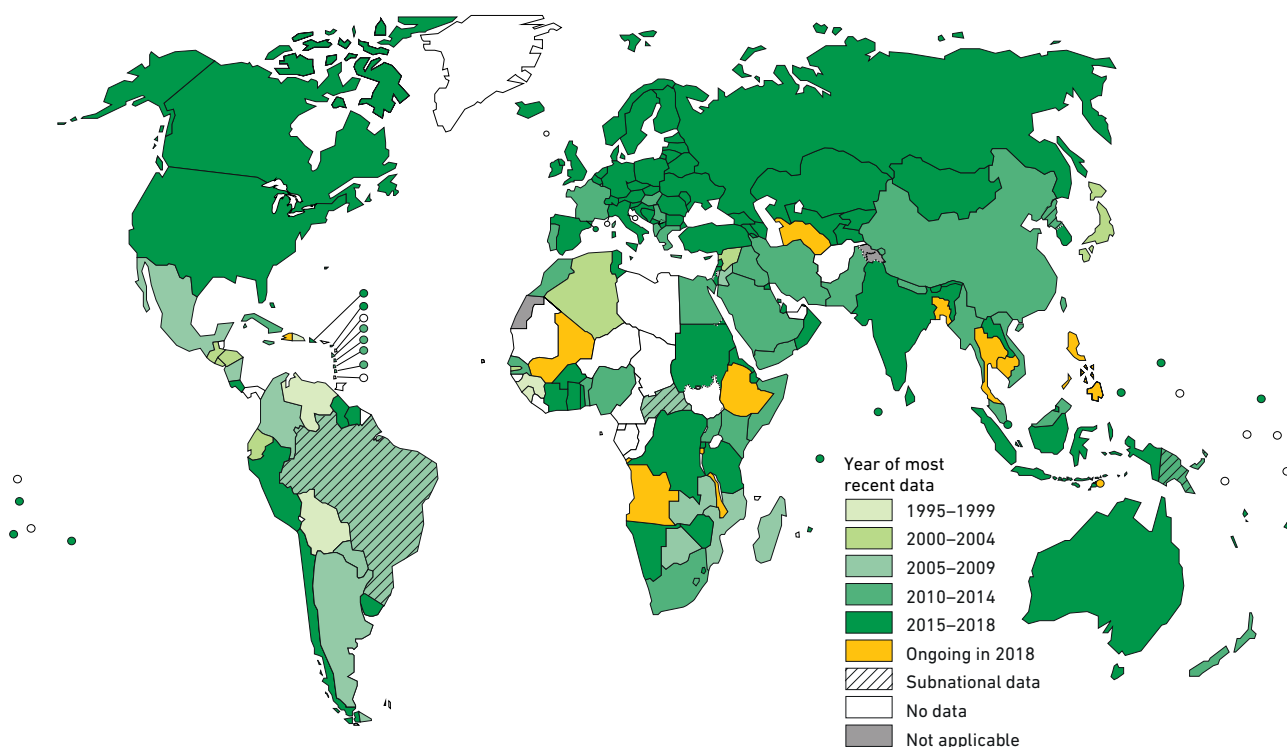


FIG. 3.19

Global coverage of surveillance data on drug resistance, 1995–2018



obtained from all TB patients, and 69 countries that rely on epidemiological surveys of bacterial isolates collected from representative samples of patients (Fig. 3.18). Surveys conducted about every 5 years represent the most common approach to investigating the burden of drug resistance in resource-limited settings where routine DST is not accessible to all TB patients owing to lack of laboratory capacity or resources.

Progress towards achieving global coverage of drug-resistance surveillance data is shown in Fig. 3.19. Among the 30 high TB burden countries and the 30 high MDR-TB burden countries (which comprise a total of 40 countries, because of overlap between the two groups¹), 37 have data on levels of drug resistance. The three countries that have never conducted a drug-resistance survey are Angola, Congo and Liberia. Angola has initiated a national survey in 2018. Among the other 37 high TB burden or high MDR-TB burden countries, four countries (Brazil, Central African Republic, Democratic People’s Republic of Korea and Papua New Guinea) rely on drug-resistance surveillance data gathered from subnational areas only.

In 2016–2017, the first-ever national drug resistance surveys were completed in Eritrea, Indonesia and Lao People’s Democratic Republic, and repeat surveys were completed in Eswatini, Sri Lanka, Togo and the United

Republic of Tanzania. In 2017–2018, drug-resistance surveys were ongoing in 12 countries, with the first nationwide surveys in five countries (Angola, Burundi, Haiti, Mali and Timor-Leste) and repeat surveys in seven countries (Bangladesh, Cambodia, Ethiopia, Malawi, the Philippines, Thailand and Turkmenistan).

3.3.2 Estimates of the disease burden caused by MDR/RR-TB

Globally in 2017, an estimated 3.6% (95% confidence interval [CI]: 2.6–4.8%) of new cases and 17% (95% CI: 5.6–33%) of previously treated cases had MDR/RR-TB (Table 3.6). The proportions of new and previously treated TB cases with MDR/RR-TB at the country level are shown in Fig. 3.20 and Fig. 3.21.

There were an estimated 558 000 (range, 483 000–639 000) incident cases of MDR/RR-TB in 2017. This is a slight downward revision from the best estimate published in the 2017 edition of the WHO global TB report, for reasons explained in Box 3.2. As before, however, the proportion of cases estimated to have MDR-TB was 82% (460 000 out of 560 000) (Table 3.6). The countries with the largest numbers of MDR/RR-TB cases (47% of the global total) were China, India and the Russian Federation (Fig. 3.22).

There were about 230 000 (range, 140 000–310 000) deaths from MDR/RR-TB in 2017, similar to the best estimate for 2016 that was published in the 2017 edition of the WHO global TB report.

¹ For a full list of the high TB burden and high MDR-TB burden countries, see Chapter 2.

TABLE 3.6

Estimated incidence of MDR/RR-TB in 2017 for 30 high MDR-TB burden countries, WHO regions and globally

	ESTIMATED % OF NEW CASES WITH MDR/RR-TB		ESTIMATED % OF PREVIOUSLY TREATED CASES WITH MDR/RR-TB		INCIDENCE OF MDR/RR-TB				
	BEST ESTIMATE ^a	UNCERTAINTY INTERVAL	BEST ESTIMATE	UNCERTAINTY INTERVAL	NUMBER (IN 1000s)	UNCERTAINTY INTERVAL	RATE ^b	UNCERTAINTY INTERVAL	% OF MDR AMONG MDR/RR-TB
Angola	2.5	1.1–4.3	14	10–18	3.9	1.6–7.2	13	5.5–24	68
Azerbaijan	12	11–14	28	27–30	2.1	1.5–2.8	22	16–28	100
Bangladesh	1.6	0.74–2.8	29	24–35	8.4	3.8–15	5.1	2.3–9.0	90
Belarus	38	36–41	67	63–70	2.5	1.8–3.3	26	19–35	97
China	7.1	5.6–8.7	24	20–28	73	55–94	5.2	3.9–6.6	75
DPR Korea	2.2	0.82–4.2	16	9.1–25	5.2	2.6–8.8	20	10–34	88
DR Congo	2.2	1.0–3.5	9.6	8.8–10	7.5	3.3–13	9.2	4.0–17	68
Ethiopia	2.7	1.6–4.1	14	6.7–25	5.5	2.9–8.9	5.2	2.8–8.4	60
India	2.8	2.0–3.5	12	10–13	135	78–208	10	5.8–16	92
Indonesia	2.4	1.8–3.3	13	9.0–18	23	16–31	8.8	6.2–12	91
Kazakhstan	26	25–28	44	42–46	7.1	4.2–11	39	23–59	91
Kenya	1.3	0.74–2.0	4.4	3.7–5.2	2.8	1.2–4.9	5.6	2.5–9.9	55
Kyrgyzstan	26	24–27	61	58–64	4.1	3.0–5.4	68	50–89	87
Mozambique	3.7	2.5–5.2	20	5.2–40	8.8	4.6–14	30	15–48	89
Myanmar	5.1	3.4–7.2	27	16–40	14	8.0–21	26	15–39	98
Nigeria	4.3	3.2–5.5	25	19–31	24	14–36	12	7.3–19	66
Pakistan	4.2	3.2–5.3	16	15–17	27	17–39	14	8.8–20	88
Papua New Guinea	3.4	1.7–5.0	26	15–36	1.9	1.2–2.8	23	15–34	77
Peru	6.3	5.9–6.7	20	19–22	3.5	2.6–4.5	11	8.1–14	84
Philippines	2.6	1.9–3.4	28	27–28	27	12–47	26	12–45	60
Republic of Moldova	28	25–30	55	51–59	2.2	1.8–2.7	54	43–66	97
Russian Federation	32	31–33	67	66–67	56	36–82	39	25–57	88
Somalia	8.7	6.1–12	47	29–65	4.2	2.3–6.5	28	16–44	62
South Africa	3.4	2.5–4.3	7.1	4.8–9.5	14	8.9–20	25	16–36	62
Tajikistan	20	19–22	23	20–27	2.3	1.6–3.1	26	18–35	100
Thailand	2.2	1.5–3.0	24	18–31	3.9	2.5–5.7	5.7	3.6–8.2	87
Ukraine	28	27–29	48	47–49	20	13–30	46	29–67	77
Uzbekistan	15	14–16	57	55–60	7.5	4.7–11	24	15–34	52
Viet Nam	4.1	2.8–5.7	17	17–18	7.1	4.6–10	7.4	4.8–11	98
Zimbabwe	4.6	3.0–6.2	14	8.9–20	2.3	1.4–3.5	14	8.5–21	40
MDR/RR HBCs	10	9.0–11	29	16–43	507	431–588	11	9.3–13	82
Africa	2.4	1.4–3.7	15	0.81–43	90	76–106	8.6	7.2–10	68
The Americas	1.9	0.89–3.2	14	5.0–25	11	9.9–13	1.1	0.99–1.3	86
Eastern Mediterranean	3.8	2.6–5.3	21	4.9–45	41	31–53	6.0	4.5–7.7	79
Europe	7.1	5.9–8.4	21	15–29	109	86–136	12	9.4–15	84
South-East Asia	3.4	2.5–4.4	19	9.6–31	192	131–264	9.7	6.7–13	92
Western Pacific	2.1	0.99–3.6	14	3.9–30	114	89–142	6.0	4.7–7.5	73
GLOBAL	3.6	2.6–4.8	17	5.6–33	558	483–639	7.4	6.4–8.5	82

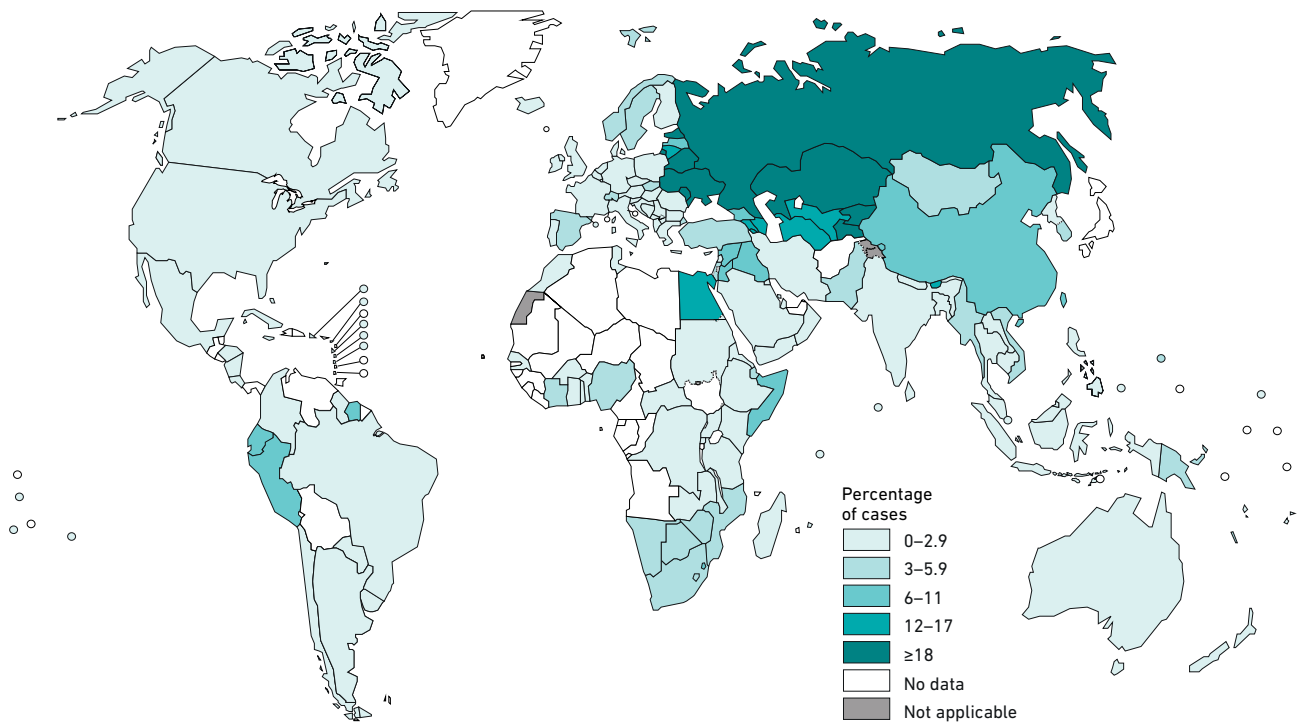
Numbers shown to two significant figures if under 100 and to three significant figures otherwise.

^a Best estimates are for the latest available year.

^b Rates are per 100 000 population.

FIG. 3.20

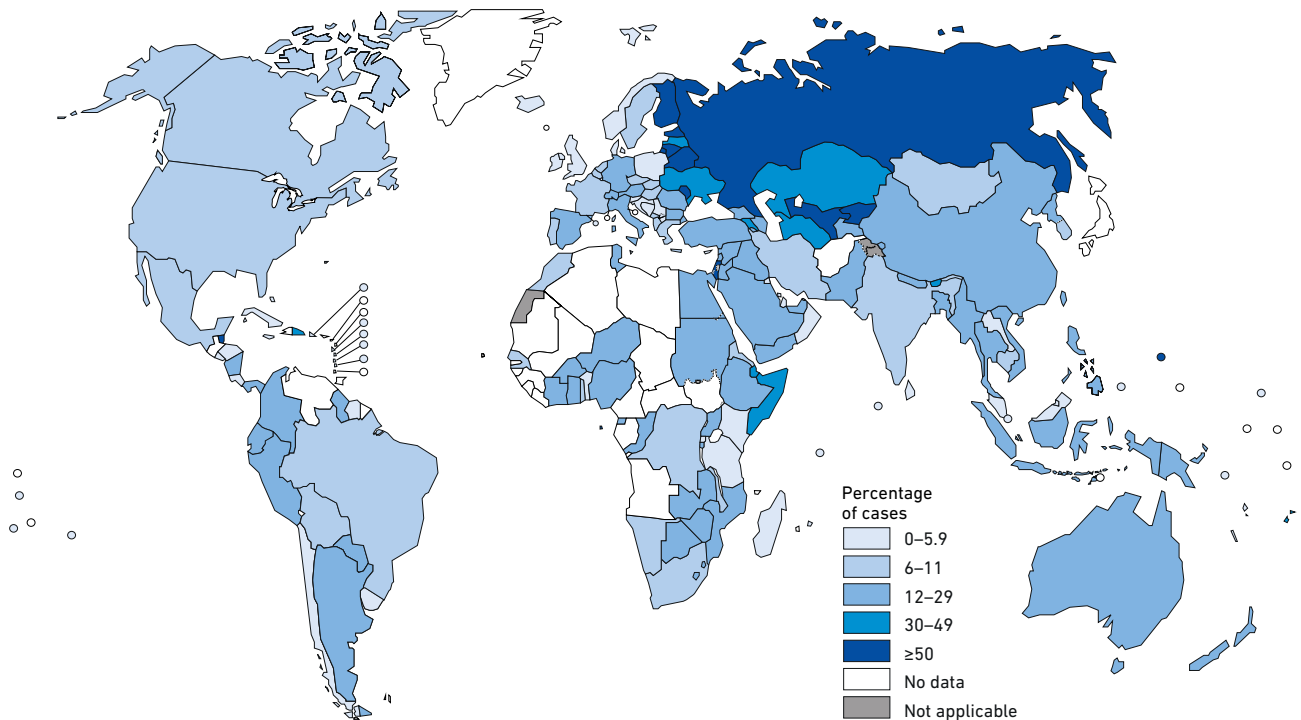
Percentage of new TB cases with MDR/RR-TB^a



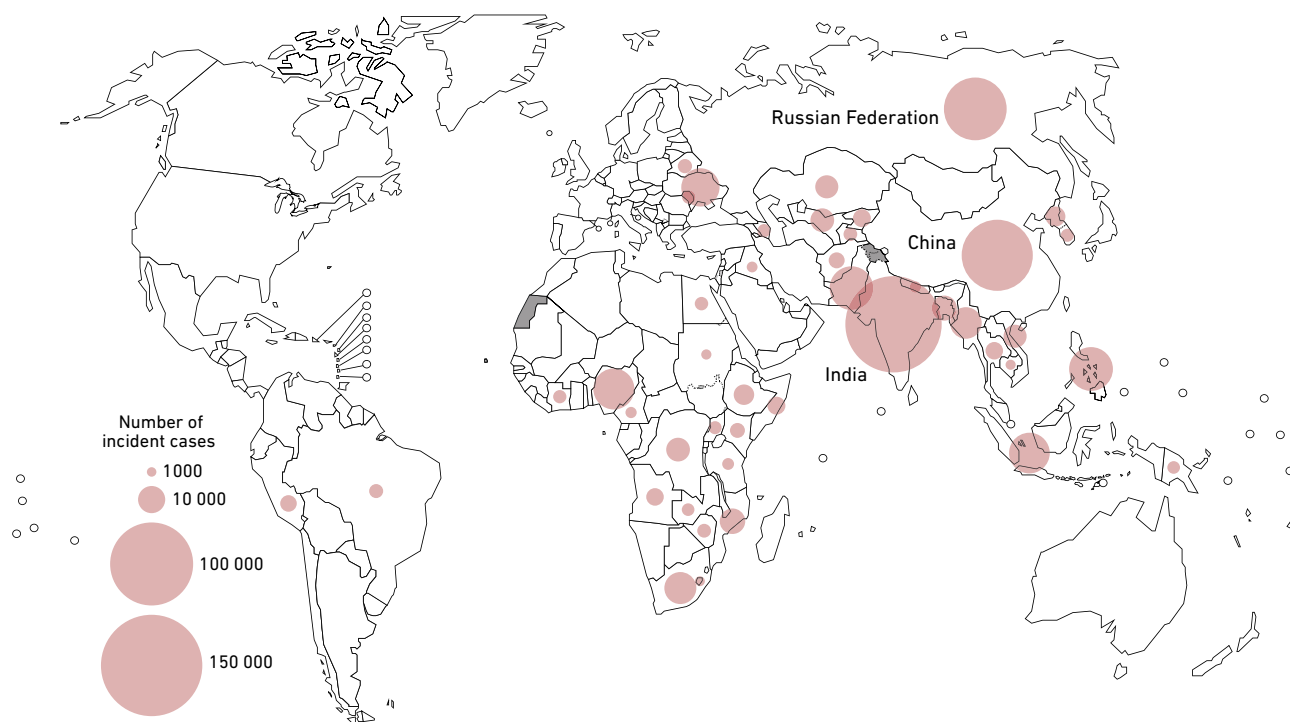
^a Figures are based on the most recent year for which data have been reported, which varies among countries. Data cover the period 2002–2018.

FIG. 3.21

Percentage of previously treated TB cases with MDR/RR-TB^a



^a Figures are based on the most recent year for which data have been reported, which varies among countries. Data cover the period 2005–2018. The high percentages of previously treated TB cases with RR-TB in Belize, Guam and Sao Tomé and Príncipe refer to only a small number of notified cases (range: 1–8 notified previously treated TB cases).

FIG. 3.22**Estimated incidence of MDR/RR-TB in 2017, for countries with at least 1000 incident cases**

Data compiled from surveys and continuous surveillance of drug resistance among TB patients also allow estimation of the number of MDR/RR-TB cases among notified TB patients with pulmonary TB. These are the MDR/RR-TB cases that could be detected if all notified patients were tested for drug resistance using WHO-recommended diagnostic tests. Globally in 2017, there were an estimated 330 000 (range, 310 000–350 000) MDR/RR-TB cases among notified TB patients.

3.3.3 Trends in drug resistance

Of the 40 countries with a high TB or MDR-TB burden (or both), only 22 have repeated a survey at least once to evaluate trends in drug resistance. Among these countries, 10 have at least 3 years of data: Belarus, Kazakhstan, Myanmar, Peru, Republic of Moldova, Russian Federation, Tajikistan, Thailand, Ukraine and Viet Nam. For these settings, **Fig. 3.23** shows trends in the number of new TB cases notified, the proportion of new TB cases with MDR, and per capita TB and MDR-TB rates. Based on these data, there is a slight trend for cases of MDR-TB to increase as a proportion of all TB cases in these countries, with the burden of MDR-TB either increasing faster or decreasing more slowly than the overall TB burden in each country.

3.3.4 Resistance to other anti-TB drugs

Data on resistance to all first-line anti-TB drugs (rifampicin, isoniazid, pyrazinamide and ethambutol)

was gathered from surveys conducted in six countries: Azerbaijan, Bangladesh, Belarus, Pakistan, South Africa and Ukraine. The average level of resistance to all first-line anti-TB drugs was 19% (95% CI: 18–20%) in new TB cases; in previously treated cases, the average was 43% (95% CI: 40–46%).

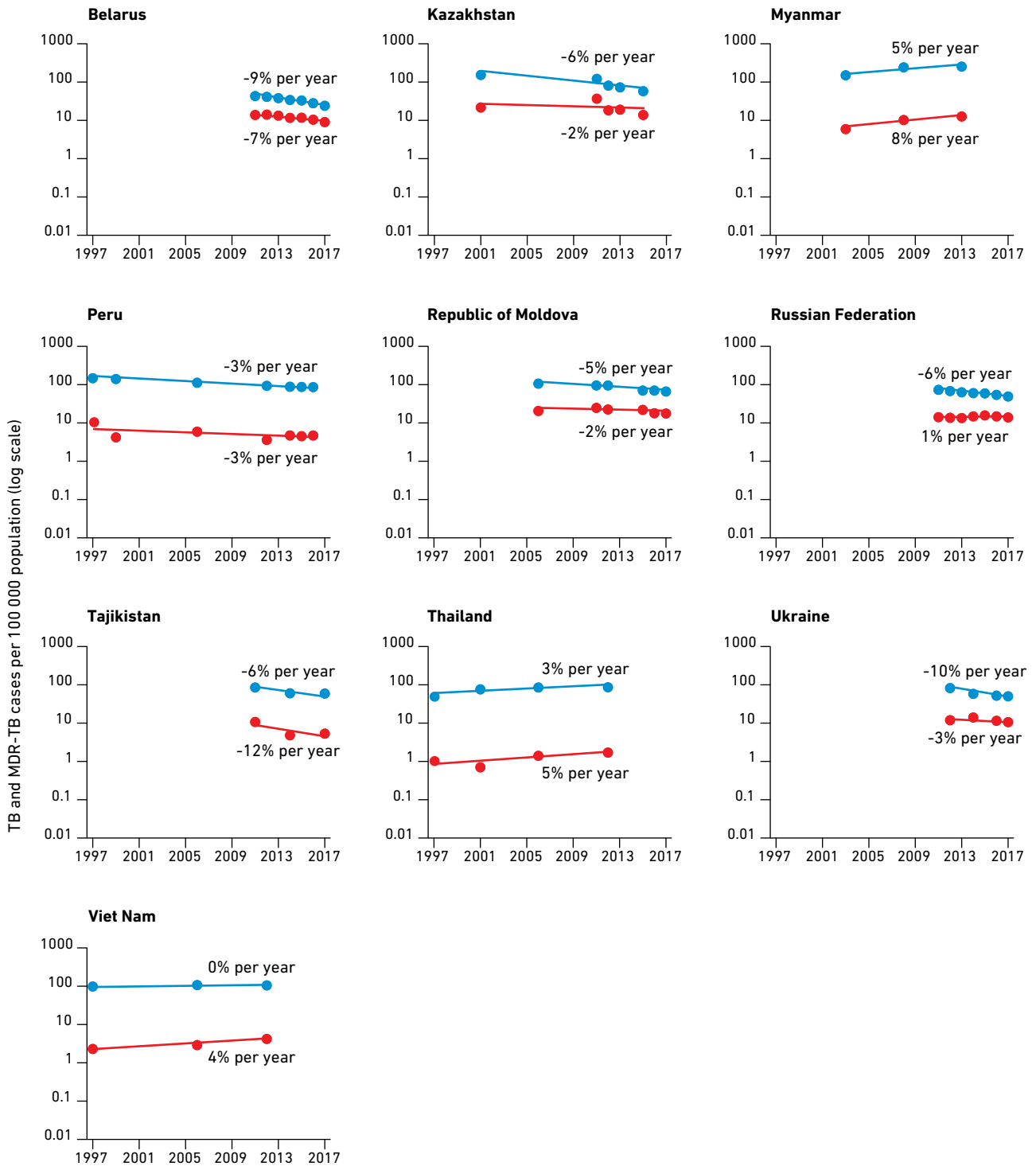
Data on levels of resistance to isoniazid without concurrent rifampicin resistance are available for 149 countries over the period 2003–2017. The proportions of TB patients resistant to isoniazid but susceptible to rifampicin in each country were weighted according to the number of new TB cases that were notified in the country, to generate a global average. The global averages of isoniazid resistance without concurrent rifampicin resistance were 7.1% (95% CI: 6.2–8.0%) in new TB cases and 7.9% (95% CI: 5.9–10%) in previously treated TB cases.

By the end of 2017, XDR-TB had been reported by 127 WHO Member States. Of these, 113 countries and five territories reported representative data from continuous surveillance or surveys regarding the proportion of MDR-TB cases that had XDR-TB. Combining their data, the average proportion of MDR-TB cases with XDR-TB was 8.5% (95% CI: 6.2–11%), an increase from the 6.2% in 2016 that was published in the 2017 edition of the WHO global TB report.

Among the 40 countries with a high TB or MDR-TB burden, 22 have surveillance data on resistance to second-line anti-TB drugs. The proportion of MDR/RR-TB

FIG. 3.23

Trends in levels of drug resistance in selected high MDR-TB burden countries with at least three years of data. The **blue** line shows the number of new notified TB cases per 100 000 population. The **red** line shows the number of MDR-TB cases among new TB patients per 100 000 population.



cases with resistance to any fluoroquinolone for which testing was done – including ofloxacin, levofloxacin and moxifloxacin – was 22% (95% CI: 17–28%).

3.4 National TB prevalence surveys

The prevalence of TB disease is not an indicator in the SDGs or a high-level indicator of the End TB Strategy, and no global target has been set for the period 2016–2035.¹ Furthermore, indirect estimates of prevalence suffer from considerable uncertainty, because they are derived from estimates of incidence and assumptions about disease duration. Hence, indirect estimates of TB prevalence (i.e. estimates that are not from a national TB prevalence survey) are not presented in this chapter.²

Nonetheless, in an important subset of countries with a large proportion of the world's TB burden, national TB prevalence surveys continue to provide the best method for measuring the burden of TB disease (both in absolute terms and to assess trends when repeat surveys are done, and by age and sex). Findings can inform assessment of actions needed to reduce this burden as well as estimates of TB incidence (Fig. 3.2), thus contributing to the monitoring of progress towards SDG and End TB Strategy targets. The WHO Global Task Force on TB Impact Measurement has retained national TB prevalence surveys within its strategic areas of work for 2016–2020 (Box 3.1), and has defined the group of countries where they continue to be relevant as those with a relatively high burden of TB (about 150 incident cases per 100 000 population) that do not yet have health, national notification and VR systems of the quality and coverage required to provide reliable and routine direct measurements of the number of TB cases and deaths.³ The most recent example of a prevalence survey that has informed understanding of trends in TB disease burden, estimates of TB incidence and identification of actions required to reduce the burden of TB disease is the 2016 survey in the Philippines.^{4,5}

¹ This is in contrast to the era of the Millennium Development Goals (MDGs) and Stop TB Strategy, when one of the global targets for reductions in TB disease burden was to halve prevalence between 1990 and 2015.

² WHO will continue to produce indirect estimates of TB prevalence. These can be provided upon request to tbdata@who.int.

³ In the Task Force's April 2016 meeting, epidemiological criteria for conducting a survey were defined for two groups of countries: those that implemented a survey in 2009–2015 and in which a repeat survey could be considered; and those that have never conducted a survey. There were 24 countries in the first group and 33 in the second group. For any of these 57 countries, it was emphasized that feasibility criteria must also be considered. In particular, the prerequisites for conducting a survey defined in the WHO handbook on national TB prevalence surveys should be met. For further details on the meeting, see World Health Organization Global Task Force on TB Impact Measurement. Report of the sixth meeting of the full Task Force; 19–21 April 2016, Glion-sur-Montreux, Switzerland. Geneva: WHO; 2016 (http://www.who.int/tb/advisory_bodies/impact_measurement_taskforce/meetings/tf6_report.pdf?ua=1, accessed 8 August 2018).

⁴ World Health Organization. Global Tuberculosis Report 2017 (WHO/HTM/TB/2017.23). Geneva: WHO; 2017 (http://www.who.int/tb/publications/global_report/archive/en/, accessed 21 June 2018).

⁵ The Philippines TB prevalence survey report (<http://ntp.doh.gov.ph/>, accessed 2 August 2018).

FIG. 3.24

Global progress in implementing national surveys of the prevalence of TB disease, actual (2000–2018) and expected (2019)

2000	China				
2001					
2002	Cambodia				
2003	Malaysia				
2004	Indonesia				
2005	Eritrea ^a				
2006	Thailand				
2007	Philippines	Viet Nam			
2008	Bangladesh ^a				
2009	Myanmar				
2010	China				
2011	Cambodia	Ethiopia	Lao PDR	Pakistan	
2012	Gambia	Nigeria	Rwanda	UR Tanzania	Thailand
2013	Ghana	Malawi	Sudan		
2014	Indonesia	Zambia	Zimbabwe		
2015	Bangladesh	Kenya	Mongolia	Uganda	
2016	DPR Korea	Philippines			
2017	Mozambique ^b	Myanmar ^b	Namibia ^c	South Africa ^b	Viet Nam ^c
2018	Botswana ^d	Eswatini ^b	Nepal ^b		
2019	India	Lesotho			

^a The surveys in Bangladesh (2008) and Eritrea (2005) collected sputum samples from all individuals (aged ≥15 years), and did not use chest X-ray and/or a symptom questionnaire to screen individuals for sputum submission.

^b Field operations are ongoing as of August 2018.

^c Field operations are completed and analysis is ongoing.

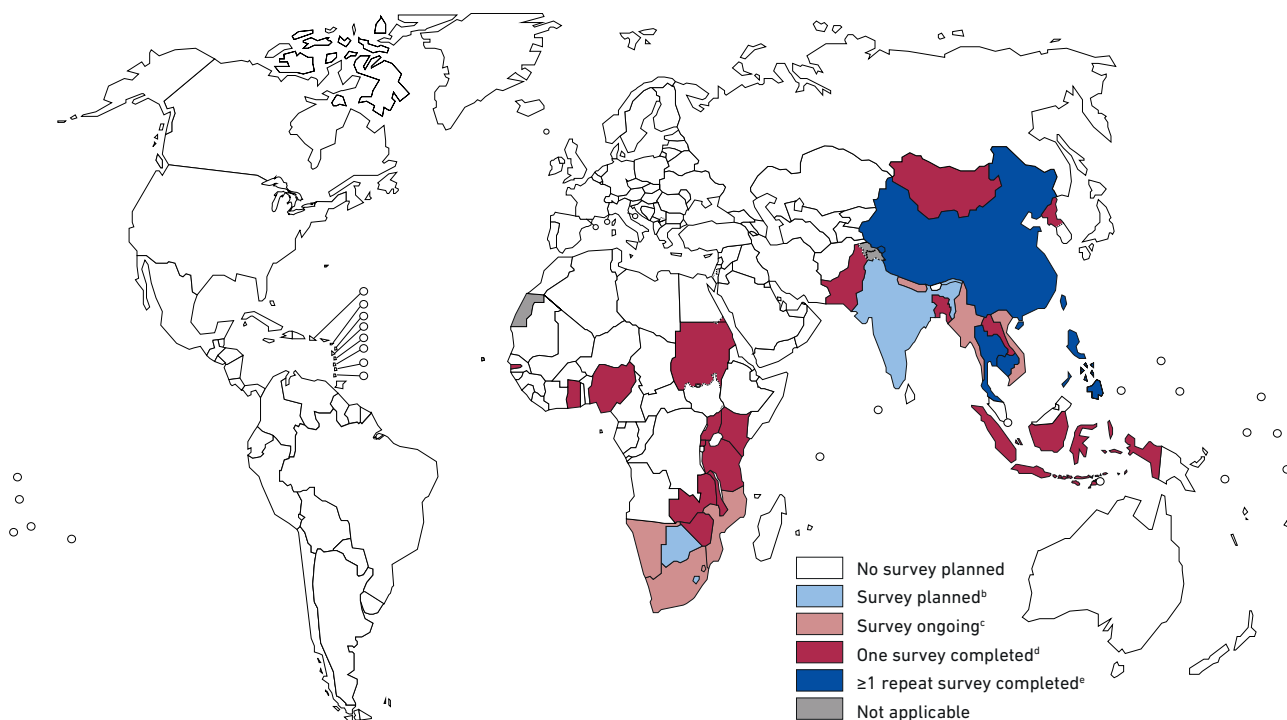
^d Field operations scheduled to start in 2018.

Countries in which national prevalence surveys were implemented in 2000–2018 or are planned for 2019 are shown in Fig. 3.24 and Fig. 3.25. An unprecedented number of surveys were implemented in 2007–2015, a period in which the WHO Global Task Force on TB Impact Measurement defined national TB prevalence surveys in 22 global focus countries as one of its three strategic areas of work (Box 3.1). Between 2007 and the end of 2017, a total of 25 surveys that used the screening and diagnostic methods recommended in the second edition of the WHO handbook on prevalence surveys⁶ were completed. This included 13 surveys in Asian countries and 12 in African countries. No surveys were completed in 2017; however, in early 2018, field operations were completed in the first national survey in Namibia and a repeat survey in Viet Nam (following a first survey in 2007). It is expected that Myanmar will finish its repeat survey (following a first survey in 2009) in late 2018. As of August 2018, field operations were ongoing in five

⁶ World Health Organization. Tuberculosis prevalence surveys: a handbook (WHO/HTM/TB/2010.17). Geneva: WHO; 2011 (www.who.int/tb/advisory_bodies/impact_measurement_taskforce/resources_documents/thelimebook/, accessed 8 August 2018).

FIG. 3.25

Countries in which national population-based surveys of the prevalence of TB disease have been implemented using currently recommended screening and diagnostic methods^a since 2000 or are planned in the future (status in August 2018)



^a Screening methods include field chest X-ray; at least culture was used to confirm diagnosis. The most recent surveys in Bangladesh, Kenya, Namibia, Myanmar, Mozambique, Nepal, the Philippines and Viet Nam used both culture and Xpert MTB/RIF (or Xpert Ultra) to confirm diagnosis.

^b A country has submitted at least a draft survey protocol and a budget plan to the WHO Global Task Force on TB Impact Measurement.

^c Countries were implementing field operations in August 2018 or were undertaking data cleaning and analysis.

^d A survey was conducted in accordance with WHO recommendations as outlined in 'Tuberculosis prevalence surveys: a handbook (2011)' and at least a preliminary report has been published.

^e A repeat national survey is one in which participants were screened with chest X-ray, and (at least) culture was used to diagnose TB cases.

countries: Eswatini, Mozambique, Myanmar, Nepal and South Africa. Surveys in Botswana, India and Lesotho are planned in 2019 or 2020.

A comparison of estimates of TB prevalence before and after the implementation of a national survey is shown for 25 countries in **Fig. 3.26**. Post-survey prevalence estimates were almost always more precise (i.e. had narrower uncertainty intervals). For 18 countries, estimates were within the pre-survey uncertainty interval, whereas for the other seven countries the survey found a burden that was either significantly above (six countries) or below (one country) the burden that had been estimated in the absence of survey data.

The distribution of TB disease by age (in adults) and sex based on prevalence survey data is shown in **Fig. 3.27** and **Fig. 3.28**. In Asia and some African countries (e.g. Ghana, Malawi, Rwanda, the United Republic of Tanzania and Zimbabwe), prevalence increases with age. However, in several African countries (e.g. Ethiopia, Gambia, Nigeria, Sudan, Uganda and Zambia), prevalence per 100 000 population peaks among those aged 35–54 years. The male to female (M:F) ratio of cases for the

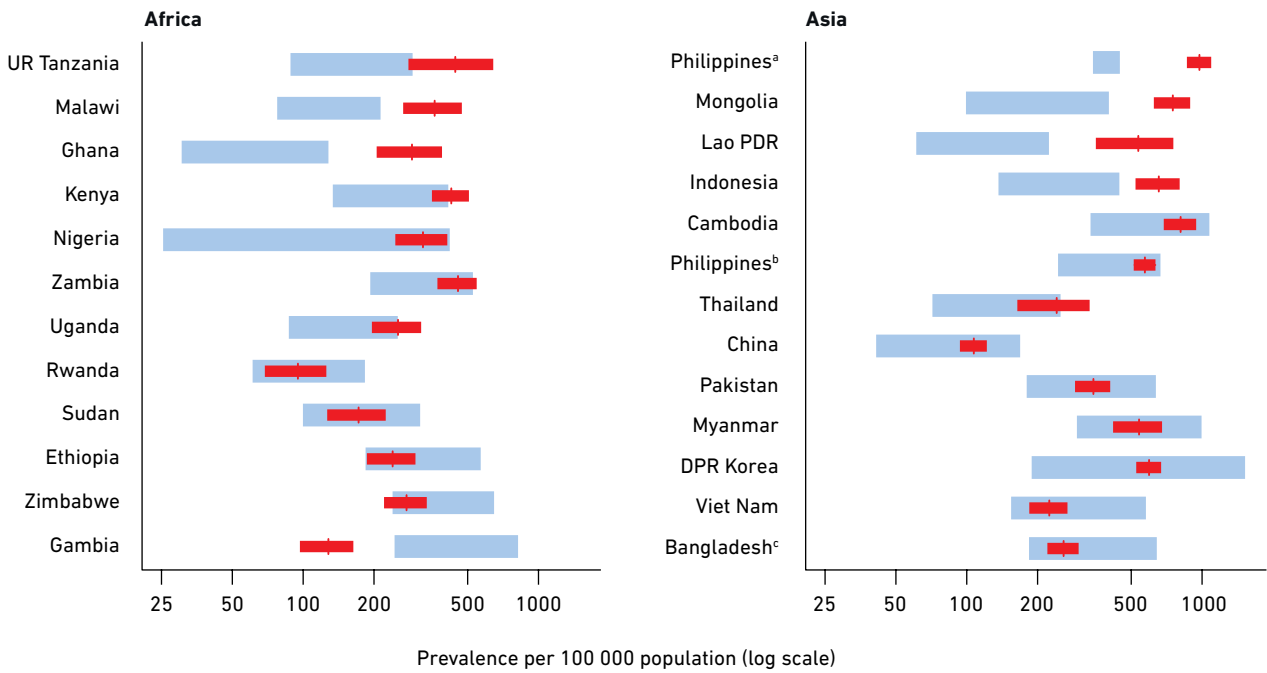
same set of surveys shows a systematically higher burden of TB disease among men, with ratios ranging from 1.2 (in Ethiopia) to 4.5 (in Viet Nam) for bacteriologically confirmed pulmonary TB. In most countries, the ratio was in the range 2–4.

The ratio of prevalence to notifications (P:N) can be used to assess detection and reporting gaps (**Fig. 3.29a**) and variation in these gaps by sex (**Fig. 3.29b**). The P:N ratios from surveys implemented in 2007–2017 suggest that women are accessing available diagnostic and treatment services more effectively than men. The higher disease burden in men, combined with larger gaps in detection and reporting, also suggests that there is a need for strategies to improve access to and use of health services among men.¹

¹ Horton KC, MacPherson P, Houben RMGJ, White RG, Corbett EL. Sex differences in tuberculosis burden and notifications in low- and middle-income countries: a systematic review and meta-analysis. Metcalfe JZ, ed. *PLoS Medicine*. 2016;13(9) (<https://www.ncbi.nlm.nih.gov/pubmed/27598345>, accessed 31 August 2018).

FIG. 3.26

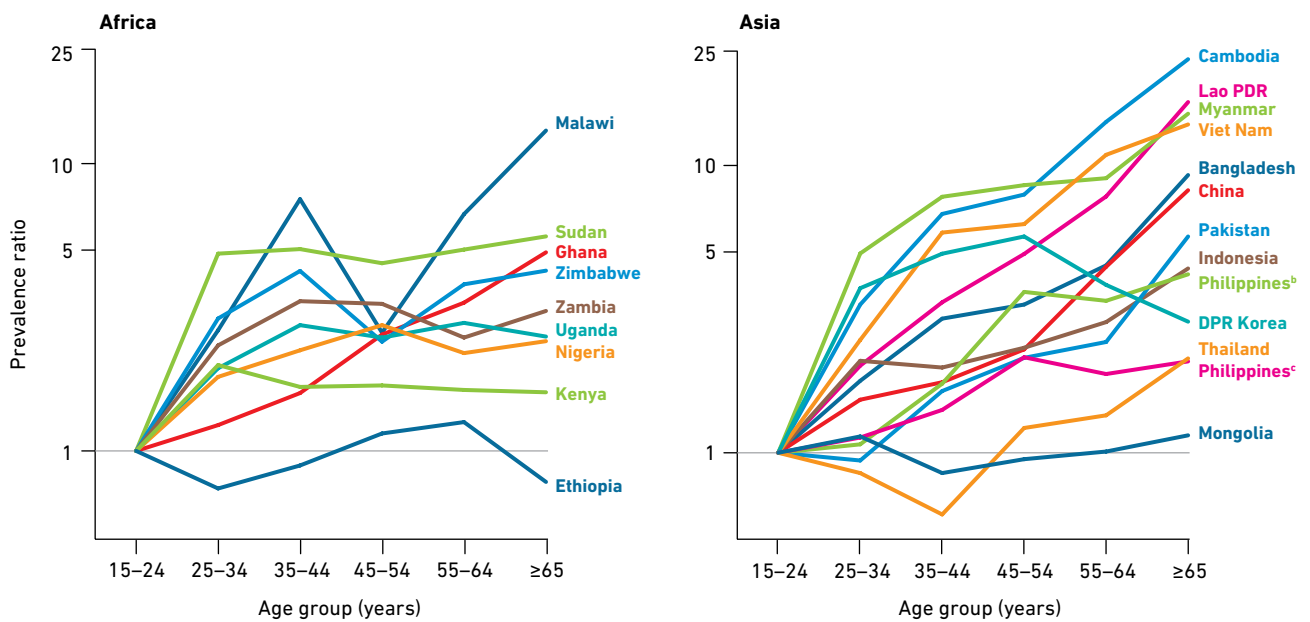
Estimates of TB prevalence (all ages, all forms of TB) for 25 surveys, before (in blue) and after (in red) results from national TB prevalence surveys became available since 2007. Countries are ordered according to the before–after difference.



^a These data relate to the repeat prevalence survey conducted in 2016.
^b These data relate to the prevalence survey conducted in 2007.
^c These data relate to the prevalence survey conducted in 2015–2016.

FIG. 3.27

Age-specific prevalence rate ratio of bacteriologically confirmed TB in surveys implemented 2007–2017^a



^a Age-specific prevalence ratios were calculated using the prevalence of the 15–24 year age group as the baseline. Data in the presented age groups were not available for Gambia and Rwanda. Data are not shown for UR Tanzania because laboratory challenges during the survey meant that it was only possible to directly estimate the prevalence of smear-positive (as opposed to bacteriologically confirmed) TB.
^b These data relate to the prevalence survey conducted in 2007.
^c These data relate to the repeat prevalence survey conducted in 2016.

3.5 Estimates of TB incidence and mortality disaggregated by age and sex

This section presents estimates of TB incidence and TB mortality disaggregated by age and sex.

3.5.1 Methods to disaggregate estimates by age and sex

Estimates of TB incidence in children (aged <15 years) were based on case notifications, adjusted for under-diagnosis and underreporting¹ and combined with estimates derived from dynamic modelling.² Results for the 0–14 year age group (0–4 and 5–14 years) in each country were then further disaggregated using outputs from an established deterministic model,² followed by disaggregation by sex using results from a meta-analysis of the M:F notification ratio. Estimates of TB incidence in adults were derived by first subtracting incidence in children from incidence in all ages. The estimate for adults was then disaggregated into six age groups (15–24, 25–34, 35–44, 45–54, 55–64 and ≥65 years) using data from national TB prevalence surveys implemented in 2007–2017 (Section 3.4). Country-specific distributions were used for countries that had implemented a survey; for other countries, the age distribution was predicted using prevalence survey data. Disaggregation by sex was based on actual M:F ratios for countries that had implemented surveys; for other countries, this disaggregation was based on regional M:F ratios from a systematic review and meta-analysis.³

TB mortality in children was estimated for the two age groups using a previously published approach derived from dynamic modelling,⁴ and then by sex, on the assumption that the pattern was the same as that for incidence. If available, data on TB deaths among adults were disaggregated for six age groups (15–24, 25–34, 35–44, 45–54, 55–64 and ≥65 years) using VR data. For countries whose mortality estimates were not based on VR data, a CFR was applied to the adult age- and sex-disaggregated incidence. This CFR accounted for differences between HIV-positive and HIV-negative TB cases, and for variation in HIV prevalence by age and sex.

FIG. 3.28

The male to female ratio of bacteriologically confirmed adult TB cases detected in prevalence surveys implemented 2007–2017^a



^a A value is not shown for UR Tanzania because laboratory challenges during the survey meant that it was only possible to directly estimate the prevalence of smear-positive (as opposed to bacteriologically confirmed) TB.

^b These data relate to the prevalence survey conducted in 2015–2016.

^c These data relate to the repeat prevalence survey conducted in 2016.

^d These data relate to the prevalence survey conducted in 2007.

¹ Jenkins HE, Tolman AW, Yuen CM, Parr JB, Keshavjee S, Pérez-Vélez CM et al. Incidence of multidrug-resistant tuberculosis disease in children: systematic review and global estimates. *The Lancet*. 2014;383(9928):1572–1579 (<http://www.ncbi.nlm.nih.gov/pubmed/24671080>, accessed 24 August 2018).

² Dodd PJ, Gardiner E, Coghlan R, Seddon JA. Burden of childhood tuberculosis in 22 high-burden countries: a mathematical modelling study. *Lancet Glob Health*. 2014;2(8):e453–459 (<http://www.ncbi.nlm.nih.gov/pubmed/25103518>, accessed 24 August 2018).

³ World Health Organization Global Task Force on TB Impact Measurement. Report of the sixth meeting of the full task force; 19–21 April 2016, Gion-sur-Montreux, Switzerland. Geneva: WHO; 2015 (http://www.who.int/tb/advisory_bodies/impact_measurement_taskforce/meetings/tf6_report.pdf?ua=1, accessed 8 August 2018).

⁴ Dodd PJ, Yuen CM, Sismanidis C, Seddon JA, Jenkins HE. The global burden of tuberculosis mortality in children: a mathematical modelling study. *Lancet Glob Health*. 2017;5(9):e898–e906 (<https://www.ncbi.nlm.nih.gov/pubmed/28807188>, accessed 24 August 2018).

FIG. 3.29a

The prevalence to notification (P:N) ratio of adult TB cases in prevalence surveys implemented 2007–2017^a

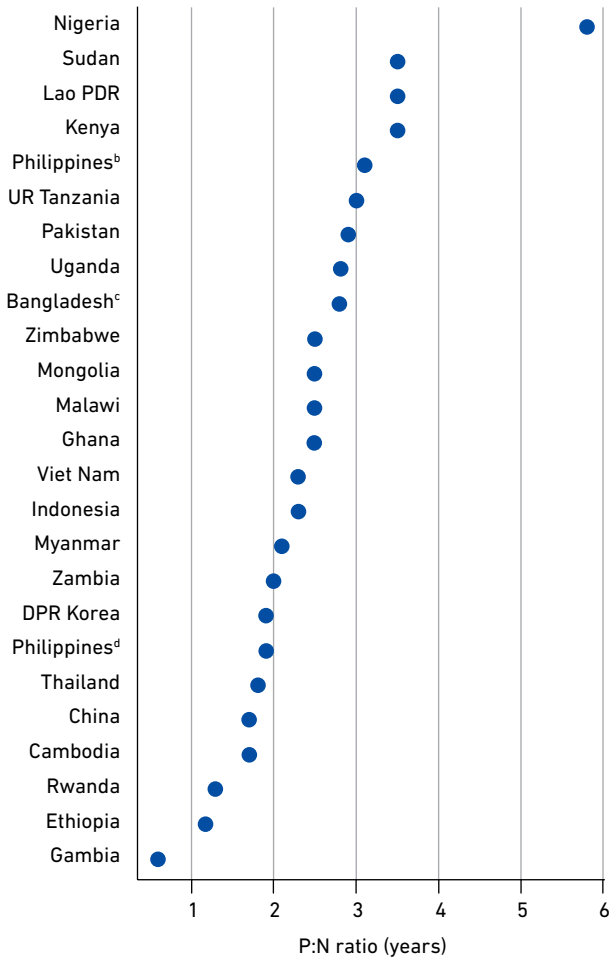
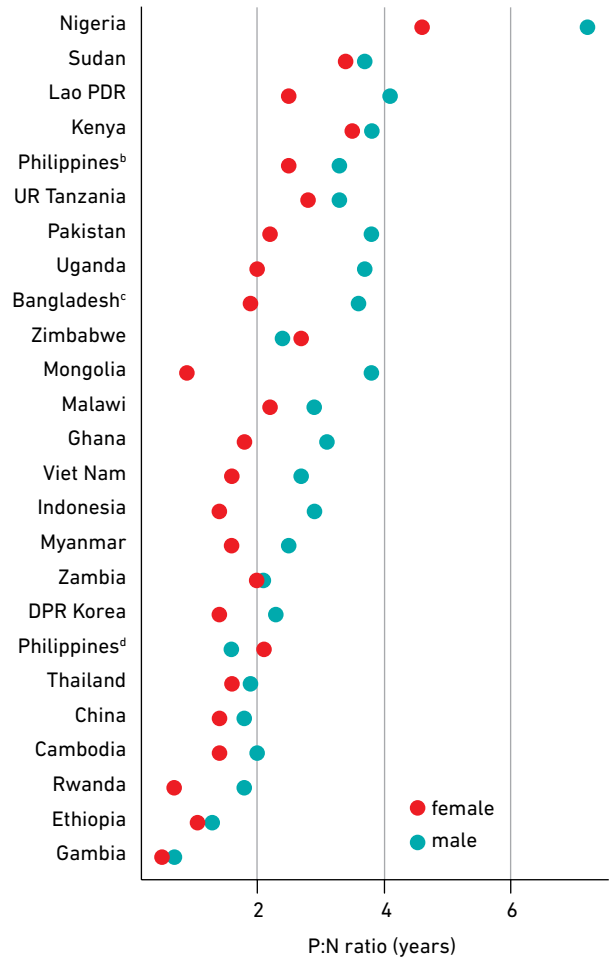


FIG. 3.29b

The prevalence to notification (P:N) ratio by sex for adult TB cases in prevalence surveys implemented 2007–2017^a



^a The P:N ratio is for smear-positive TB, except for Uganda and Zimbabwe where it is based on bacteriologically confirmed TB. Prevalence estimates are from a cross-sectional survey, and therefore only represent one point in time. Notification data are from the main year of the survey (shown in Fig. 3.24).
^b These data relate to the repeat prevalence survey conducted in 2016.
^c These data relate to the prevalence survey conducted in 2015–2016.
^d These data relate to the prevalence survey conducted in 2007.

Details of the methods used are provided in the [online technical appendix](#).¹

3.5.2 TB incidence disaggregated by age and sex

Estimates of TB incidence disaggregated by age and sex are shown in [Fig. 3.30](#) (global), [Fig. 3.31](#) (WHO regions) and [Fig. 3.32](#) (30 high TB burden countries), and in [Table 3.7](#). There were cases in all age groups but, overall, 90% were adults and 10% were children (aged <15 years). Globally, 64% of cases were among men and boys, and 36% were among women and girls.² The higher shares of TB cases among men shown in [Fig. 3.33](#), [Fig. 3.34](#) and [Fig. 3.35](#) are consistent with evidence from prevalence surveys, which show that TB disease affects men more than women ([Fig. 3.28](#)), and that gaps in case detection and reporting are higher among men ([Fig. 3.29](#)).

The M:F ratio of incident TB cases for all ages ranged from 1.3 in the WHO Eastern Mediterranean Region to 2.1 in the WHO Western Pacific Region. In children, the M:F ratio was close to 1.

FIG. 3.30

Global estimates of TB incidence (black line) and case notifications disaggregated by age and sex (female in red; male in green), 2017

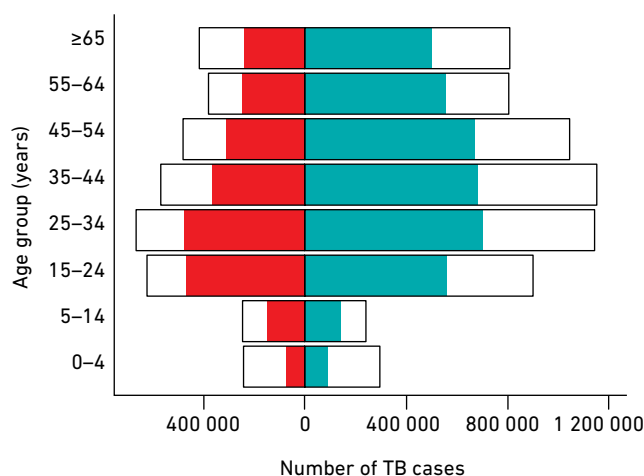
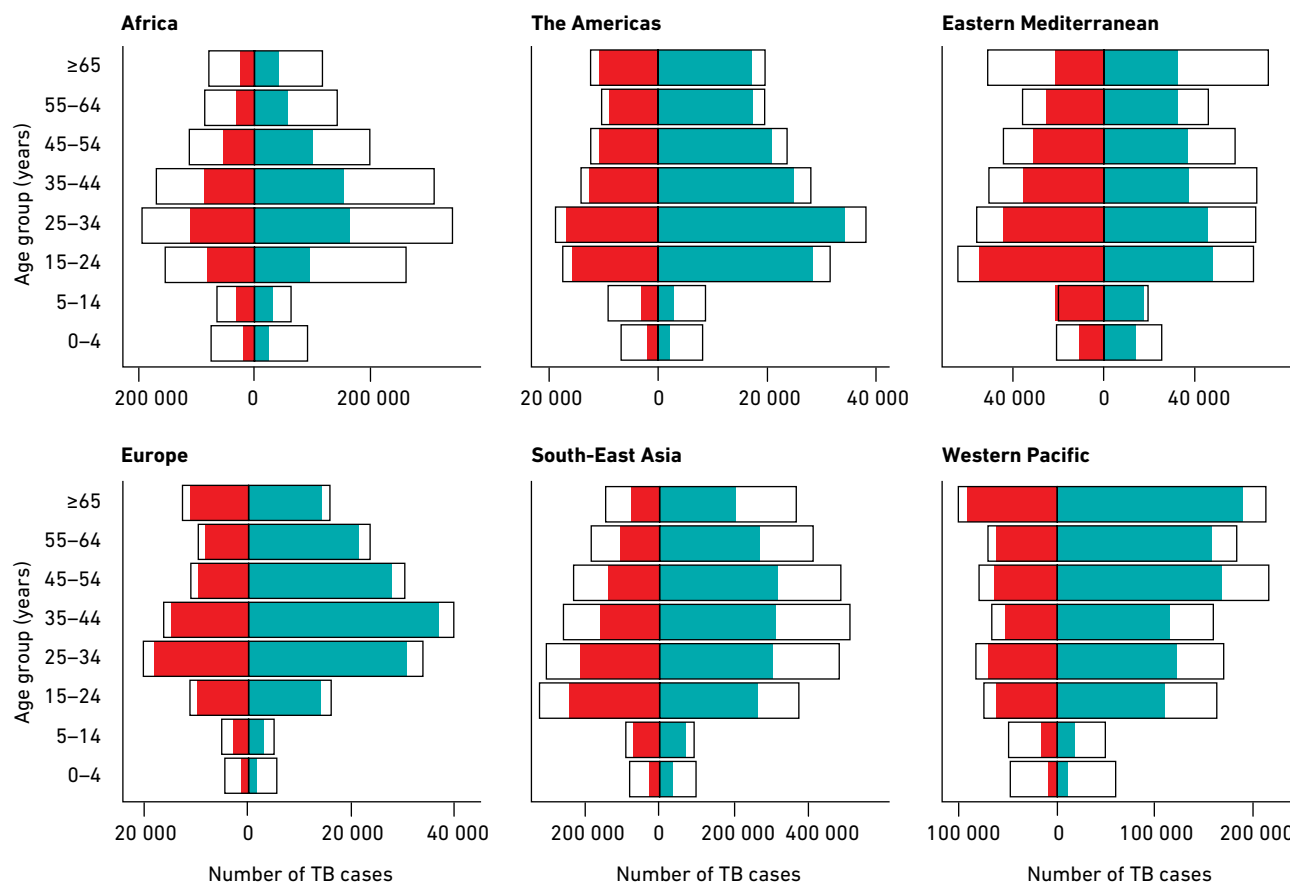


FIG. 3.31

Regional estimates of TB incidence (black line) and case notifications disaggregated by age and sex (female in red; male in green), 2017

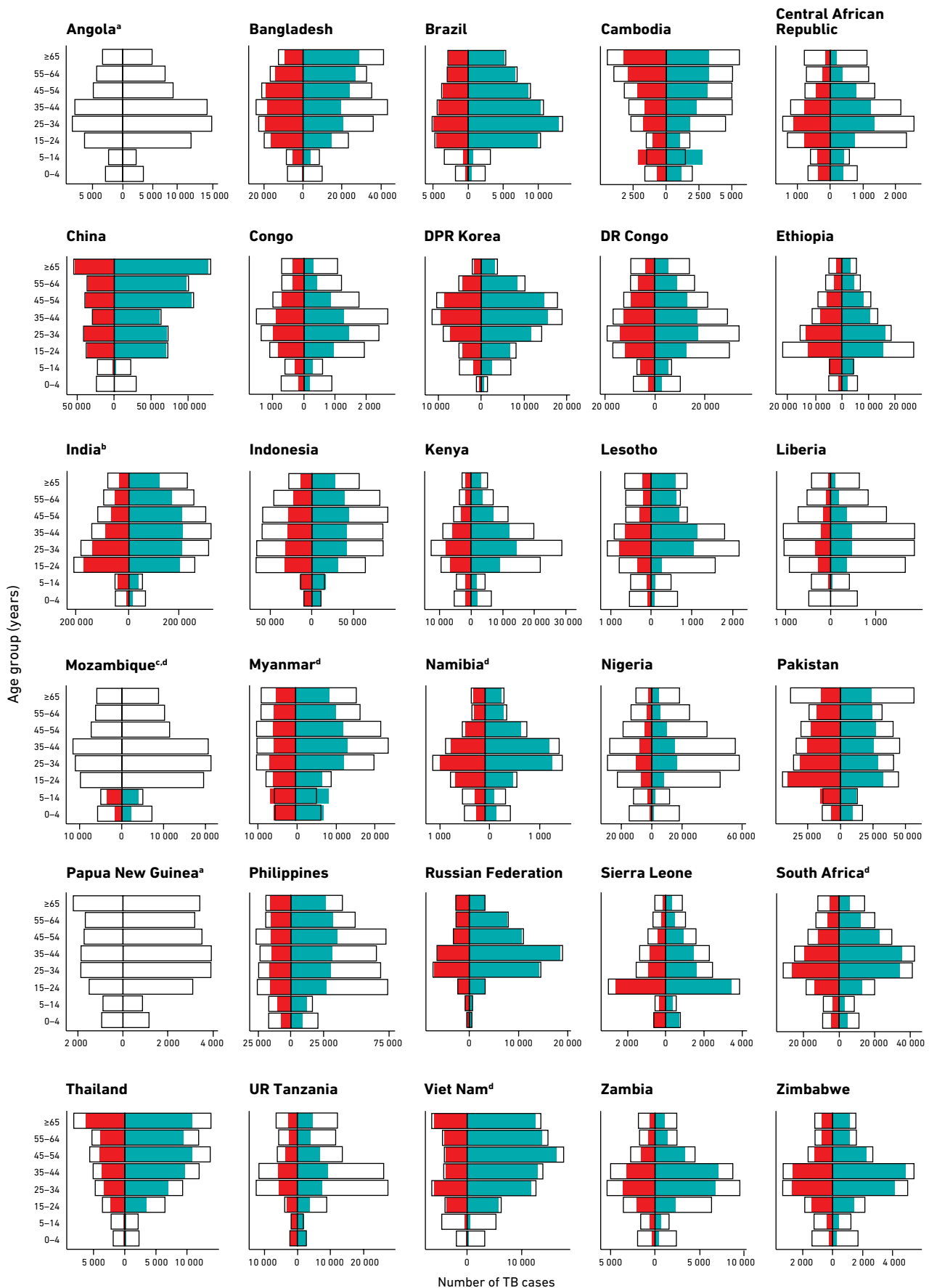


¹ The [online technical appendix](#) is available at www.who.int/tb/data.

² Further breakdowns by HIV status are not possible, because data on the HIV status of TB cases by age and sex are not available.

FIG. 3.32

Estimates of TB incidence (black line) and case notifications disaggregated by age and sex (female in red, male in green), 2017, in the 30 high TB burden countries



^a Age and sex disaggregated case notifications were not available.

^b Estimates of TB incidence for India are interim, pending results from the national TB prevalence survey planned for 2019/2020.

^c No age and sex disaggregated case notifications from 15 years and above were available for Mozambique.

^d Estimates of TB incidence for Mozambique, Myanmar, Namibia, South Africa and Viet Nam will be reviewed after final results from their respective national TB prevalence surveys are available in 2019.

TABLE 3.7
Estimated number of TB cases (in thousands) in children and adults, globally and for WHO regions, 2017^a

WHO REGION	TOTAL		MALE		FEMALE	
	BEST ESTIMATE	UNCERTAINTY INTERVAL	BEST ESTIMATE	UNCERTAINTY INTERVAL	BEST ESTIMATE	UNCERTAINTY INTERVAL
Africa	2 480	2 210–2 760	1 540	1 310–1 770	941	798–1 080
The Americas	282	262–302	179	162–197	102	92–112
Eastern Mediterranean	771	611–950	425	294–557	346	239–453
Europe	273	236–313	177	144–211	96	78–114
South-East Asia	4 440	3 530–5 450	2 820	1 990–3 660	1 620	1 140–2 090
Western Pacific	1 800	1 490–2 130	1 220	927–1510	577	439–715
GLOBAL	10 000	9 000–11 100	6 360	5 440–7 290	3 680	3 140–4 210

WHO REGION	TOTAL ≥15 YEARS		MALE ≥15 YEARS		FEMALE ≥15 YEARS	
	BEST ESTIMATE	UNCERTAINTY INTERVAL	BEST ESTIMATE	UNCERTAINTY INTERVAL	BEST ESTIMATE	UNCERTAINTY INTERVAL
Africa	2 180	1 910–2 450	1 380	1 150–1 620	800	665–936
The Americas	249	228–269	162	144–180	86	77–96
Eastern Mediterranean	685	517–853	380	249–511	305	200–410
Europe	252	214–290	166	133–200	86	68–103
South-East Asia	4 080	3 120–5 040	2 630	1 790–3 470	1 440	984–1 910
Western Pacific	1 590	1 270–1 910	1 110	815–1400	479	352–606
GLOBAL	9 030	7 980–10 100	5 830	4 900–6 760	3 200	2 690–3 710

WHO REGION	TOTAL 0–14 YEARS		MALE 0–14 YEARS		FEMALE 0–14 YEARS	
	BEST ESTIMATE	UNCERTAINTY INTERVAL	BEST ESTIMATE	UNCERTAINTY INTERVAL	BEST ESTIMATE	UNCERTAINTY INTERVAL
Africa	296	260–333	156	129–182	141	117–164
The Americas	33	30–36	17	15–19	16	14–18
Eastern Mediterranean	87	65–108	45	30–61	41	27–55
Europe	21	18–24	11	8.7–13	10	8.2–12
South-East Asia	362	277–447	190	130–251	171	117–226
Western Pacific	207	166–249	109	80–138	98	72–124
GLOBAL	1 010	888–1 120	529	445–613	478	401–554

^a Numbers shown to two significant figures if under 100 and to three significant figures otherwise.

3.5.3 TB mortality disaggregated by age and sex

Estimates of TB mortality in 2017 disaggregated by age and sex are shown in [Fig. 3.33](#) (global), [Fig. 3.34](#) (WHO regions) and [Fig. 3.35](#) (30 high TB burden countries), and in [Table 3.8](#). Estimates are shown for HIV-positive and HIV-negative people separately, given that the cause of TB deaths among HIV-positive people is classified as HIV in ICD-10 (see also [Section 3.2](#)).

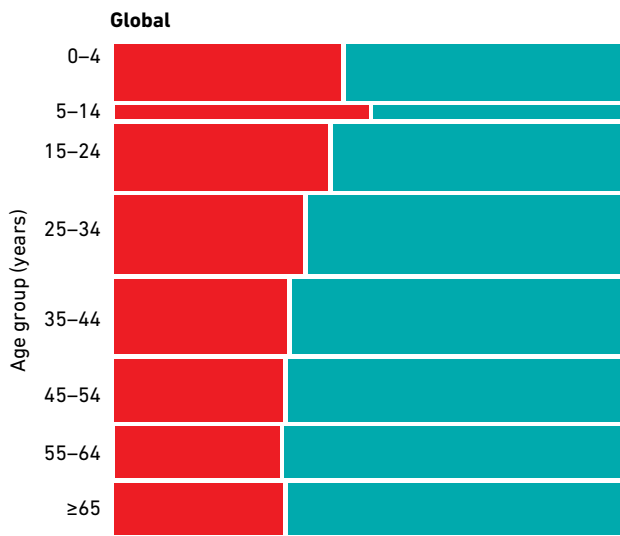
Globally in 2017, 63% of the TB deaths among HIV-

negative people were men and boys, and 37% were women and girls. Children (aged <15 years) accounted for 15% of total deaths, higher than their share of estimated cases, suggesting poorer access to diagnosis and treatment.

Globally in 2017, 57% of HIV-positive TB deaths were men and boys and 43% were women and girls. Children (aged <15 years) accounted for 10% of total deaths in HIV-positive people.

FIG. 3.33

Global distribution of TB mortality in HIV-negative people by age group and sex (female in red; male in green), 2017^a



^a The total area represents global TB mortality and all rectangles are proportional to their share of total TB mortality.

FIG. 3.34

Regional distribution of TB mortality in HIV-negative people by age group and sex (female in red; male in green), 2017^a



^a The total area represents TB mortality and all rectangles are proportional to their share of total TB mortality by region.

FIG. 3.35

Estimated distribution of TB mortality in HIV-negative people in the 30 high TB burden countries by age group and sex (female in red; male in green), 2017^a



^a The total area represents TB mortality and all rectangles are proportional to their share of total TB mortality by country.

^b Estimates of TB mortality for India are interim in nature, pending results from the national TB prevalence survey planned for 2019/2020.

^c Estimates of TB mortality for Mozambique, Myanmar, Namibia, South Africa and Viet Nam will be reviewed after final results from their respective national TB prevalence surveys are available in 2019.

TABLE 3.8

Estimated number of TB deaths (in thousands) by HIV status in children and adults, globally and for WHO regions, 2017^a

HIV-NEGATIVE										
WHO REGION	TOTAL		MALE 0-14 YEARS		FEMALE 0-14 YEARS		MALE ≥15 YEARS		FEMALE ≥15 YEARS	
	BEST ESTIMATE	UNCERTAINTY INTERVAL	BEST ESTIMATE	UNCERTAINTY INTERVAL	BEST ESTIMATE	UNCERTAINTY INTERVAL	BEST ESTIMATE	UNCERTAINTY INTERVAL	BEST ESTIMATE	UNCERTAINTY INTERVAL
Africa	413	348-485	31	23-39	27	20-33	229	170-289	127	94-160
The Americas	18	17-19	2.0	1.8-2.3	1.7	1.5-1.9	9.6	8.6-11	4.6	4.1-5.1
Eastern Mediterranean	89	75-104	7.8	5.7-9.8	6.5	4.8-8.3	41	30-53	33	24-42
Europe	24	23-25	2.4	2.2-2.6	2.3	2.1-2.4	13	12-14	6.1	5.6-6.5
South-East Asia	638	598-679	41	37-45	35	32-38	363	328-398	199	180-218
Western Pacific	92	85-100	21	18-24	17	15-20	37	31-43	17	15-20
GLOBAL	1 270	1 190-1 360	105	95-116	89	80-98	693	623-763	387	348-426

HIV-POSITIVE										
WHO REGION	TOTAL		MALE 0-14 YEARS		FEMALE 0-14 YEARS		MALE ≥15 YEARS		FEMALE ≥15 YEARS	
	BEST ESTIMATE	UNCERTAINTY INTERVAL	BEST ESTIMATE	UNCERTAINTY INTERVAL	BEST ESTIMATE	UNCERTAINTY INTERVAL	BEST ESTIMATE	UNCERTAINTY INTERVAL	BEST ESTIMATE	UNCERTAINTY INTERVAL
Africa	252	219-287	20	15-24	17	13-21	116	91-142	99	77-121
The Americas	6.0	5.3-6.7	0.25	0.21-0.29	0.21	0.18-0.24	4.1	3.5-4.8	1.4	1.2-1.6
Eastern Mediterranean	3.0	1.8-4.5	0.11	0.04-0.17	0.09	0.03-0.15	2.0	0.71-3.3	0.78	0.28-1.3
Europe	5.0	3.8-6.4	0.07	0.05-0.09	0.06	0.04-0.08	3.7	2.5-5.0	1.1	0.77-1.5
South-East Asia	28	22-36	0.68	0.45-0.91	0.58	0.39-0.78	20	13-26	7.4	4.9-9.9
Western Pacific	5.0	3.8-6.4	0.22	0.15-0.30	0.18	0.12-0.24	3.6	2.4-4.8	1.1	0.70-1.4
GLOBAL	300	266-335	21	17-25	19	15-22	149	121-176	111	91-132

^a Numbers shown to two significant figures if under 100 and to three significant figures otherwise.



A health officer teaching patients about TB and multidrug-resistant TB detection and treatment in a hospital in Lima, Peru

WHO / Pan American Health Organization

Chapter 4. Diagnosis and treatment: TB, HIV-associated TB and drug-resistant TB

KEY FACTS AND MESSAGES

Globally in 2017, 6.7 million people with tuberculosis (TB) were notified to national TB programmes (NTPs) and reported to WHO. Of these, just over 6.4 million had an incident episode (new or relapse) of TB. The global number of new and relapse TB cases notified and the notification rate per 100 000 population have both been increasing since 2013, following 4 years (2009–2012) in which 5.7–5.8 million new cases were notified annually. This increase is mostly explained by increased notifications in India (+44% between 2013 and 2017) and Indonesia (+21% between 2016 and 2017).

In 2017, 30% (2.0 million) of the 6.6 million new and previously treated TB cases notified globally were reported to have been tested for resistance to rifampicin. Coverage was 24% for new TB patients and 70% for previously treated TB patients. Globally, 160 684 cases of multidrug-resistant TB and rifampicin-resistant TB (MDR/RR-TB) were notified in 2017 (up from 153 119 in 2016), and 139 114 cases were enrolled in treatment (up from 129 689 in 2016).

Globally in 2017, 60% of notified TB patients had a documented HIV test result, up from 58% in 2016 and representing a 23-fold increase since 2004. In the WHO African Region, where the burden of HIV-associated TB is highest, 86% of TB patients had a documented HIV test result. A total of 464 633 TB cases among HIV-positive people were reported; of these, 84% were on antiretroviral therapy (ART).

Despite increases in notifications of TB, progress in closing detection and treatment gaps is slow and large gaps remain. Globally in 2017, there was a gap of 3.6 million between notifications of new and relapse cases and the best estimate of the number of incident cases (10.0 million). Ten countries accounted for around 80% of this global gap, with India (26%), Indonesia (11%), Nigeria (9%) and the Philippines (7%) accounting for more than half of the global total.

Gaps between the estimated number of new cases and the number actually reported are due to a mixture of underreporting of detected cases, and underdiagnosis (either because people do not access health care or because they are not diagnosed when they do). An informative example is Indonesia. In 2017, a national

study found that about 20% of new cases were not diagnosed; of the approximately 80% of new cases that were detected, 41% were not reported. Actions to correct underreporting are being put in place..

There are also large gaps in detection and treatment of MDR/RR-TB and HIV-associated TB. In 2017, the number of MDR/RR-TB cases started on treatment was only 25% of the estimated incidence of 558 000 cases, while the number of notified HIV-positive TB cases was only 51% of the estimated 920 000 new cases of TB among people living with HIV.

Ten countries accounted for 75% of the gap between enrolments in MDR-TB treatment in 2017 and the estimated number of incident MDR/RR-TB cases in 2017; two countries – China and India – accounted for 39% of the total gap. Most of the gaps in detection of HIV-positive TB cases and provision of ART in 2017 were accounted for by the WHO African Region.

The global male:female (M:F) ratio for notifications in 2017 was 1.7. Results from national TB prevalence surveys of adults show higher M:F ratios, indicating that notification data understate the share of the TB burden accounted for by men in some countries. Globally, children (aged <15 years) accounted for 7.1% of the new and relapse cases that were notified in 2017.

The WHO-recommended rapid diagnostic (WRD) test for detection of TB and rifampicin resistance currently available is the Xpert MTB/RIF® assay. Of the 48 countries in at least one of the lists of high burden countries, 32 had adopted national algorithms positioning the WRD as the initial diagnostic test for all people suspected of having pulmonary TB by the end of 2017.

The latest treatment outcome data show treatment success rates of 82% for TB (2016 cohort), 77% for HIV-associated TB (2016 cohort), 55% for MDR/RR-TB (2015 cohort) and 34% for extensively drug-resistant TB (XDR-TB) (2015 cohort). As part of efforts to improve outcomes for MDR/XDR-TB, 68 countries and territories reported that they had started using bedaquiline and 42 reported that they had started using delamanid by the end of 2017.

Prompt and accurate diagnosis of tuberculosis (TB), HIV-associated TB and drug-resistant TB, followed by provision of treatment in line with international standards, prevents deaths and limits ill-health among people who develop TB. It also prevents further transmission of infection to others. The 2020 and 2025 milestones for reductions in TB incidence and TB deaths set in the End TB Strategy ([Chapter 2](#)) require the case fatality ratio (i.e. the proportion of people with TB who die from the disease) to fall to 10% by 2020 and to 6.5% by 2025. The latter is only feasible if all people with TB are promptly diagnosed and effectively treated. Patient-centred care and prevention – backed by bold policies and supportive systems such as universal health coverage (UHC) and social protection – are Pillars 1 and 2 of the End TB Strategy ([Box 4.1](#)).

This chapter provides the latest national data reported to WHO on the diagnosis and treatment of TB, HIV-associated TB and drug-resistant TB. [Section 4.1](#) presents and discusses data for 2017 on notifications of TB cases and associated coverage of diagnostic testing, as well as trends since 2000. It includes data on the contribution of community engagement and public–public and public–private mix (PPM) initiatives to case-finding efforts. [Section 4.2](#) focuses on treatment coverage (and on detection and treatment gaps) for patients with TB, HIV-associated TB and drug-resistant TB, comparing numbers detected and treated with underlying estimates of disease burden. [Section 4.3](#) provides the most recent data on treatment outcomes, for new and relapse TB patients, TB patients living with HIV and patients with multidrug-resistant TB and rifampicin-resistant TB (MDR/RR-TB), as well as global trends for these three groups between 2012 and 2016.

Throughout the chapter, data are presented at global, regional and country levels, giving particular attention to high burden countries (HBCs).¹ Further country-specific details for all of the indicators covered in this chapter are provided in [Annex 2](#) and [Annex 4](#).

4.1 Case notifications and testing coverage

4.1.1 TB case notifications and bacteriological confirmation

In 2017, 6.7 million people with TB were notified to national TB programmes (NTPs) and reported to WHO ([Table 4.1](#)). Of these, 6.4 million had a new or relapse (incident) episode of TB (shown as the total of new and relapse cases), and an additional 260 000 had been previously diagnosed with TB but their treatment was changed to a retreatment regimen.

The number of new and relapse TB cases notified increased between 2000 and 2009, was then stable at around 5.7–5.8 million annually during 2009–2012, and

¹ The three lists of HBCs (for TB, HIV-associated TB and MDR-TB) are explained in [Chapter 2](#).

BOX 4.1

Pillars 1 and 2 of the End TB Strategy

Pillar 1 of the End TB Strategy is “Integrated, patient-centred care and prevention”. It has four components:

- early diagnosis of TB including universal drug susceptibility testing (DST), and systematic screening of contacts and high-risk groups;
- treatment of all people with TB, including drug-resistant TB, and patient support;
- collaborative TB/HIV activities, and management of comorbidities; and
- preventive treatment of persons at high risk, and vaccination against TB.

The fourth component of Pillar 1 is the topic of [Chapter 5](#).

Pillar 2 of the End TB Strategy is “Bold policies and supportive systems”. This pillar also has four components:

- political commitment with adequate resources for TB care and prevention;
- engagement of communities, civil society organizations, and providers of public and private care;
- UHC policy and regulatory frameworks for case notification, vital registration, quality and rational use of medicines, and infection control; and
- social protection, poverty alleviation and actions on other determinants of TB.

The components of Pillar 2 are primarily discussed in [Chapter 7](#).

For an overview of all aspects of the End TB Strategy, see [Chapter 2](#).

has subsequently increased ([Fig. 4.1](#)).² The increase since 2013 is mostly explained by a continuous increase in notifications in India (+44% between 2013 and 2017), following the introduction of a national policy of mandatory notification in 2012 and the rollout (also since 2012) of a nationwide web-based and case-based reporting system (called “Nikshay”) that facilitates reporting of detected cases by care providers in the public and private sectors. There was also an upturn in notifications in Indonesia in 2017 (an increase of 21% compared with 2016), for reasons discussed later in this chapter ([Section 4.2](#)).

² A similar pattern exists for case notification rates (new and relapse cases per 100 000 population per year). See [Fig. 3.7](#) and [Fig. 3.8](#) in [Chapter 3](#).

TABLE 4.1

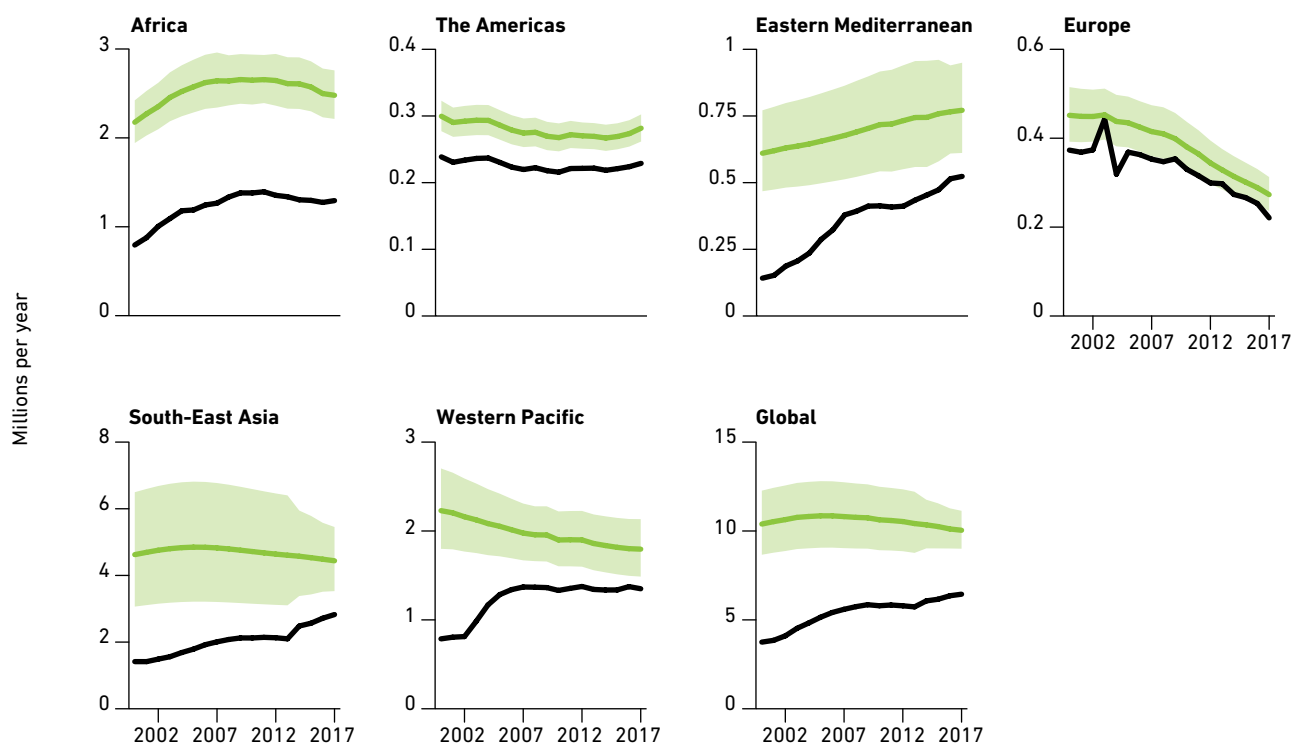
Notifications of TB, HIV-positive TB and MDR/RR-TB cases, globally and for WHO regions, 2017

	TOTAL NOTIFIED	NEW AND RELAPSE ^a	PULMONARY NEW AND RELAPSE		EXTRA-PULMONARY NEW AND RELAPSE (%)	HIV-POSITIVE NEW AND RELAPSE	MDR/RR-TB	XDR-TB
			NUMBER	OF WHICH BACTERIOLOGICALLY CONFIRMED (%)				
Africa	1 323 450	1 294 321	1 088 909	66%	16%	344 093	26 845	867
The Americas	243 064	228 943	194 228	78%	15%	20 487	4 084	121
Eastern Mediterranean	536 185	523 494	396 945	53%	24%	1 327	4 969	168
Europe	264 563	220 832	185 826	64%	16%	24 722	48 299	6 758
South-East Asia	2 965 311	2 826 486	2 403 901	59%	15%	61 832	51 788	2 755
Western Pacific	1 375 550	1 350 202	1 244 213	39%	8%	12 172	24 699	131
Global	6 708 123	6 444 278	5 514 022	56%	14%	464 633	160 684	10 800

^a New and relapse includes cases for which the treatment history is unknown. It excludes cases that have been re-registered as *treatment after failure*, as *treatment after lost to follow up* or as *other previously treated* (whose outcome after the most recent course of treatment is unknown or undocumented).

FIG. 4.1

Notifications of TB cases (new and relapse cases, all forms) (black) compared with estimated TB incident cases (green), 2000–2017, globally and for WHO regions. Shaded areas represent uncertainty bands.

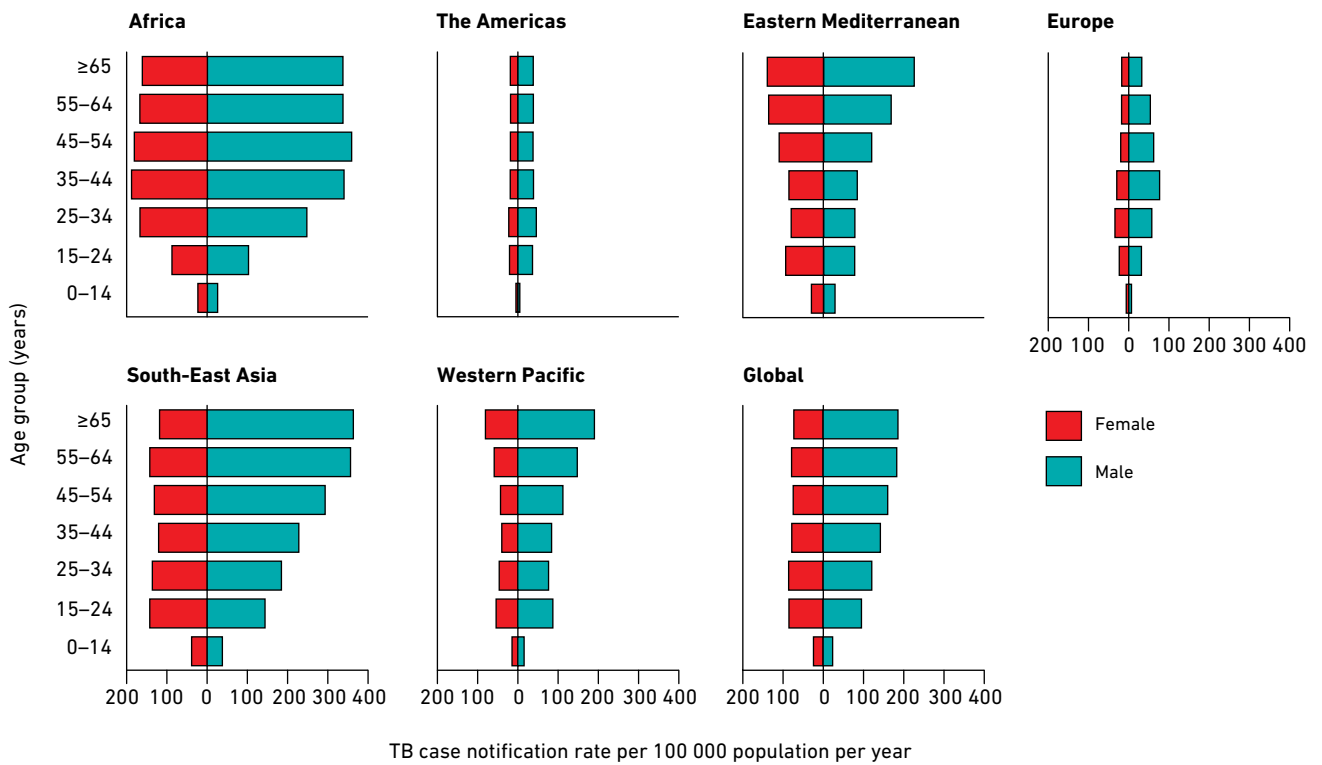


The distribution of notified cases in 2017 by age and sex is shown globally and for WHO regions in [Fig. 4.2](#). The global male:female (M:F) ratio for notifications was 1.7. In contrast, the M:F ratio in 25 national TB disease prevalence surveys of adults in African and Asian countries implemented in 2007–2017 was about 2.5 overall, and reached 4.5 in Viet Nam. This indicates that notification data understate the share of the TB burden accounted for by men in some countries (see [Chapter 3](#) for further details). Children (aged <15 years) accounted for 7.1% of the new and relapse cases that were notified

globally. In the WHO Eastern Mediterranean, South-East Asia and Western Pacific regions, the TB epidemic is a markedly ageing one, with a progressive increase in the notification rate with age, and a peak among those aged 65 years or over. Elsewhere, notification rates were highest among adults, most noticeably in the WHO African Region (in the 45–54 year age category) and the WHO European Region (in the 35–44 year age category). In several central and eastern European countries, as well as three high TB burden countries in Asia – China, Thailand and Viet Nam – less than 2% of notified cases

FIG. 4.2

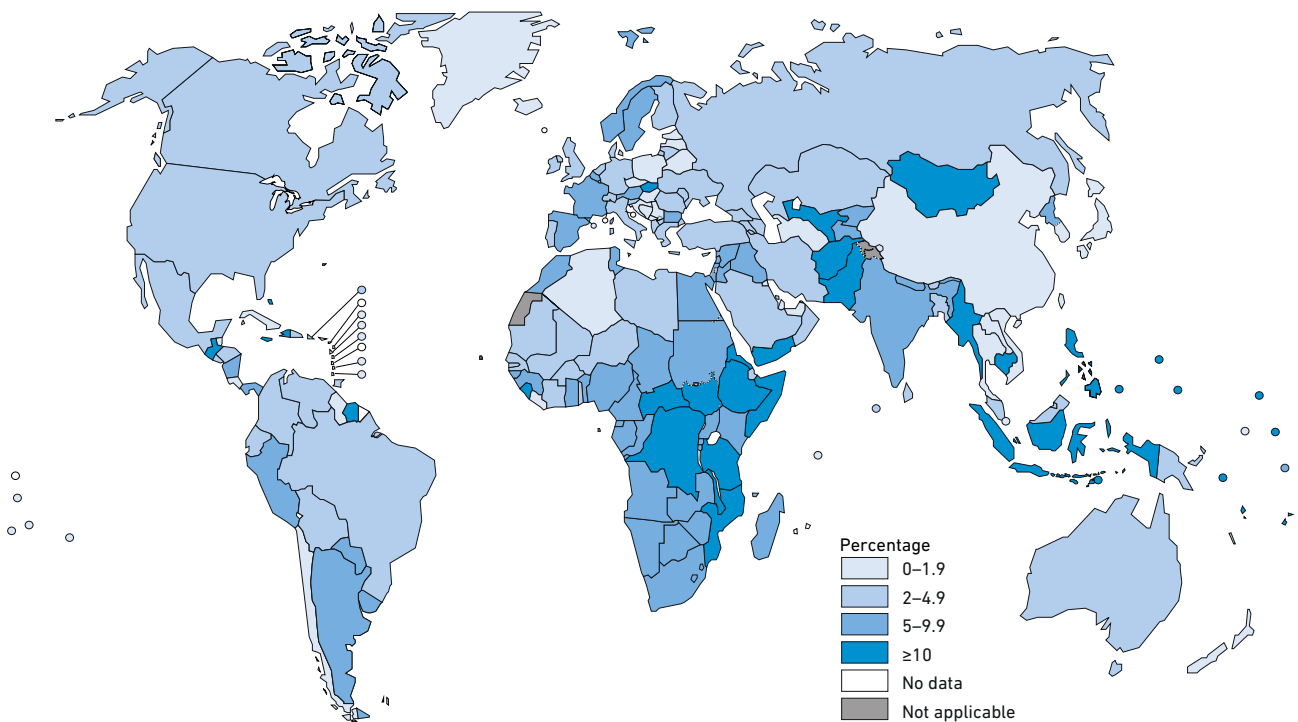
New and relapse TB case notification rates by age and sex^a in 2017, globally and for WHO regions



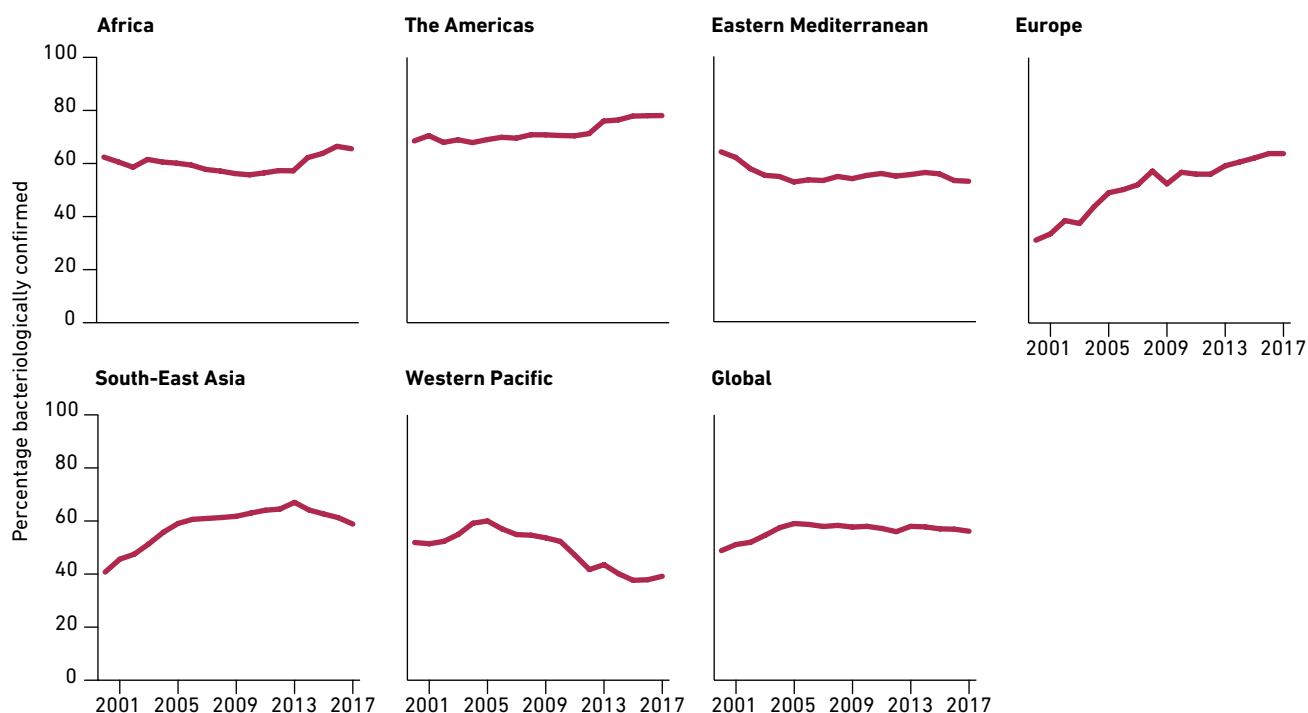
^a Countries not reporting cases in these categories are excluded. Cases included account for 89% of reported cases.

FIG. 4.3

Percentage of new and relapse TB cases that were children (aged <15), 2017^a



^a 2016 data were used for 23 countries.

FIG. 4.4**Percentage of new and relapse^a pulmonary TB cases with bacteriological confirmation, globally and for WHO regions, 2000–2017**

^a The calculation is for new pulmonary cases in years prior to 2013 based on smear results, except for the European Region where data on confirmation by culture was also available for the period 2002–2012.

were children (Fig. 4.3). Variation among countries in the child:adult and M:F ratios of cases may reflect real differences in epidemiology, differential access to or use of health-care services, or differential reporting practices.

Of the 5.5 million new and relapse pulmonary TB patients notified globally in 2017, 56% were bacteriologically confirmed.¹ The remaining patients were diagnosed clinically; that is, based on symptoms, abnormalities on chest radiography or suggestive histology. The percentage of pulmonary cases with bacteriological confirmation worldwide has declined slightly since 2013 (Fig. 4.4), mainly reflecting trends in the WHO South-East Asia Region (67% to 59%) and the WHO Western Pacific Region (43% to 39%). There was an improvement in the WHO African Region (57% to 66%) and the WHO European Region (59% to 64%). Considerable variation in the percentage of new and relapse pulmonary TB patients that are bacteriologically confirmed is evident, even among countries with a similar epidemiological profile (Fig. 4.5).

Reasons for a low proportion of cases being bacteriologically confirmed should be assessed at country level, as should reductions over time. The microbiological detection of TB is critical for infection control because it allows patients to be correctly diagnosed and started on

the most effective treatment regimen as early as possible. Most clinical features of TB and abnormalities on chest radiography or histology results generally associated with TB have low specificity, which may lead to false diagnoses of TB, and hence to people being enrolled in TB treatment unnecessarily.

Extrapulmonary TB represented 14% of the 6.4 million incident cases that were notified in 2017, ranging from 8% in the WHO Western Pacific Region to 24% in the WHO Eastern Mediterranean Region (Fig. 4.6 and Table 4.1).

Engagement of all care providers in the public and private sectors should be integral components of national TB strategies, and PPM initiatives have particular relevance to HBCs in Africa and Asia. The contribution of PPM to total notifications in countries that have reported PPM data for several years is shown in Box 4.2.

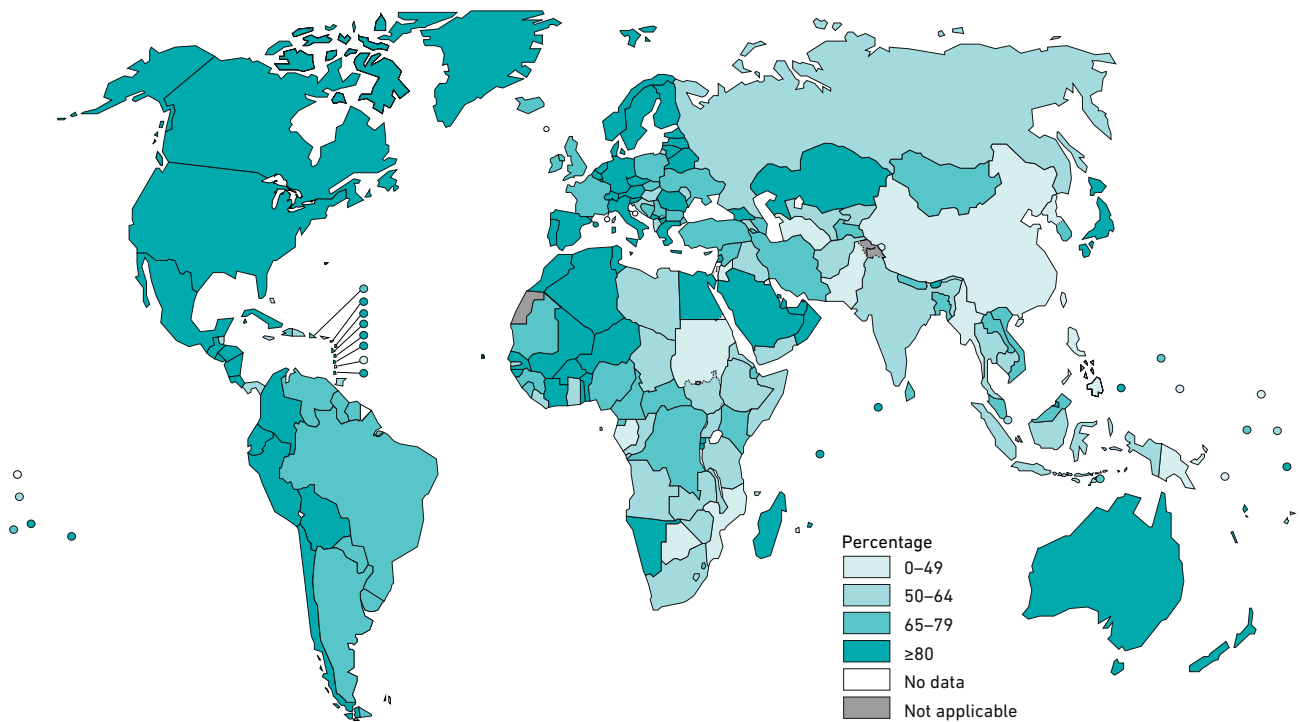
4.1.2 HIV testing for TB patients, detection of HIV-associated TB and screening for TB among people living with HIV

In 2017, 168 countries reported 3.8 million notified new and relapse TB patients with a documented HIV test result (a 4% increase from 2016), equivalent to 60% of notified TB cases. This represented a 23-fold increase in the number of people with TB tested for HIV since 2004, when WHO first asked countries to report data (Fig. 4.7). In 125 countries and territories, at least 75% of TB cases knew their HIV status (Fig. 4.8). Documentation of

¹ A bacteriologically confirmed case is one from whom a biological specimen is positive by smear microscopy, culture or WHO-recommended rapid diagnostic test, such as Xpert MTB/RIF.

FIG. 4.5

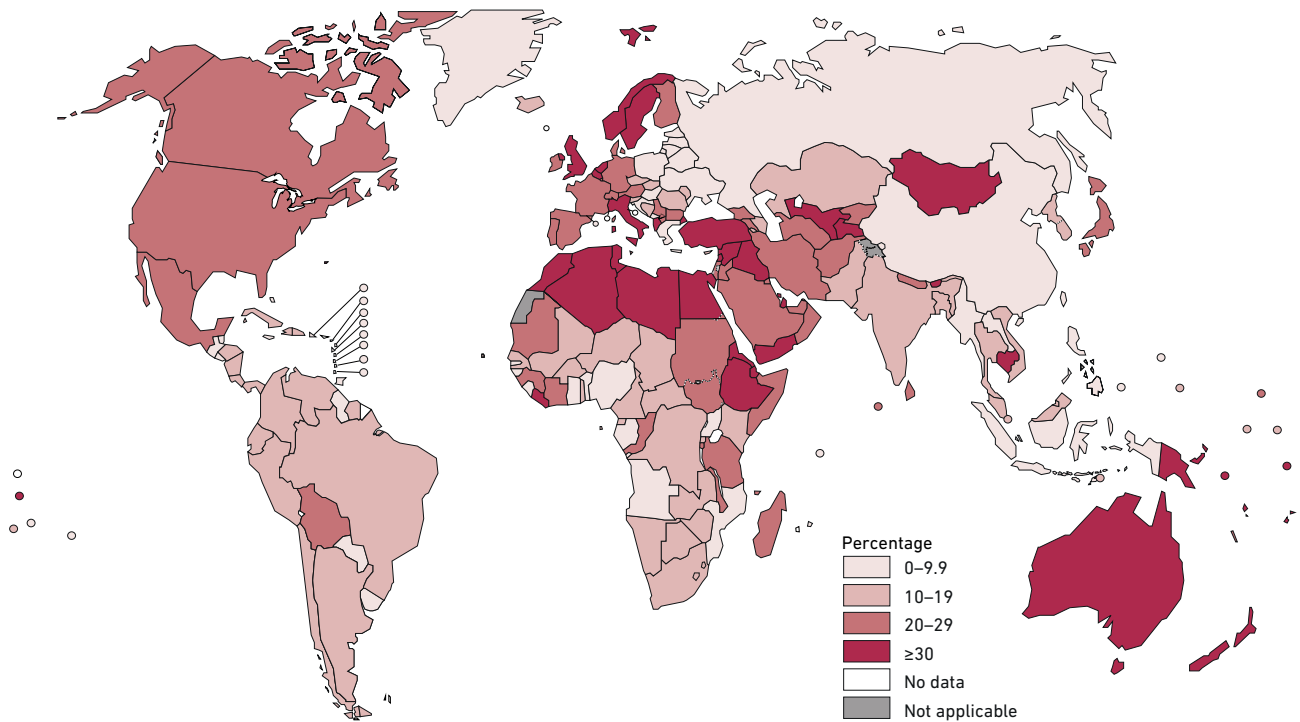
Percentage of new and relapse pulmonary TB cases with bacteriological confirmation, 2017^a



^a 2016 data were used for 18 countries.

FIG. 4.6

Percentage of extrapulmonary cases among new and relapse TB cases, 2017^a



^a 2016 data were used for 18 countries.

BOX 4.2

Trends in the contribution of public–private mix approaches to TB case notifications

One of the major challenges in ensuring universal access to quality TB services is the lack of systematic engagement of all health-care providers, especially those in the private sector, which is where people with TB often seek care initially. In 2017, about 3.6 million of the estimated 10 million people with TB worldwide were either not reported or not diagnosed (see [Section 4.2](#)); many of these people are likely to have been treated by public and private providers who are not linked to the NTP. The quality of care provided in these settings is difficult to ascertain and may be substandard.

WHO policies and global and national TB strategies have long acknowledged the need to engage all care providers, through PPM approaches. PPM encompasses diverse collaborative strategies. Public–public mix refers to engagement by the NTP of public sector providers of TB care that are not under the direct purview of the NTP (e.g. public hospitals, public medical colleges, prisons or detention centres, military

facilities, railways and public health insurance organizations). PPM refers to engagement by the NTP of private sector providers of TB care (e.g. private individual and institutional providers, the corporate or business sector, mission hospitals, nongovernmental organizations and faith-based organizations).

Many countries have made some progress in engaging non-NTP public providers and major non-profit facilities, especially those managed by faith-based organizations that have historically had close relationships with ministries of health. Gaps in engagement are typically more evident in the case of for-profit private providers.

Trends in the contribution of PPM to notifications in selected countries where PPM has been recognized as a priority and from which data have been reported to WHO for each year during 2012–2017 are shown in [Fig. B4.2.1](#) and [Fig. B4.2.2](#).

The contribution of PPM to case notifications quantifies the engagement of different providers in the delivery of TB care, and the alignment of their TB management practices with national and international standards.

Countries that have prioritized PPM – such as Bangladesh, India, Indonesia, Kenya and Pakistan – clearly show an increasing trend in the contribution of public non-NTP or private sector engagement to TB case notifications. As electronic case notification systems and digital technologies begin to be set up and rolled out in countries (see also [Box 4.3](#)), including for PPM approaches, contributions to case notifications from the private sector and from the currently unengaged parts of the public sector will continue to increase.

Inventory studies that quantify the underreporting of detected TB cases in both public and private sectors can help to identify where further PPM efforts are needed. An excellent recent example, from Indonesia, is highlighted in [Box 4.4](#).

Electronic systems for online reporting of a limited number of variables can facilitate more complete reporting of cases from the private sector. Recent examples include Bangladesh and Pakistan.

FIG. B4.2.1

Contribution of public–public mix to TB case notifications in eight countries, 2012–2017

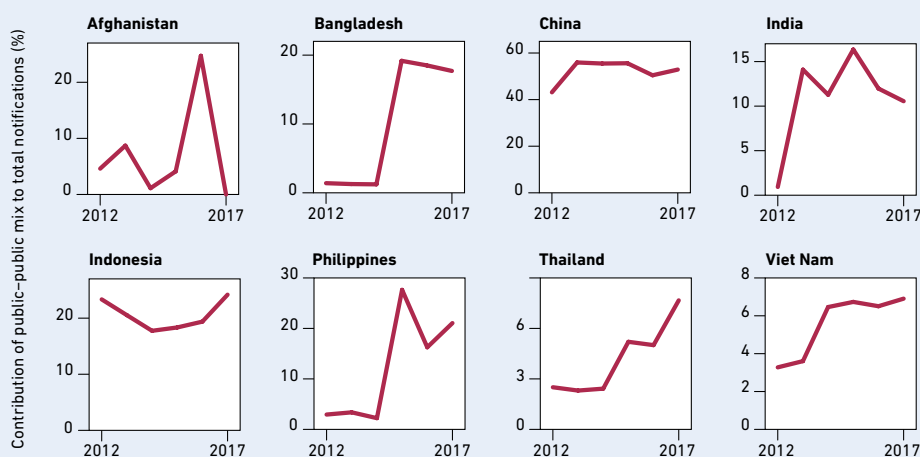


FIG. B4.2.2

Contribution of public–private mix to TB case notifications in eight countries, 2012–2017

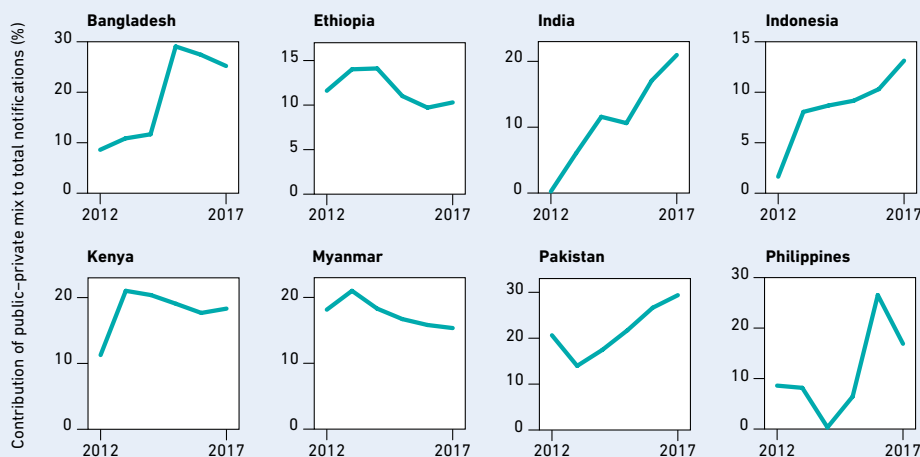
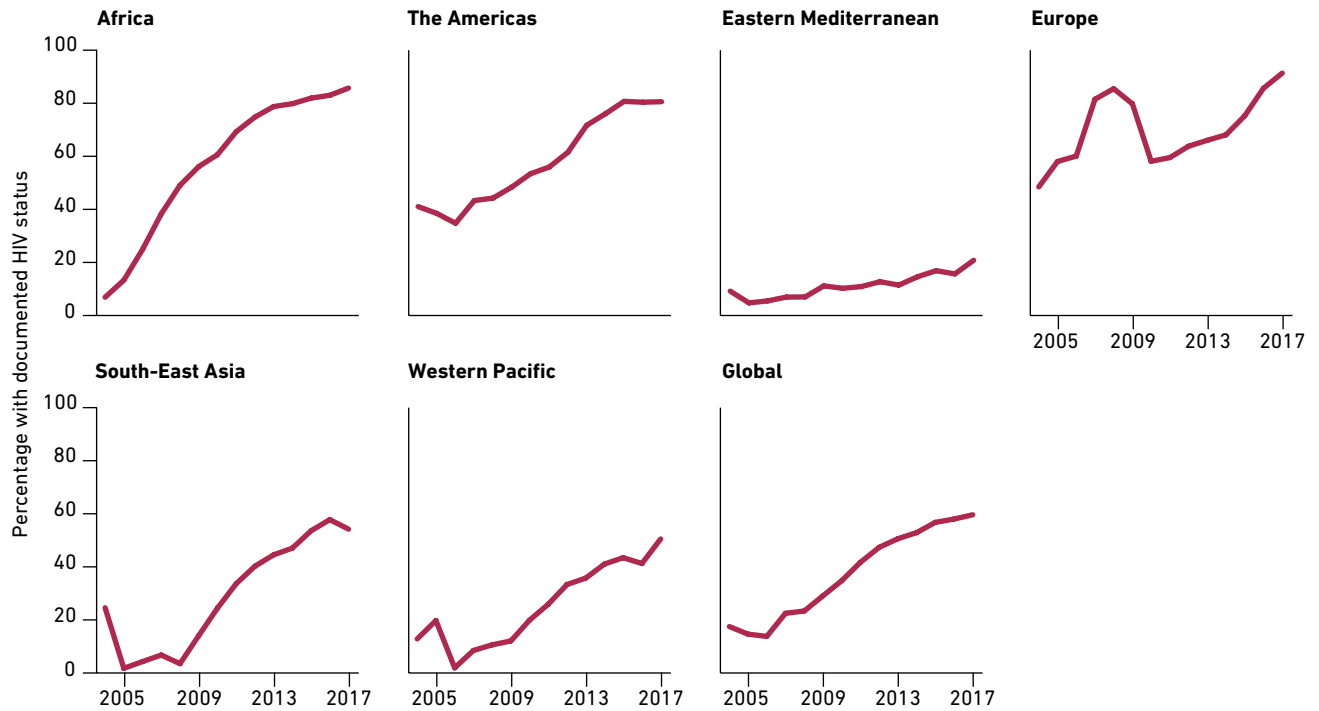


FIG. 4.7

Percentage of new and relapse^a TB cases with documented HIV status, 2004–2017, globally and for WHO regions^b

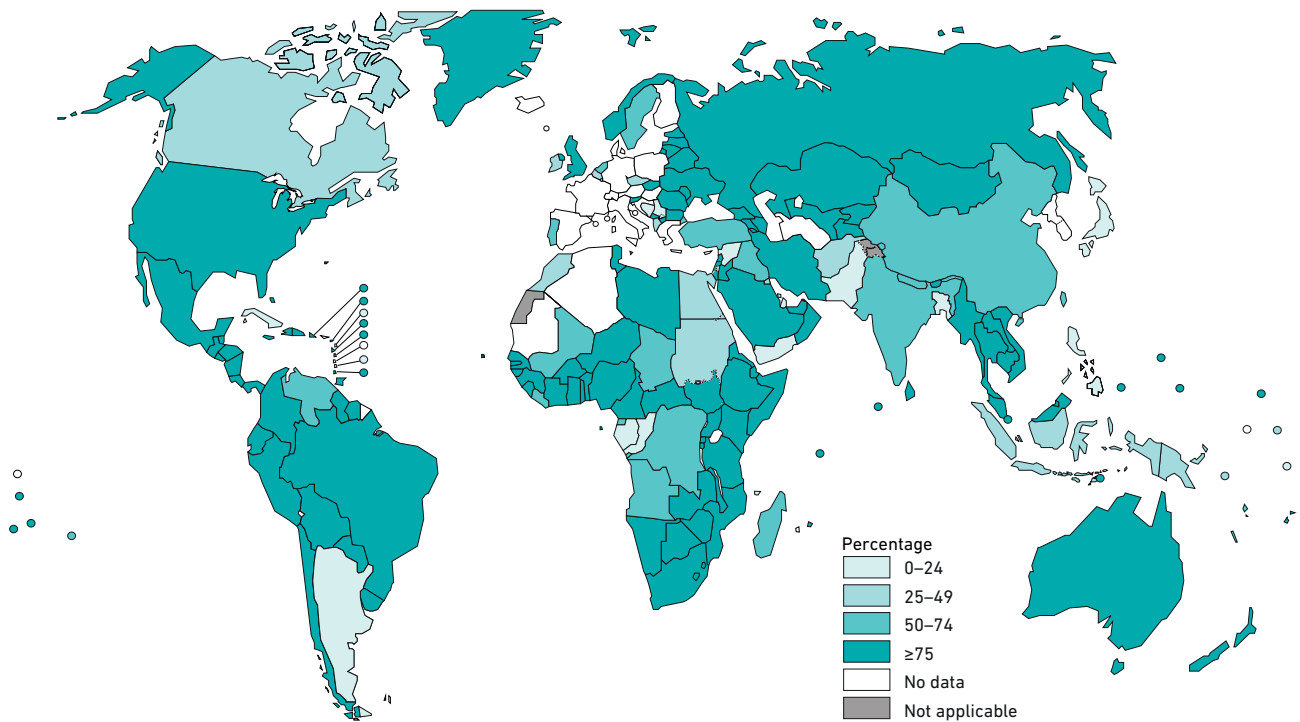


^a The calculation is for all cases in years prior to 2015.

^b Countries were excluded if the number with documented HIV status was not reported to WHO.

FIG. 4.8

Percentage of new and relapse TB cases with documented HIV status, 2017^a



^a 2016 data were used for 15 countries.

HIV status averaged 66% of TB patients in the 30 high TB/HIV burden countries, but varied considerably, from 13% in Congo to above 80% in 16 high TB/HIV burden countries in the WHO African Region.¹ In the WHO African Region, which accounted for 72% of the global burden of HIV-associated TB in 2017 (Chapter 3), 86% of TB patients knew their HIV status.

Globally, 464 633 cases of TB among people living with HIV were notified in 2017 (Table 4.1), equivalent to 12% of TB patients with an HIV test result. The number notified was only 51% of the estimated number of incident cases among people living with HIV (Fig. 4.9),² but an increase from 49% in 2016.

Overall, the percentage of TB patients testing HIV-positive has been falling globally since 2008. This decline is evident in all WHO regions with the exception of the WHO European Region, where the proportion of TB patients testing HIV-positive increased from 3% in 2008 to 13% in 2017.

Systematic symptom screening for TB among people living with HIV is recommended by WHO as an essential component of the HIV care package, together with linkage to diagnostic services, as necessary. In 2017, 92 countries reported annual data on the number of TB cases notified among those newly enrolled in HIV care (up from 90 countries in 2016). In total, 8% of the 1.5 million people who were reported to be newly enrolled in HIV care in 2017 were diagnosed with TB during the same year; data for the 17 high TB/HIV burden countries that reported data are shown in Table 4.2.

4.1.3 Rapid testing for TB

Increasing patient access to early and accurate diagnosis using a WHO-recommended rapid diagnostic (WRD³) is one of the three main objectives of TB laboratory strengthening efforts under the End TB Strategy. As a first step towards reaching this objective, countries should adopt policies that include diagnostic algorithms in which a WRD is the initial diagnostic test for all people with signs or symptoms of TB. They should also adopt such policies as part of working towards the first indicator of the *Framework of indicators and targets for laboratory strengthening under the End TB Strategy*,⁴ which was launched in 2016. Such policies should be an especially high priority for the 48 countries included in one or more of the lists of high TB, TB/HIV and MDR-TB burden

¹ Although the national figure for China was 55%, in the counties defined as having a high burden of HIV-associated TB the figure was 87%.

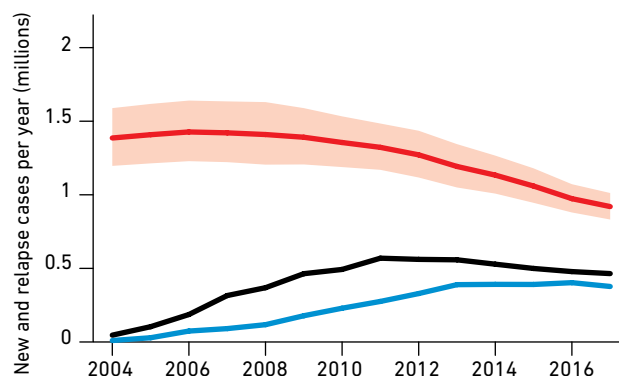
² See also Table 3.2 in Chapter 3 for the global estimate of TB incidence among people living with HIV. The best estimate was 920 000 cases in 2017 (9% of the total number of incident cases).

³ WRDs use molecular techniques to detect TB among people with signs or symptoms of TB. They include the XpertMTB/RIF[®] assay (Cepheid, United States) and the LoopampTM MTBC Detection Kit (Eiken Chemical Company Ltd, Japan).

⁴ World Health Organization. Framework of indicators and targets for laboratory strengthening under the End TB Strategy (WHO/HTM/TB/2016.18). Geneva: WHO; 2016 (<http://www.who.int/tb/publications/labindicators/en/>, accessed 15 August 2018).

FIG. 4.9

Global numbers of notified new and relapse cases^a known to be HIV-positive (black), number started on antiretroviral therapy (blue) and estimated number of incident HIV-positive TB cases (red), 2004–2017. Shaded areas represent uncertainty bands.



^a The calculation is for all cases in years prior to 2015.

TABLE 4.2

Number of people newly enrolled in HIV care who were also notified as a TB case in 2017, 17 high TB/HIV burden countries that reported annual data^a

	NUMBER OF PEOPLE NEWLY ENROLLED IN HIV CARE	NUMBER NOTIFIED AS A TB CASE	NOTIFIED TB CASES AS A PERCENTAGE OF THOSE NEWLY ENROLLED IN HIV CARE
Angola	22 137	2 337	11
Central African Republic	8 180	2 673	33
Congo	29 060	726	2.5
Eswatini	23 271	238	1.0
Ethiopia	39 126	2 375	6.1
Ghana	31 058	1 456	4.7
Guinea-Bissau	2 247	528	23
India	173 174	19 403	11
Indonesia	49 215	10 752	22
Kenya	155 377	8 155	5.2
Malawi	168 972	1 286	0.76
Myanmar	37 402	4 621	12
Namibia	21 213	3 795	18
Nigeria	232 822	19 533	8.4
Papua New Guinea	3 788	482	13
Thailand	11 077	1 436	13
Zambia	199 278	20 323	10
Total	1 207 397	100 119	8.3

^a Zimbabwe reported data for July–December only and was excluded from the table.

countries; of these 48 countries, 32 reported having policies that included such an algorithm by the end of 2017 (Table 4.3). The second indicator of the framework is the percentage of new and relapse TB cases actually tested with a WRD as the initial diagnostic test. Of 190 reporting countries and territories, 133 indicated that their routine surveillance system captures the data required to monitor this indicator. Among the 48 HBCs, 12 countries reported that a WRD had been used as the initial diagnostic test for more than half of their notified TB cases.

The Xpert MTB/RIF® assay (Cepheid, United States) is currently the WRD used most frequently by countries worldwide; it simultaneously detects both TB and resistance to rifampicin. The assay is performed using the GeneXpert® platform, a modular testing device that can detect multiple diseases.¹ Between 2010 (when the test was initially recommended by WHO) and 2017, a cumulative total of 9449 GeneXpert instruments, comprising 42 392 modules and 34.4 million cartridges, were procured by the public sector in 130 of 145 countries eligible for concessional pricing.

4.1.4 Drug susceptibility testing and detection of drug-resistant TB

Drug-resistant TB threatens global TB care and prevention, and it remains a major public health concern in many countries. Three categories are used for global surveillance and treatment: RR-TB, MDR-TB and extensively drug-resistant TB (XDR-TB). MDR-TB is TB that is resistant to both rifampicin and isoniazid, the two most powerful anti-TB drugs; it requires treatment with a second-line regimen. RR-TB also requires treatment with second-line drugs.² With increasing use of Xpert MTB/RIF for simultaneous detection of TB and resistance to rifampicin, a growing number of RR-TB cases (without further testing for isoniazid resistance) are being detected and notified.³ XDR-TB is defined as MDR-TB plus resistance to at least one fluoroquinolone and a second-line injectable agent (amikacin, capreomycin or kanamycin).

The End TB Strategy calls for universal access to drug susceptibility testing (DST).

Drug susceptibility testing for first-line drugs and detection of MDR/RR-TB

Fig. 4.10 shows progress in DST coverage since 2009, when WHO intensified efforts to track progress in the

programmatic response to drug-resistant TB.⁴ In 2017, 2.0 million (30%) of the 6.7 million new and previously treated TB cases notified globally were tested for rifampicin resistance, with coverage of 24% for new TB patients and 70% for previously treated TB patients. These figures represent an improvement since 2015, when 12% of new and 53% of previously treated TB cases had a test result for rifampicin resistance. DST coverage increased in four of the six WHO regions between 2016 and 2017, with a high of 57% in the WHO European Region in 2017. DST coverage varied substantially between countries (even within the same region) and among the 30 high MDR-TB burden countries (Fig. 4.11).

Globally, 160 684 cases of MDR/RR-TB were detected and notified in 2017 (Table 4.1). This was a small increase from 153 119 in 2016 (Fig. 4.12), although aggregate global trends conceal faster progress in some countries (Fig. 4.13). Between 2016 and 2017, the number of reported MDR/RR-TB cases increased by more than 30% in six of the 30 high MDR-TB burden countries (Angola, Democratic People's Republic of Korea, Indonesia, Nigeria, Somalia and Thailand).

The global number of MDR/RR-TB cases notified in 2017 was 29% of the estimated 558 000 incident cases in 2017 (Fig. 4.12; incidence estimates are discussed in more detail in Chapter 3) and 49% of the estimated 330 000 cases of MDR/RR-TB among notified TB cases. Closing these large detection gaps will require improvements in both overall TB detection (Section 4.2) and coverage of diagnostic DST. The latter requires further strengthening of laboratory capacity and wider uptake of new rapid diagnostics.

Drug susceptibility testing for second-line drugs and detection of XDR-TB

Among MDR/RR-TB patients notified in 2017, 50% were tested for resistance to both fluoroquinolones and second-line injectable agents, a considerable increase from the 39% tested in 2016. Coverage varied widely among countries (Fig. 4.14). A total of 10 800 cases of XDR-TB were reported by 77 countries, (up from 8014 cases reported by 72 countries in 2016), with 88% of cases being from the WHO European and South-East Asia regions (Table 4.1). The five countries that reported the largest numbers of cases were Belarus (525), India (2650), the Russian Federation (3661), South Africa (747) and Ukraine (1097).

¹ World Health Organization (2017). Considerations for adoption and use of multidisease testing devices in integrated laboratory networks (WHO/HTM/TB/2017.05). Geneva: WHO (http://www.who.int/tb/publications/2017/considerations_multidisease_testing_devices_2017/en/, accessed 15 August 2018).

² World Health Organization. WHO treatment guidelines for drug-resistant tuberculosis (2016 update) (WHO/HTM/TB/2016.04). Geneva: WHO; 2016 (<http://www.who.int/tb/areas-of-work/drug-resistant-tb/treatment/resources/en/>, accessed 3 July 2018).

³ Surveillance and survey data show that about 82% of RR-TB cases have MDR-TB. Further details are provided in Chapter 3.

⁴ This happened following a ministerial conference for high MDR-TB burden countries, held in Beijing, China, in April 2009. The conference preceded a World Health Assembly resolution the following month: WHA62.15: Prevention and control of multidrug-resistant tuberculosis and extensively drug-resistant tuberculosis. Sixty-second World Health Assembly, Geneva, 18–22 May 2009, Resolutions and decisions; annexes; 2009 (http://apps.who.int/gb/ebwha/pdf_files/WHA62-REC1/WHA62_REC1-en.pdf, accessed 15 August 2018).

TABLE 4.3

National policies and their implementation to increase access to rapid TB testing and universal DST,^a 2017

	■ YES ■ NO	HIGH TB BURDEN	HIGH TB/HIV BURDEN	HIGH MDR-TB BURDEN	NATIONAL POLICY AND ALGORITHM INDICATE A WRD AS THE INITIAL DIAGNOSTIC TEST FOR ALL PEOPLE PRESUMED TO HAVE TB	PERCENTAGE OF NOTIFIED NEW AND RELAPSE TB CASES TESTED WITH A WRD AS THE INITIAL DIAGNOSTIC TEST	NATIONAL POLICY AND ALGORITHM INDICATE UNIVERSAL ACCESS TO DST	PERCENTAGE OF NOTIFIED BACTERIOLOGICALLY CONFIRMED TB CASES WITH DST RESULTS FOR RIFAMPICIN ^b	PERCENTAGE OF NOTIFIED RIFAMPICIN-RESISTANT TB CASES WITH DST RESULTS FOR FLUOROQUINOLONES AND SECOND-LINE INJECTABLE AGENTS
Angola	■	■	■	■	■	—	■	1.7	0
Azerbaijan	□	□	■	■	■	55	■	93	61
Bangladesh	■	□	■	□	□	0.25	□	32	38
Belarus	□	□	■	■	■	88	■	100	97
Botswana	□	■	□	■	■	—	■	3.5	43
Brazil	■	■	□	■	■	26	■	46	16
Cambodia	■	□	□	□	□	—	□	4.6	74
Cameroon	□	■	□	■	■	70	□	28	65
Central African Republic	■	■	□	□	□	2.8	■	5.0	2.2
Chad	□	■	□	□	□	9.9	□	21	1.7
China	■	■	■	□	□	—	■	45	—
Congo	■	■	□	□	□	5.3	□	12	0
DPR Korea	■	□	■	□	□	—	□	4.5	0
DR Congo	■	■	■	□	□	—	□	10	16
Eswatini	□	■	□	■	■	86	■	98	85
Ethiopia	■	■	■	□	□	—	■	—	30
Ghana	□	■	□	■	■	5.8	■	99	0
Guinea-Bissau	□	■	□	□	□	—	□	1.7	0
India	■	■	■	■	■	40	■	70	69
Indonesia	■	■	■	■	■	2.4	■	51	36
Kazakhstan	□	□	■	■	■	82	■	>100	—
Kenya	■	■	■	■	■	47	■	79	51
Kyrgyzstan	□	□	■	■	■	65	■	80	51
Lesotho	■	■	□	■	■	—	■	89	—
Liberia	■	■	□	■	■	17	■	100	22
Malawi	□	■	□	□	□	—	■	—	0
Mozambique	■	■	■	■	■	—	■	—	24
Myanmar	■	■	■	□	□	—	□	78	5.0
Namibia	■	■	□	■	■	100	■	—	19
Nigeria	■	■	■	■	■	41	■	55	30
Pakistan	■	□	■	□	□	2.6	■	32	83
Papua New Guinea	■	■	■	■	■	—	■	72	45
Peru	□	□	■	■	■	—	■	>100	55
Philippines	■	□	■	□	□	26	□	61	37
Republic of Moldova	□	□	■	■	■	93	■	83	74
Russian Federation	■	□	■	■	■	78	■	73	87
Sierra Leone	■	□	□	■	■	2.9	□	—	63
Somalia	□	□	■	■	■	12	■	81	40
South Africa	■	■	■	■	■	66	■	>100	—
Tajikistan	□	□	■	■	■	52	■	89	55
Thailand	■	■	■	■	■	12	■	52	20
Uganda	□	■	□	■	■	37	■	58	15
Ukraine	□	□	■	□	□	—	■	89	100
UR Tanzania	■	■	□	■	■	14	□	39	22
Uzbekistan	□	□	■	■	■	67	■	>100	80
Viet Nam	■	□	■	□	□	26	■	62	48
Zambia	■	■	□	■	■	28	■	37	0
Zimbabwe	■	■	■	■	■	—	■	—	—

Blank cells indicate data not reported. — indicates value that cannot be calculated.

^a The 48 countries shown in the table are the countries that are in one or more of the three lists of high TB, TB/HIV and MDR-TB burden countries (see also Chapter 2, Figure 2.5 and Table 2.4).

^b Testing in cases with unknown previous treatment history is not included. The percentage may exceed 100% for several reasons, e.g. samples rather than cases are counted in the numerator; laboratory specimen results are not linked to the denominator data source when enumerated; or there is incomplete reporting of bacteriologically confirmed cases in the denominator. Bacteriologically confirmed extrapulmonary cases are not included in the denominator because they cannot be differentiated from clinically diagnosed ones in the way data are reported to WHO.

FIG. 4.10

Percentage of all TB cases tested for RR-TB, globally and for WHO regions, 2009–2017

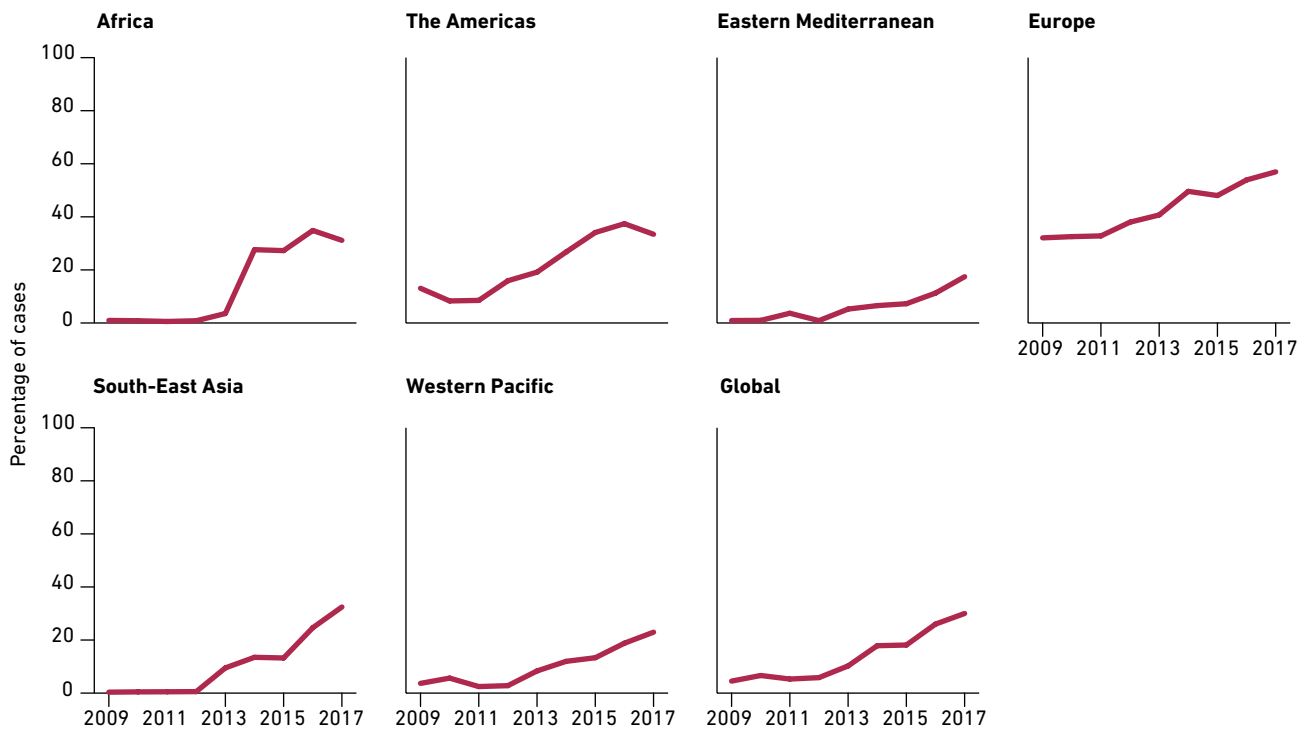
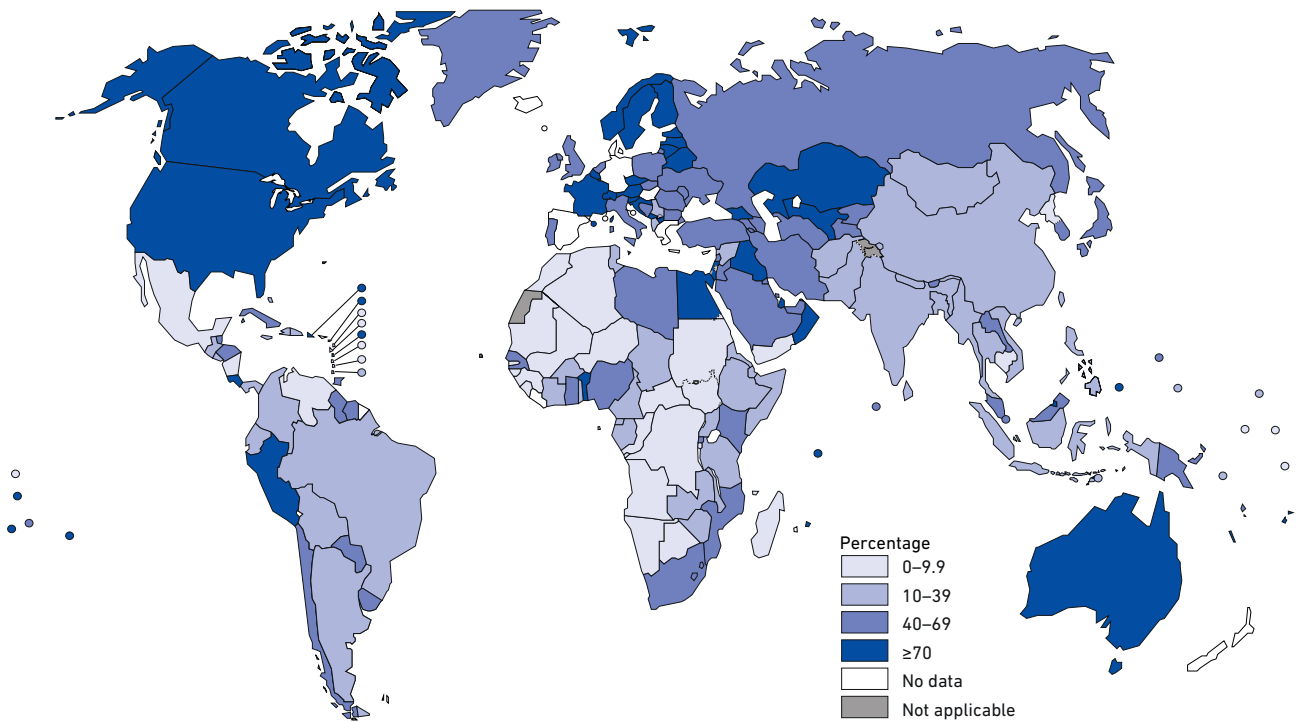


FIG. 4.11

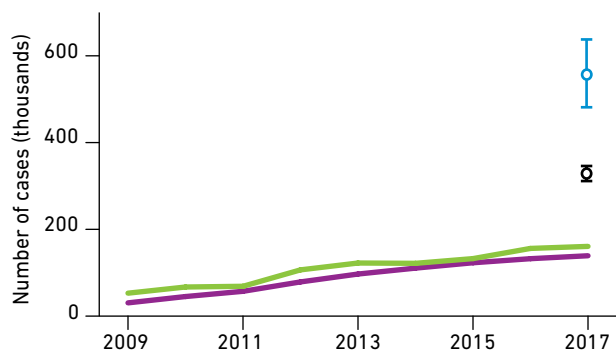
Percentage of all TB cases tested for RR-TB, 2017^a



^a 2016 data were used for 18 countries.

FIG. 4.12

Global number of MDR/RR-TB cases detected (green) and number enrolled on MDR-TB treatment (purple), 2009–2017, compared with estimate for 2017 of the number of incident cases of MDR/RR-TB (uncertainty interval shown in blue) and the number of MDR/RR-TB cases among notified pulmonary cases (uncertainty interval shown in black)



4.1.5 Electronic, case-based surveillance for TB

A growing number of countries are capturing data for notified TB cases in electronic case-based surveillance systems. These systems have several advantages compared with more traditional paper-based reporting of aggregated data, including enabling more timely access to data (up to “real-time”) and the availability of data at the level of individual patients up to national level. They also greatly facilitate data analysis (including by age, sex and location) to inform adaptation and targeting of response efforts, both geographically and for specific population groups. Further details, including of global efforts to support the adoption of case-based surveillance for TB, are provided in [Box 4.3](#).

4.2 Treatment coverage

The Sustainable Development Goals (SDGs) include a target to “Achieve universal health coverage, including financial risk protection, access to quality essential health-care services and access to safe, effective, quality and affordable essential medicines and vaccines for all” ([Chapter 2](#)). One of the indicators for Target 3.8 of SDG 3 is the coverage of essential health services; this is a composite indicator based on 16 tracer indicators, one of which is TB treatment. Achieving UHC is a fundamental requirement for achieving the milestones and targets of the End TB Strategy; hence, priority indicators for monitoring progress in implementing the End TB Strategy include both TB treatment coverage and the percentage of TB patients and their households that face catastrophic costs as a result of TB disease ([Chapter 2](#)).

TB treatment coverage is defined as the number of new and relapse cases detected and treated in a given year, divided by the estimated number of incident TB

cases in the same year, expressed as a percentage. In this section, numbers of notified new and relapse cases in 2017 are used as the numerator for the indicator, because these are the data that are available. However, as discussed further below, there are people with TB who are treated but not notified to national authorities (and in turn are not notified to WHO), and people who are notified but who may not be started on treatment.

ART is recommended for all HIV-positive TB patients, and a second-line MDR-TB treatment regimen is recommended for people with MDR/RR-TB. This section includes estimates of treatment coverage for these two interventions as well.

4.2.1 TB treatment coverage

Trends in notifications of new and relapse cases and estimated incidence are shown for the 30 high TB burden countries in [Fig. 4.15](#). Estimates of TB treatment coverage in 2017 (calculated as notifications of new and relapse cases divided by estimated TB incidence) are shown globally, for WHO regions and the 30 high TB burden countries, in [Fig. 4.16](#).

Globally, TB treatment coverage was 64% (range, 58–72%) in 2017, up from 53% (range, 46–64%) in 2010 and 35% (range, 30–43%) in 2000. Three WHO regions achieved levels above 75%: the WHO Region of the Americas, the WHO European Region and the WHO Western Pacific Region. High TB burden countries with high levels of treatment coverage in 2017 (>80%) included Brazil, China, Namibia, the Russian Federation and Viet Nam. The lowest levels, with best estimates of 50% or less, were in Central African Republic, Lesotho, Nigeria and the United Republic of Tanzania.

Globally in 2017, there was a gap of about 3.6 million cases between the 6.4 million new and relapse cases that were notified, and the estimated 10 million incident TB cases in the same year ([Fig. 4.1](#)). The global gap has been narrowing, especially in the WHO Eastern Mediterranean Region and the WHO Western Pacific Region, and to a lesser extent in the WHO South-East Asia Region.¹ Ten countries account for 80% of the total estimated global gap between incidence and notifications ([Fig. 4.17](#)), with India (26%), Indonesia (11%), Nigeria (9%) and the Philippines (7%) accounting for more than half of the global total.

There are three main reasons for a gap between notifications and estimated incidence:

- **Underreporting of detected TB cases.** In many countries, levels of underreporting may be high; this is especially the case for those countries that lack policies on mandatory notification and other measures to ensure reporting of detected cases by all care providers and large private health sectors.

¹ Time trends in countries and regions are shown in [Annex 2](#) and [Annex 3](#), respectively.

FIG. 4.13

Number of MDR/RR-TB cases detected (green) and enrolled on MDR-TB treatment (purple), 2009–2017, 30 high MDR-TB burden countries

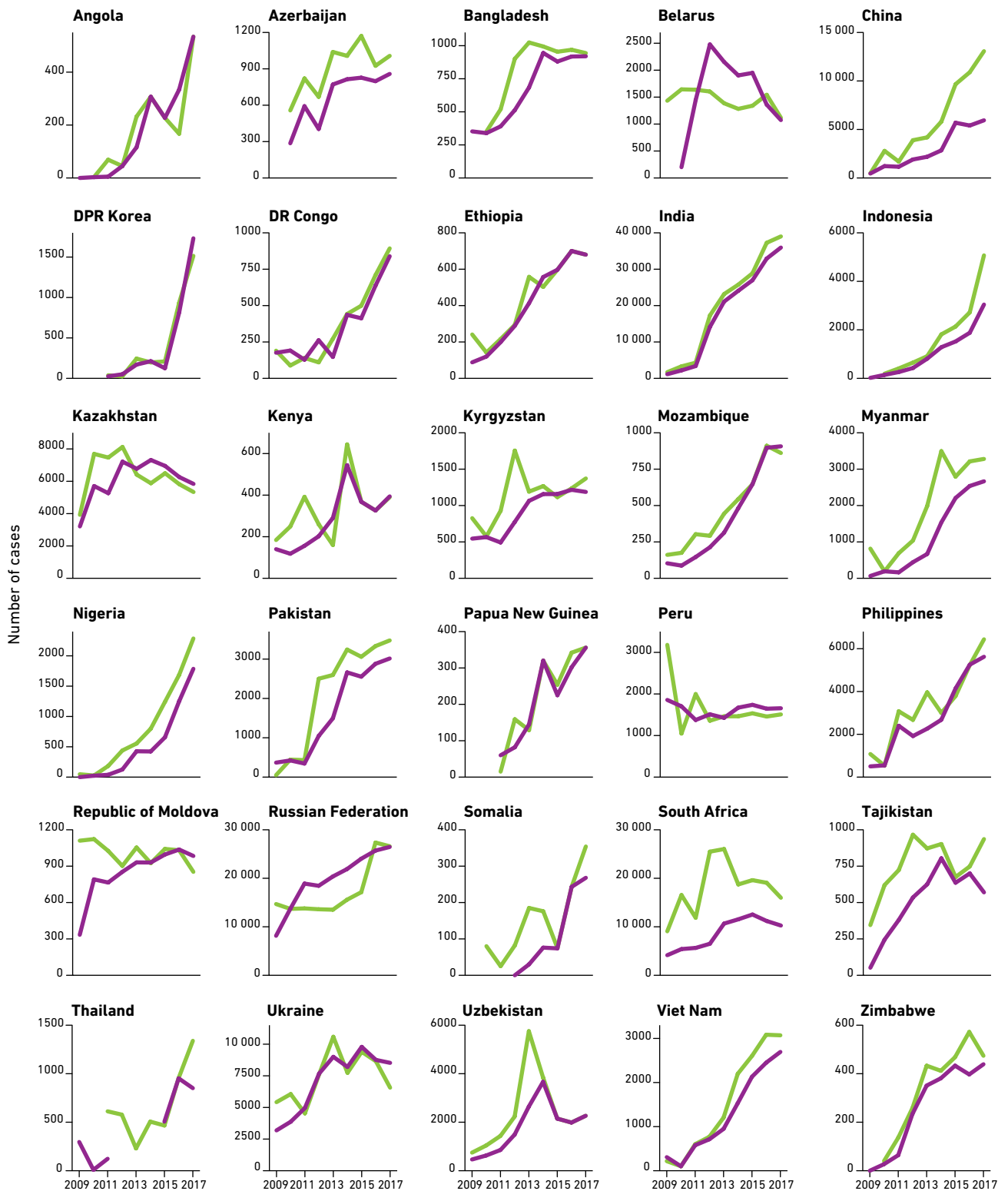
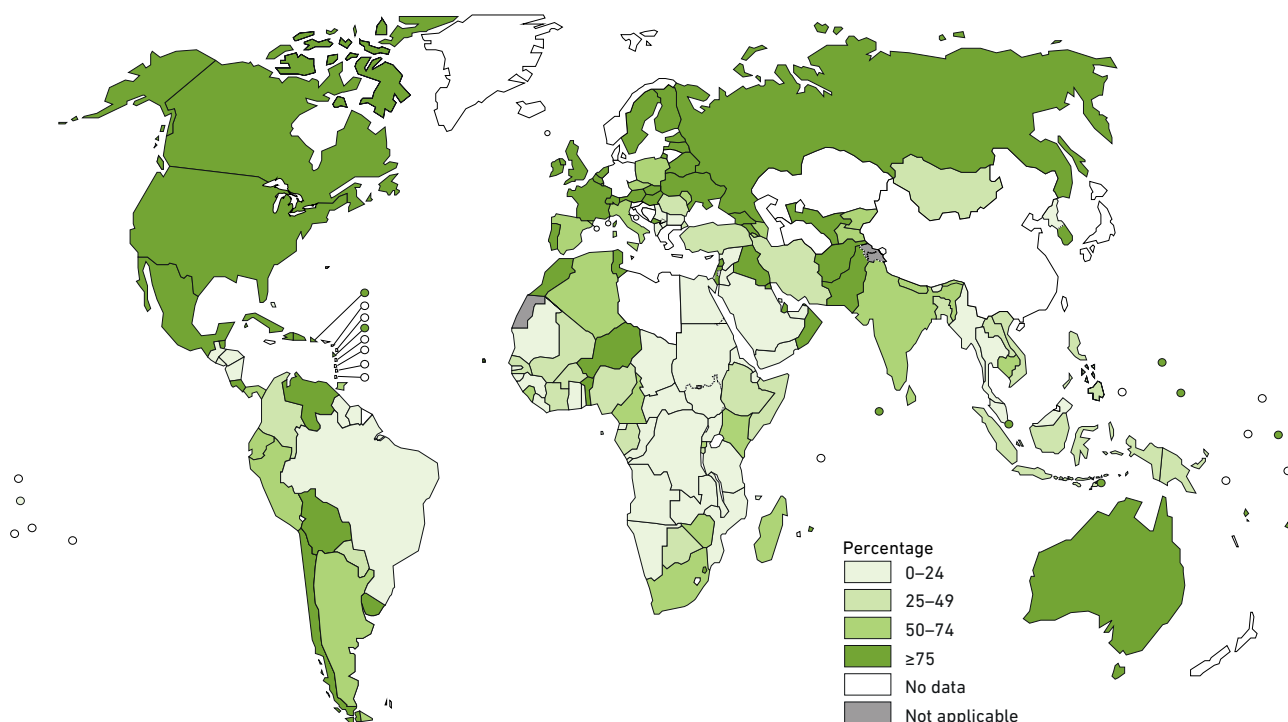


FIG. 4.14

Percentage of MDR/RR-TB cases tested for susceptibility to second-line drugs, 2017^a



^a 2016 data were used for 18 countries.

- **Underdiagnosis of people with TB.** Underdiagnosis can occur for reasons such as poor geographical and financial access to health care; lack of or limited symptoms that delay seeking of health care; failure to test for TB when people do present to health facilities; and diagnostic tests that are not sufficiently sensitive or specific to ensure accurate identification of all cases.
- **Overestimation of the level of TB incidence.** In this report, estimates of TB incidence for 44 countries with 19% of the world's estimated cases are based on expert opinion about levels of underreporting and underdiagnosis, as opposed to direct measurements from surveillance or survey data (Chapter 3). Also, the uncertainty intervals around the best estimates of TB incidence can be wide, and gaps may be lower or higher than the best estimates quoted in this section.

In some of the countries with the largest estimated gaps between notifications and TB incidence, there is already good evidence about the reasons for such gaps, and actions to address them are being taken or are planned.

An excellent recent example is Indonesia. The 2013–2014 national TB prevalence survey showed high levels of underreporting of detected TB cases,¹ leading to recommendations such as a mandatory policy on notification (enacted in January 2017), and intensified en-

¹ This confirmed earlier data on extensive sales of drugs in the private sector.

gagement with public and private hospitals where many people with TB were being treated.² The results from the national TB prevalence survey also led to a decision to implement a national inventory study to measure the level of underreporting of detected TB cases. This was implemented in the first quarter of 2017, and the study proved to be large and robust enough to allow the use of capture–recapture analysis to reassess estimates of incidence.³ The key study results were that 41% of detected cases were not reported to the national surveillance system, and that about 20% of incident cases were not diagnosed. In other words, although about 80% of incident cases were estimated to be detected, only 59% of these detected cases were officially reported. Based on capture–recapture analysis (see also Chapter 3),

² For further details, see Box 2.4 in World Health Organization. Global tuberculosis report 2015 (WHO/HTM/TB/2015.22). Geneva: WHO; 2015 (http://apps.who.int/iris/bitstream/10665/191102/1/9789241565059_eng.pdf, accessed 21 June 2018).

³ Capture–recapture modelling is possible if six assumptions are met: (i) all cases should be observable; (ii) the proportion of mismatches and matching failures in record-linkage is low, which typically requires a large sampling fraction; (iii) a closed population during the study period (typically 3–6 months); (iv) if S represents the number of case lists or data sources available, then at least three data sources should be available ($S \geq 3$) and their dependencies must be accounted for in the model design, while the full S-way interaction between sources is assumed null; (v) homogeneity of within-source observation probabilities across subpopulation groups, such as those defined by socioeconomic and demographic characteristics; (vi) consistent case definitions across data sources. Few high TB burden countries are expected to be able to implement inventory studies that will meet these 6 assumptions to a sufficient degree.

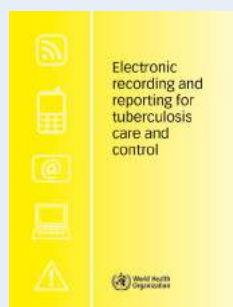
BOX 4.3

National case-based electronic surveillance systems for TB: status of progress, global efforts to support expansion, broader context

Case-based electronic surveillance systems for TB with national coverage have several advantages over more traditional paper-based reporting of aggregated data. For example, they enable more timely access to data (up to “real-time”) and the availability of data at the level of individual patients up to national level;

also, they greatly facilitate data analysis (including by age, sex and location) to inform adaptation and targeting of response efforts, both geographically and for specific population groups.

WHO has promoted case-based electronic surveillance for TB for several years, with guidance issued in 2012.^a



Status of progress in national case-based electronic surveillance for TB, August 2018

In August 2018, data on the type of TB surveillance system in place at national level were available for 206 countries (Fig. B4.3.1). Of these, 136 had a case-based surveillance system that covered all TB cases (both drug-susceptible and drug-resistant TB). These countries accounted for 67% of global TB notifications in 2017.

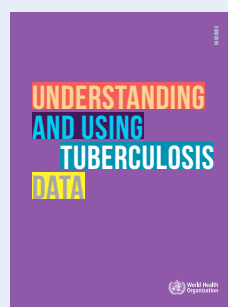
A further 24 countries, mainly in the WHO African Region and the WHO South-East Asia Region, had a case-based surveillance system for all cases of drug-resistant TB. These countries are in a transition phase between aggregate paper-based reporting and case-based electronic surveillance. The initial prioritization of MDR-TB is explained by the additional complexity of monitoring treatment and treatment outcomes compared with drug-susceptible TB, which is much easier to manage with case-based surveillance; and by the fact that

often the numbers of treatment centres and laboratories that need to be involved are smaller, making introduction more feasible from a logistics perspective. About half of the countries in the WHO African Region still have paper-based systems for recording and reporting of data.

Global efforts to support further expansion of national case-based electronic surveillance for TB

Building on the WHO guidance issued in 2012,^a WHO's Global TB Programme has been working with other WHO departments and the University of Oslo to develop modules in the DHIS2 software^b for electronic management of data in both aggregated format (as an interim step for countries not yet ready to transition to case-based surveillance) and case-based data. Both modules are based on the WHO recording and reporting framework^c and both allow extensive data analysis at different levels of the health system. To facilitate the routine analysis and use of TB data, standard dashboards have been developed as part of these DHIS2 modules; these

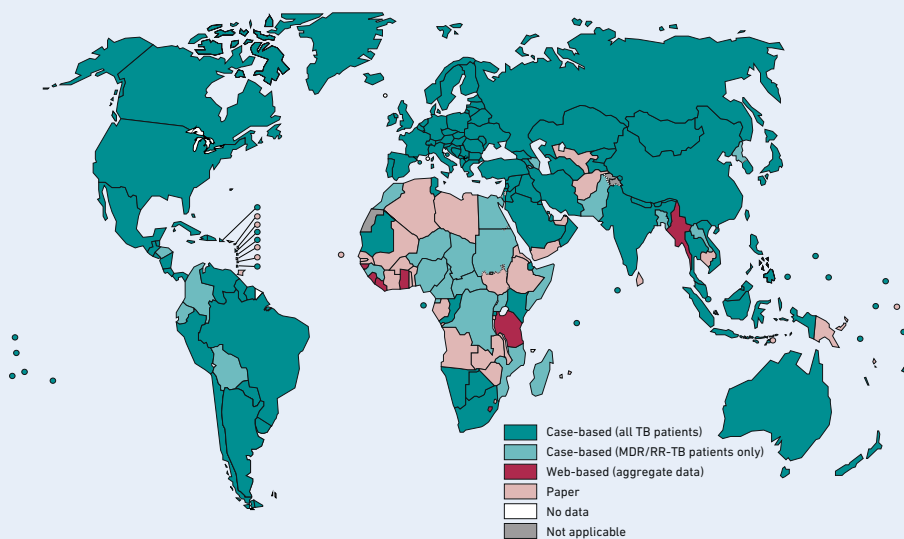
include graphs, tables and maps for core surveillance (e.g. notifications, coverage of testing for drug resistance and HIV, and treatment outcomes) indicators and data quality indicators (e.g. completeness and internal consistency), based on WHO guidance on analysis and use of TB data,^d and the WHO TB surveillance checklist of standards and benchmarks.^e



The modules were developed in DHIS2 because many countries are already using this software within their health information systems, it is open-source software with no license fees, and it is being supported by a wide range of technical and funding partners.

FIG. B4.3.1

National TB surveillance systems in place in August 2018



WHO and the University of Oslo have already supported six countries to implement, optimize and use their own TB module for aggregated data in DHIS2: Ghana, Guinea-Bissau, Liberia, Myanmar, Uganda and the United Republic of Tanzania. An additional 24 countries are being supported to use the DHIS2 module for aggregated data. By August 2018, aggregated data down to the regional or district level for at least five years had been compiled and stored in a DHIS2 TB platform developed by WHO^f for 44 countries (Fig. B4.3.2), with more limited data available for an additional four countries. These data can be easily transferred into national DHIS2 databases once these have been adopted and implemented.

Data source: WHO global TB database 2018, supplemented with information provided during WHO workshops on analysis and use of TB data and national TB epidemiological reviews.

A DHIS2 TB module for case-based data that enables electronic management of data for both drug-susceptible and drug-resistant TB in one system is currently in the pilot-testing phase in the Lao People's Democratic Republic.

Broader context: the WHO Global Task Force on TB Impact Measurement and the Health Data Collaborative

Efforts to expand national case-based electronic surveillance systems for TB are part of wider efforts to support the strengthening of national health information systems, and the associated analysis and use of data.

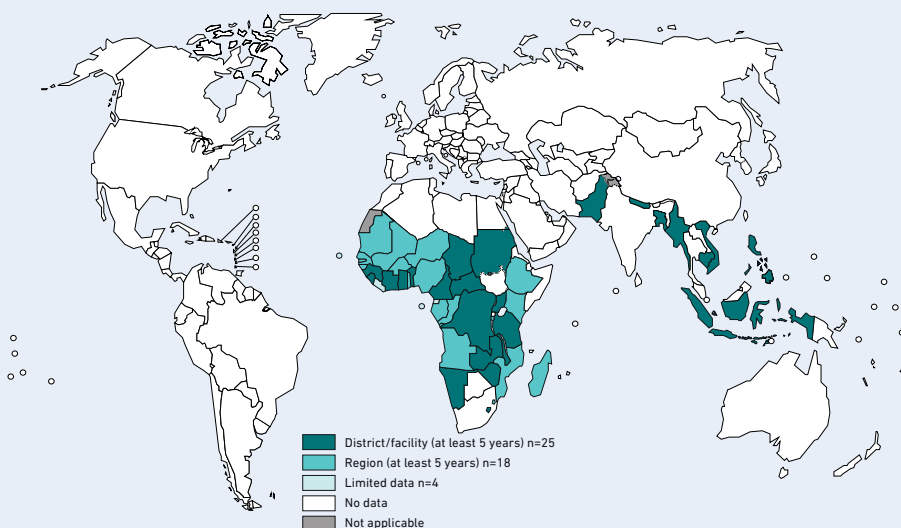
Two of the five strategic areas of work of the WHO Global Task Force on TB Impact Measurement are strengthening national notification systems for direct measurement of TB incidence (within which one of the priorities is case-based electronic surveillance), and analysis and use of TB-relevant data at country level (for further details, see [Box 3.1](#) in [Chapter 3](#)).

The Health Data Collaborative (HDC)⁹ is a joint effort of multiple global health partners to work alongside countries to improve the availability, quality and use of data for local decision-making and the tracking of progress towards the health-related SDGs. The Secretariat sits with the Department of Information, Evidence and Research (IER) at WHO headquarters. The role of the HDC is to build on existing efforts by establishing a network of working groups that will address specific technical issues and identify and fill technical gaps.

Under the umbrella of the HDC, WHO's Global TB Programme is working with IER as well as the Global HIV and Hepatitis Programme, the Global Malaria Programme, the Department of Immunisation and the University of Oslo on a 3-year workplan (2017–2020). The workplan covers three major topics:

- Strengthening country health information systems (in general and for specific diseases). This includes support for the development and management of Health Management Information Systems (HMIS) and DHIS2 modules. Examples include WHO health data apps for DHIS2 (cross-cutting and disease specific), which include standard indicators, dashboards and outputs, and capacity-building for case-based surveillance.
- Strengthening analysis and use of the data generated by country information systems. Examples include a curriculum for routine analysis and use of health facility data (curriculum content on TB has already been

FIG. B4.3.2
Subnational, aggregated data stored in the DHIS2 platform developed by WHO (status in August 2018)



developed), national epidemiological reviews, and national and regional workshops.

- Ensuring quality in the data generated by country health information systems and in the analysis and use of data. Examples include guidance documents and tools; reviews of plans and investments related to monitoring and evaluation; establishment of a pool of trained consultants, national professional officers and in-country staff and institutions to provide technical assistance; and peer review of the quality of tools, plans, investments and reviews.

^a World Health Organization. Electronic recording and reporting for tuberculosis care and control. Geneva: WHO; 2012 (http://www.who.int/tb/publications/electronic_recording_reporting/en/, accessed 25 July 2018).

^b <https://www.dhis2.org/overview>

^c World Health Organization. Definitions and reporting framework for tuberculosis – 2013 revision (updated December 2014) (WHO/HTM/TB/2013.2). Geneva: WHO; 2013 (www.who.int/iris/bitstream/10665/79199/1/9789241505345_eng.pdf, accessed 21 June 2018).

^d World Health Organization. Understanding and using TB data: a handbook (WHO/HTM/TB/2014.09). Geneva: WHO; 2014 (http://www.who.int/tb/publications/understanding_and_using_tb_data/en/, accessed 24 August 2016).

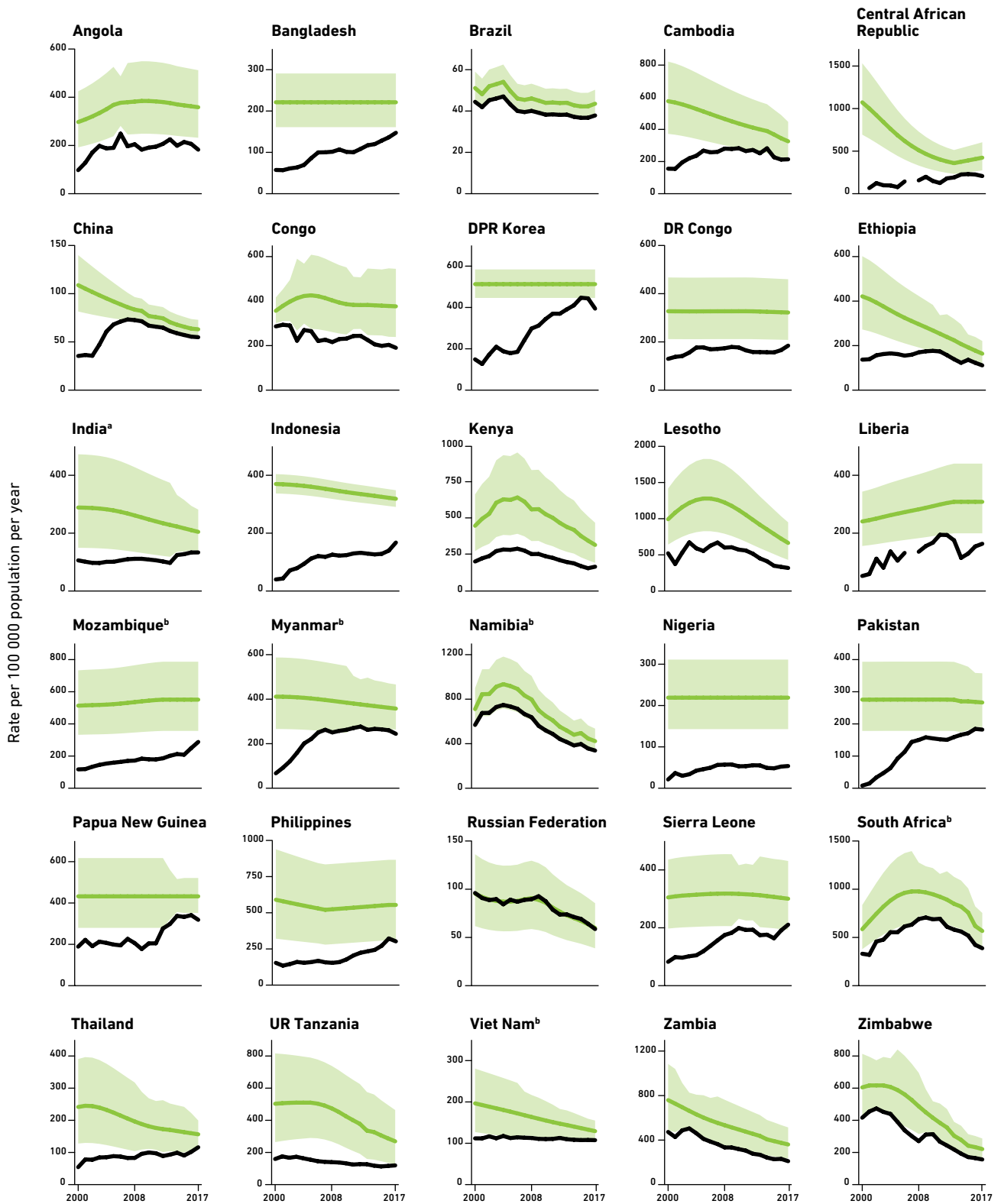
^e World Health Organization. Standards and benchmarks for tuberculosis surveillance and vital registration systems: checklist and user guide. Geneva: WHO; 2014 (<http://www.who.int/tb/publications/standardsandbenchmarks/en/>, accessed 25 July 2018).

^f <https://tbhistorical.org>

^g <https://www.healthdatacollaborative.org>

FIG. 4.15

Case notification rates (new and relapse cases, all forms) (black) compared with estimated TB incidence rates (green), 2000–2017, 30 high TB burden countries. Shaded areas represent uncertainty bands.

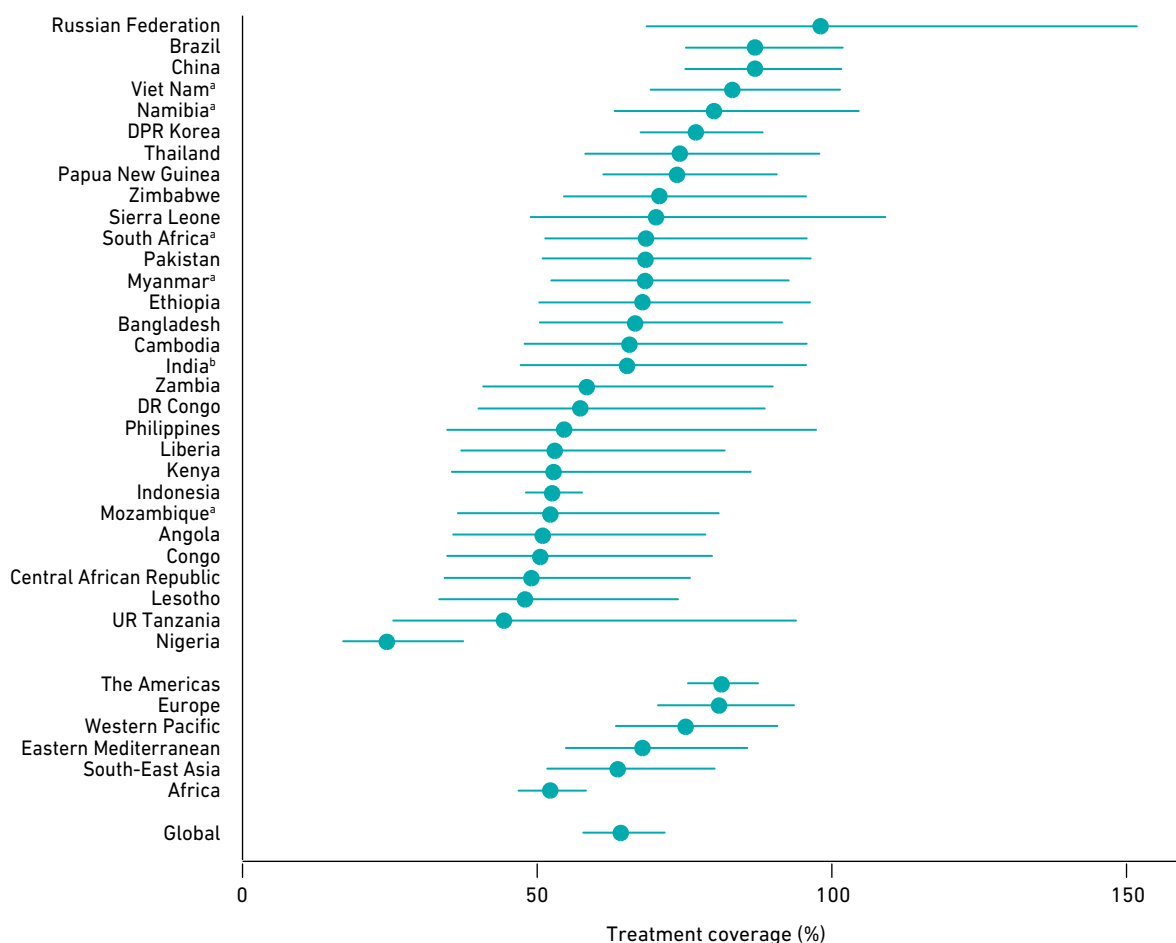


^a Estimates of TB incidence for India are interim, pending results from the national TB prevalence survey planned for 2019/2020.

^b Estimates of TB incidence for Mozambique, Myanmar, Namibia, South Africa and Viet Nam will be reviewed after final results from their respective national TB prevalence surveys are available in 2019.

FIG. 4.16

Estimated TB treatment coverage (new and relapse patients as a percentage of estimated TB incidence) in 2017, 30 high TB burden countries, WHO regions and globally



^a Estimates of TB incidence for Mozambique, Myanmar, Namibia, South Africa and Viet Nam will be reviewed after final results from their respective national TB prevalence surveys are available in 2019.

^b Estimates of TB incidence for India are interim in nature, pending results from the national TB prevalence survey planned for 2019/2020.

estimates of incidence were revised to a best estimate of 842 000 cases (range, 767 000–919 000) in 2017 (less than the previous best estimate of around 1 million, although still consistent with the range that was estimated based on the prevalence survey).¹ Actions are now being taken to correct the high level of underreporting, with results already evident from the notifications reported for the whole of 2017 (Fig. 4.15). A fuller description of the study, including methods, results and lessons learned, is provided in Box 4.4.

In India, multiple sources of evidence from surveys and surveillance have shown large underreporting of detected TB cases, especially in the private sector.² Three examples of actions that have been taken to close reporting gaps are mandatory notification, a simplified national electronic reporting system that facilitates reporting of

cases, and further efforts to engage all care providers through PPM schemes, all of which have already had an effect on notification trends (Section 4.1.1; Fig. 4.15).

A good example of a country where underdiagnosis is a major challenge is Nigeria. The 2012 prevalence survey in Nigeria found that 75% of the smear-positive cases detected had symptoms that met national screening criteria but had not been previously diagnosed, demonstrating high levels of underdiagnosis and a need to strengthen access to screening, diagnostic and treatment services of high quality.³

In countries where underreporting is thought to exist, inventory studies in which electronic lists of notified cases are compared with electronic lists of TB cases detected by all care providers, ideally employing unique identifiers, can be used to quantify levels of underreport-

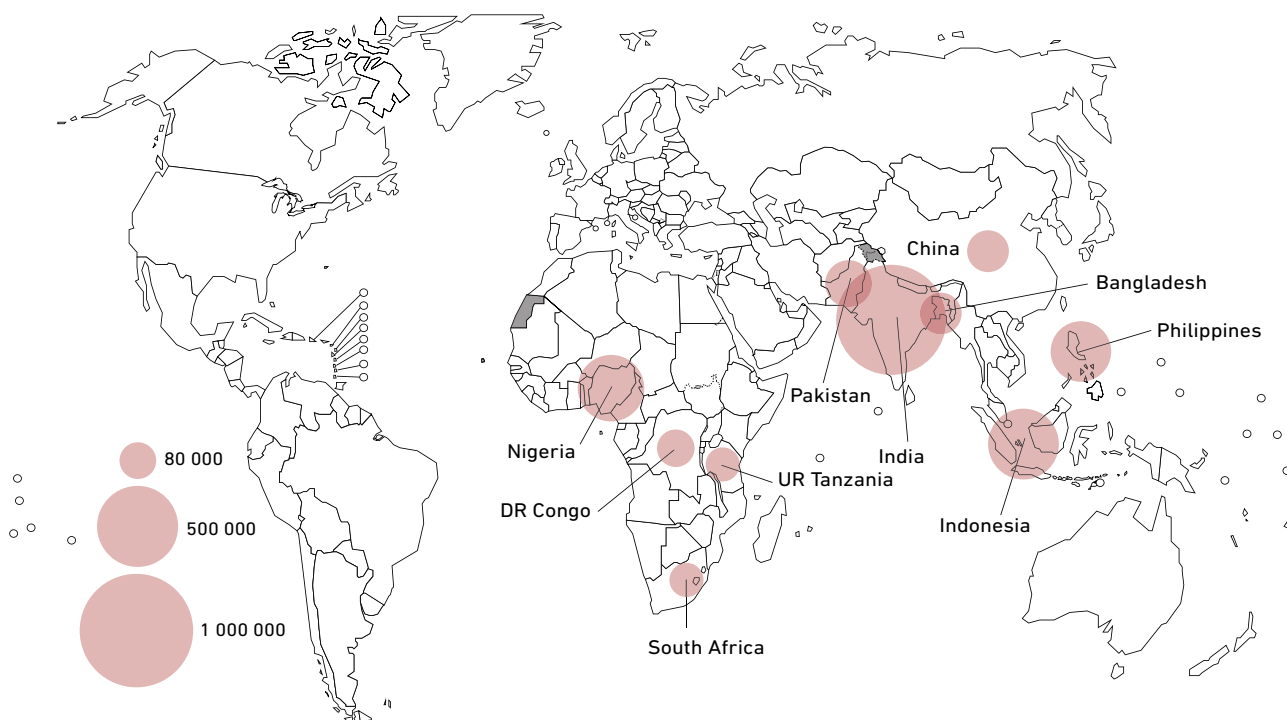
¹ See also Box 3.2 in Chapter 3.

² For further details, see Box 3.3 in World Health Organization. Global tuberculosis report 2016 (WHO/HTM/TB/2016.13). Geneva: WHO; 2016 (<http://apps.who.int/iris/bitstream/10665/250441/1/9789241565394-eng.pdf>, accessed 11 July 2018).

³ For further details, see Box 2.2 in World Health Organization. Global tuberculosis report 2014 (WHO/HTM/TB/2014.08). Geneva: WHO; 2014 (<http://www.searo.who.int/indonesia/documents/9789241564809/en/>, accessed 15 August 2018).

FIG. 4.17

The ten countries with the largest gaps between notifications of new and relapse (incident) TB cases and the best estimates of TB incidence, 2017^a



^a The ten countries ranked in order of the size of the gap between notified cases and the best estimates of TB incidence in 2017 are India, Indonesia, Nigeria, the Philippines, Pakistan, Bangladesh, China, DR Congo, South Africa and UR Tanzania. Estimates of TB incidence for India are interim in nature, pending results from the national TB prevalence survey planned for 2019/2020. Estimates of TB incidence for South Africa will be reviewed after final results from its national TB prevalence survey is available in 2019.

ing, as in Indonesia.¹ Other high TB burden countries that have already implemented an inventory study are India, Kenya, Pakistan and Viet Nam.² Four more are underway or being planned, in China, the Philippines, South Africa and the United Republic of Tanzania. When this type of study is done prospectively (as opposed to retrospectively, using electronic databases that are already available), the mapping of providers that is required at the beginning can subsequently help with efforts to engage all care providers, including in reporting (Box 4.4).

Examples of mechanisms to ensure reporting of all detected cases include linking reimbursement from health insurance schemes to notification of cases (as is done in the Philippines and the Republic of Korea) and linking the supply of first-line drugs to notification of cases (as is done in Brazil).

Recent national TB prevalence surveys³ have also shown that, in both Africa and Asia, detection and reporting gaps are systematically higher for men than for

women (Chapter 3). This suggests that specific efforts are needed to improve access to TB diagnosis and treatment for men.

Systematic screening for active TB among specific populations can also help to ensure early diagnosis and reduce levels of underdiagnosis. WHO recommends such screening for contacts of bacteriologically confirmed cases, people living with HIV and people exposed to silica dust.^{4,5} Other individuals at risk should be considered for systematic screening based on an assessment of TB epidemiology in each setting. To date, there have been few assessments of the implementation and outcomes of systematic screening in countries that are currently introducing or scaling up systematic screening. However, this is expected to become a more prominent part of national programme monitoring and evaluation efforts in future. Engaging communities can also add value to efforts to improve case detection and patient support (Box 4.5).

¹ For a guide to inventory studies, see World Health Organization. Assessing tuberculosis under-reporting through inventory studies. Geneva: WHO; 2012 (http://www.who.int/tb/publications/inventory_studies/en/, accessed 15 August 2018).

² Results from these studies have been used to inform estimates of TB incidence.

³ See http://www.who.int/tb/advisory_bodies/impact_measurement_taskforce/meetings/tf6_p06_prevalence_surveys_2009_2015.pdf

⁴ World Health Organization. Systematic screening for active tuberculosis: principles and recommendations (WHO/HTM/TB.2013.04). Geneva: WHO; 2013 (<http://www.who.int/tb/tbscreening/en/>, accessed 15 August 2018).

⁵ The data requested as part of WHO's global monitoring focus on screening among people living with HIV and close contacts. Hence, the data requested in WHO's annual round of global TB data collection focus on screening among people living with HIV and close contacts. These data are presented in Chapter 5.

BOX 4.4

The 2017 national TB inventory study in Indonesia: main results, lessons learned, policy and programmatic implications

Indonesia is a country with a large private health sector that is not yet firmly linked to the reporting network of the NTP. Furthermore, some of the secondary and tertiary level facilities of the public health sector do not have functioning and sustained reporting links with the NTP.

In 2016, a total of 360 565 TB cases were notified to national authorities, while the estimated TB incidence was 1 020 000 (95% confidence interval [CI]: 660 000–1 460 000). To address underreporting, a Health Ministerial Decree was enacted in 2017, making notification of TB mandatory nationwide.

A national TB inventory study was implemented in 2017 under the leadership of the NTP and the National Institute of Health Research and Development. The aim was to directly measure the level of underreporting of detected TB cases to the national TB surveillance system (SITT, System Informasi Tuberkulosis Terpadu) maintained by the NTP for the different types of health facilities, and to identify best-practice methods for addressing TB underreporting.

Methods and main results

The study methods from design through implementation, analysis and reporting of results followed the recommendations of the WHO Global Task Force on TB Impact Measurement.^a

The study employed a random, nationally representative sample of 23 districts (out of a total of 514) from across the country, using a sampling design with stratification by urban or rural status and three regions – Sumatera, Java/Bali and “other provinces” (Fig. B4.4.1). The 23 districts accounted for 10% of the national population of 260 million.

In each of the sampled districts, all health-care providers of TB services were mapped, starting from existing but outdated lists of health-care providers, which were then confirmed by study enumerators (Fig. B4.4.2). A total of 4207 health-care providers were enumerated in the 23 districts, of which 1687 were eligible (defined as a facility that reported having diagnosed or treated at least one TB patient in the

previous 3 months). Of the eligible health-care providers, 99% participated in the study – a testament to the success of district-level workshops and meetings with all stakeholders. Pharmacies were not included in the sampling frame of the study.

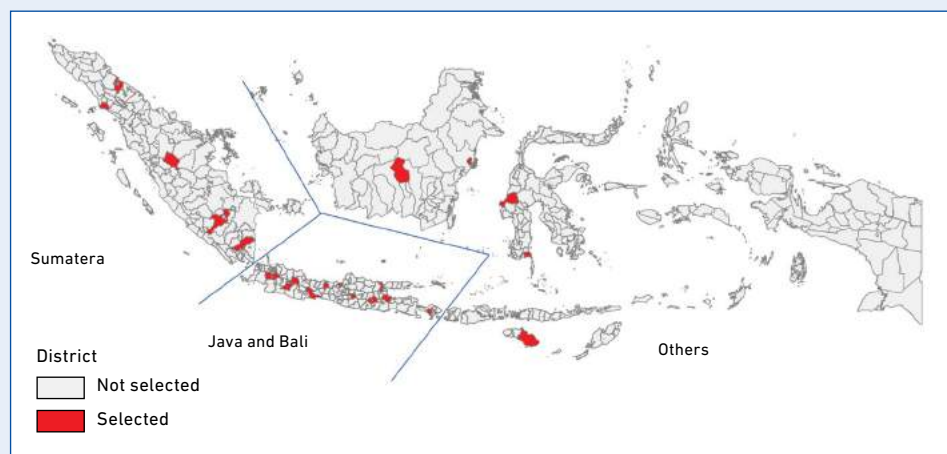
During the study period (the first quarter of 2017), patient records for a total of 21 320 unique TB patients were detected overall. Of these, 13 211 unique TB patients were notified and registered in SITT. Probabilistic matching allowed linkage of records from the TB case lists, measuring their overlap and corresponding underreporting (Fig. B4.4.3).

The overall level of underreporting of detected TB cases (i.e. the proportion of detected cases not in SITT) was estimated to be 41% (95% CI: 36–46%), ranging from 15% (95% CI: 11–20%) underreporting by primary health-care “puskesmas” units (which are part of the existing NTP network) to 65% underreporting by hospitals to 96% underreporting by the combined category of general practitioners (GPs), clinics and laboratories. Patients with clinically diagnosed and extrapulmonary TB, as well as children, were more likely to be underreported (Table B4.4.1).

Incidence was estimated using capture–recapture modelling of the three TB case lists (SITT, study public and study private), according to recommendations (see also section 4.2.1).^a It was estimated that 18% (95% CI: 15–21%) of incident cases were not detected. The annual incidence rate for 2017 was estimated at 319 (95% CI: 290–349) per 100 000 population per year.^b This updated best estimate of TB incidence is lower but statistically consistent with the previously published estimate derived from the 2013/2014 prevalence survey (Fig. B4.4.4).

Study results and updated estimates were discussed and agreed upon in a national consensus meeting held in April 2018.^c The excellent participation rate of health-care providers, in addition to the consistency of key results through sensitivity analyses, show that the study was implemented to a high standard and has produced robust results.

FIG. B4.4.1
Districts selected for the national inventory study



During the course of 2017, the NTP already started to take corrective action to reduce underreporting. This led to a substantial increase in TB notifications in 2017 compared with 2016. While the measure of underreporting in the study during the first quarter of 2017 was 41%, in the subsequent three quarters of 2017 the NTP reduced this; the overall level of underreporting for 2017 was 36% (Fig. B4.4.5).

FIG. B4.4.2
Health facility mapping

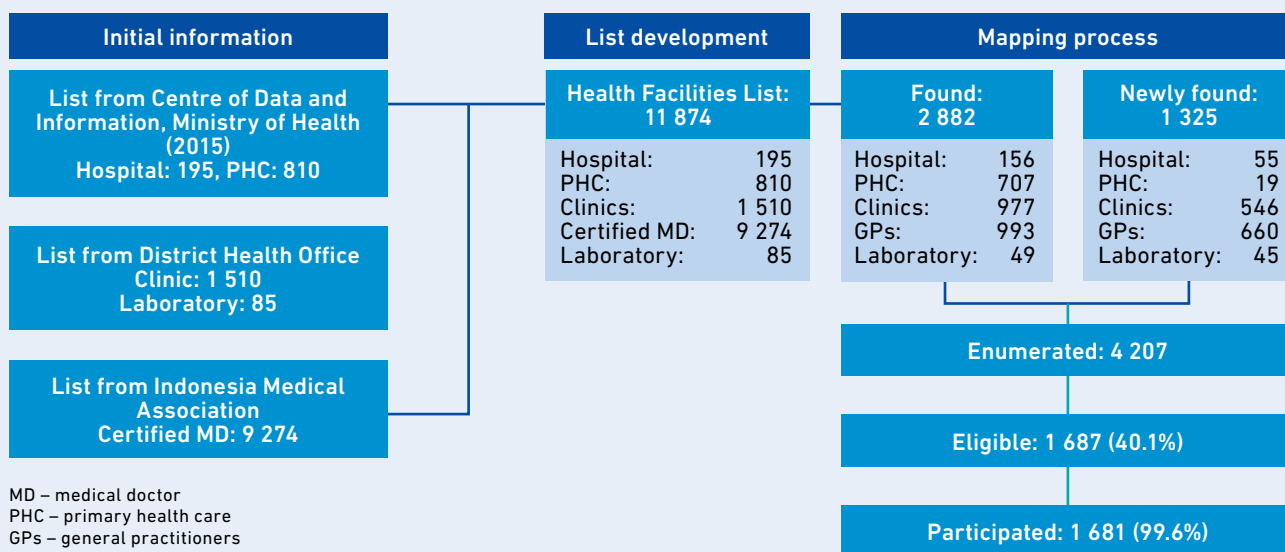
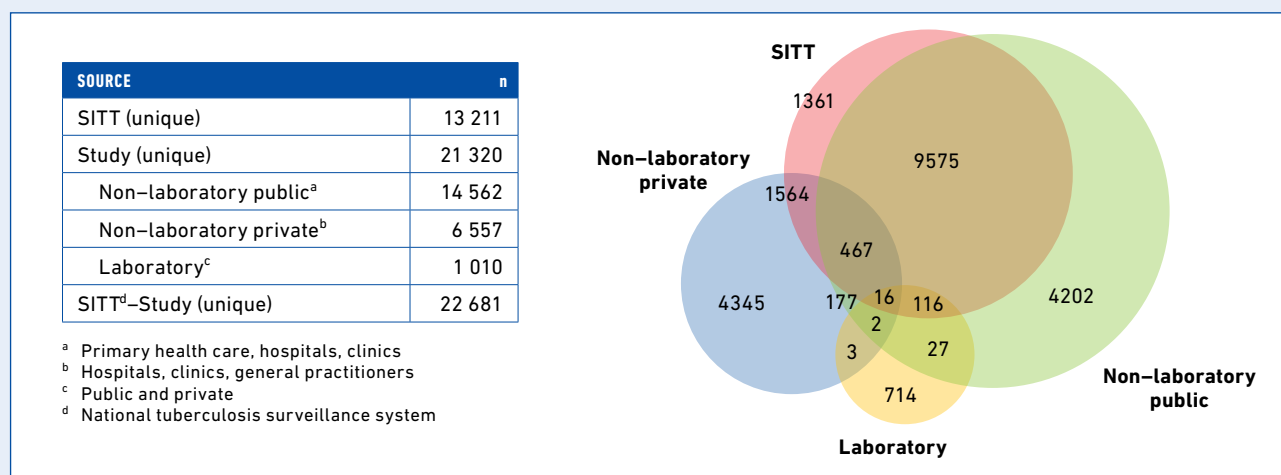


FIG. B4.4.3
Matching results of TB cases found from the study and registered in the national TB surveillance system (SITT) during the first quarter of 2017



Lessons learned

The key lessons learned from the study were as follows:

1. TB underreporting in Indonesia up to the first quarter of 2017 was very high.
2. Out of incident TB cases missed from the TB surveillance system, two thirds were not reported and the remaining one third were not detected.
3. An up-to-date master list of health facilities (both for the public and private sectors) needs to be maintained for efficient monitoring of the effectiveness of the policy of mandatory case notification.
4. Record linkage should occur routinely (at least once a year) between the database of the NTP (SITT) and other databases of TB cases.
5. A unique identifier (e.g. health insurance number) strongly facilitates disease surveillance in general, and record linkage in particular.

6. More than 95% of TB cases identified during the study were treated using a drug regimen that was consistent with national TB guidelines, indicating that a high quality of TB care is provided by different types of health providers. The low level of drug resistance among TB cases found in the recent national survey of drug resistance (see [Chapter 3](#)) is consistent with this finding.

Policy and programmatic implications

The major implications of study results, some of which require high-level policy action, include the following:

1. All health-care providers need to be made aware that all TB cases must be reported. Previously, emphasis was given to reporting of pulmonary, bacteriologically confirmed cases.
2. Human resource capacity for data collection and management needs to be increased. Dedicated staff are required to work on data entry and reporting, especially

FIG. B4.4.4

Trends in estimated rates of TB incidence and case notifications, 2000–2017. Updated time series of incidence rates for 2000–2017 are shown in **blue**, previous estimates for 2000–2016 that were based on the 2013–2014 national TB prevalence survey are shown in **orange** and case notifications are shown in **black**. Shaded areas represent uncertainty bands.

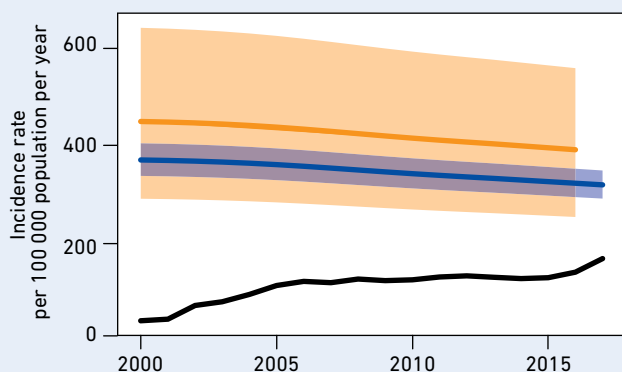
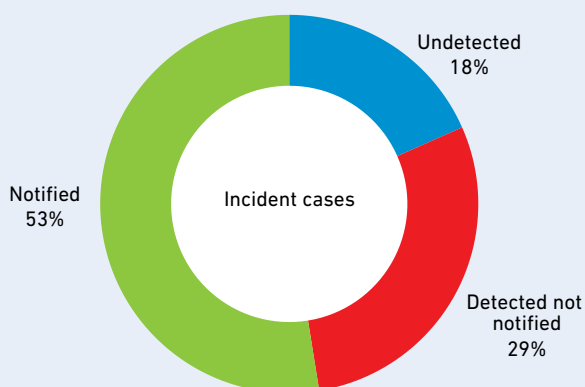


FIG. B4.4.5

TB incidence in Indonesia, 2017 broken down into those cases detected and notified, those detected and not notified (or underreported) and those not detected at all



in support of engagement with private sector health-care providers.

3. Data recording and reporting needs to be simplified, especially for private sector health-care providers. Challenge TB Indonesia has developed an android application called WifiTB, which is designed to simplify and facilitate case notification by the private sector.^d
4. Establishment of integrated information systems, especially in hospitals, needs to be promoted.
5. The NTP needs to increase the involvement and engagement of professional organizations and laboratory networks at district level, to promote adherence with national diagnostic and reporting requirements.

In 2017 and 2018, the NTP, with partners including Challenge TB Indonesia and the Global Fund, has been strengthening models of engagement with all care providers. This has been based on the model of district PPM, in which district staff engage with hospitals and puskesmas staff engage with local

TABLE B4.4.1

Level of TB under-reporting (%) by type of health care provider, type of TB case, age, sex and strata, accounting for sampling design

	MEAN PERCENTAGE (95% CONFIDENCE INTERVAL)
Total	41 (36–46)
By type of health provider	
Primary health care (“puskesmas”)	15 (11–20)
Non-primary health care	71 (61–79)
Hospital	62 (52–72)
Other ^a	96 (92–98)
By TB case type	
Bacteriologically confirmed	21 (16–26)
Clinically diagnosed	55 (49–61)
By site of disease	
Pulmonary	38 (33–44)
Extra-pulmonary	58 (49–66)
By age	
<15 years	54 (44–64)
≥15 years	39 (34–44)
By sex	
Female	41 (36–47)
Male	41 (36–46)
By strata	
Sumatera	40 (24–59)
Bali/Java	42 (18–47)
Other	39 (28–51)

^a Clinics, general practitioners, laboratories.

GP networks. The inventory study provided additional ideas for how to strengthen approaches to working with these care providers.

Conclusions and next steps

The 2017 national TB inventory study in Indonesia is the largest study of its kind ever conducted globally. It generated high-quality data, as well as important evidence with clear policy and programmatic implications.

Following official and wide dissemination of findings in April 2018, results are now being used to help to develop national and district level responses for further rollout. A study report is being finalized and results will be summarized in a paper for a peer-reviewed journal.

^a World Health Organization. Assessing tuberculosis underreporting through inventory studies. Geneva: WHO; 2012 (available on http://www.who.int/tb/publications/inventory_studies/en, accessed 15 August 2018).

^b When using population estimates from the Statistics Bureau, Indonesia – instead of population estimates from the United Nations Population Division – the estimated TB incidence rate is 322 (95% CI: 294–352) per 100 000 population per year.

^c Further discussions were also held between WHO and the NTP during the drafting and finalization of this box.

^d Challenge TB Indonesia. Notification through WiFi TB application for private providers: Update and expansion plan. https://pdf.usaid.gov/pdf_docs/PA00SWTN.pdf

BOX 4.5

Community contributions to TB notifications and treatment support

The WHO End TB Strategy calls for close collaboration with affected communities and civil society organizations in planning, implementation, and monitoring and evaluation.

Community-based TB activities include a wide range of activities that contribute to prevention, diagnosis, treatment and care, and can positively influence TB outcomes. They are delivered primarily by community health workers (CHWs) and community volunteers (CVs)^a who are drawn from within the community and are, therefore, both accessible and acceptable to community members.

In the current era of the SDGs and in the context of UHC, primary health care is receiving greater attention. An increasing number of countries are taking steps to absorb cadres of CHWs into the workforce of national health systems. This will help to strengthen community-based service delivery and also help to reach those who are otherwise unable to access health services.

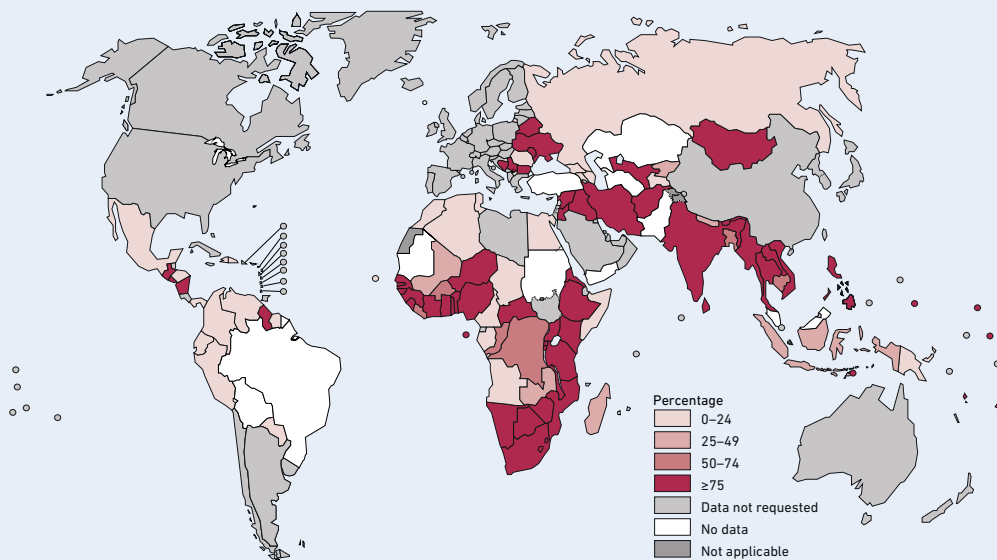
In WHO's 2018 round of global TB data collection, 61 countries reported data about the contribution of communities through CHWs or CVs to TB notifications or treatment support. This represents nearly a fivefold increase in reporting since 2013, when data were first collected on the two core indicators (contributions to new notifications and to treatment support) used to monitor community contributions to TB outcomes.

The contribution of community referrals to TB notifications was reported by 57 countries. The percentage of notified TB patients attributed to community referrals averaged 27% (ranging from less than 1% in Iran and Sri Lanka, to 100% in Bulgaria, Congo, Gabon, Haiti, Paraguay, and Sao Tome and Principe). The treatment success rates for people who benefited from any form of community treatment support could be calculated for 44 of these 57 countries. The average treatment success rate was 87%, ranging from 39% in Burkina Faso to 100% in Burundi, Honduras and Serbia.

In 2018, 112 countries were asked to report data on community contributions to TB care, of which 91 (81%) reported implementing community-based TB activities. In these 91 countries, the mean coverage of community-based TB activities was 74% of all basic management units

FIG. B4.5.1

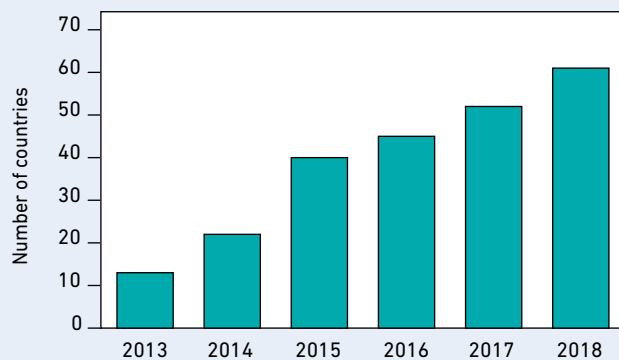
Percentage of basic management units in which there is community contribution to new case finding and/or to treatment adherence support, 2017^a



^a Data only requested from 111 countries.

FIG. B4.5.2

Number of countries reporting on WHO community engagement indicators, 2013–2018



(Fig. B4.5.1). However, 30 of the 91 countries did not have data on community contributions to TB outcomes (Fig. B4.5.2). Further efforts are needed to include the core community indicators in national data recording systems.

^a CHWs and CVs are defined here: World Health Organization. ENGAGE-TB Approach: Operational guidance – integrating community-based tuberculosis activities into the work of nongovernmental and other civil society organizations (WHO/HTM/TB/2012.8). Geneva: WHO; 2012 (http://www.who.int/tb/publications/2012/engage_tb_policy/en/, accessed 14 August 2018). The guidance is currently being revised.

4.2.2 Treatment coverage of ART for HIV-positive TB cases

WHO recommends ART for all HIV-positive TB patients within the first 8 weeks of starting TB treatment, and within two weeks in profoundly immunosuppressed HIV-positive TB patients with low CD4 counts less than 50. The number of notified HIV-positive TB patients on ART has grown in recent years (Fig. 4.18); it reached 377 040 in 2017, equivalent to 84% of the notified TB patients known to be HIV-positive (Table 4.1).¹ In the 30 high TB/HIV burden countries, overall, 85% of the TB patients known to be HIV-positive were on ART. Eight of these countries (Eswatini, Kenya, Malawi, Mozambique, Namibia, Papua New Guinea, Uganda and the United Republic of Tanzania) maintained coverage of at least 90% in both 2016 and 2017. In contrast, there were six high TB/HIV burden countries (Angola, Botswana, Brazil, Guinea-Bissau, Indonesia and Liberia) in which less than 50% of HIV-positive TB patients were started on ART in 2017. Cameroon, Chad and China did not report data on ART for TB patients for 2017.

ART treatment coverage for people with TB can also be assessed by comparing the number of HIV-positive TB patients on ART with the estimated number of HIV-positive incident TB cases (Fig. 4.19). This comparison revealed larger gaps. Globally in 2017, the number of HIV-positive TB patients on ART was 41% of the estimated global number of incident HIV-positive TB cases; this is lower than the global ART coverage of 59% among all people living with HIV in 2017. There was considerable variation among the high TB/HIV burden countries and, according to best estimates, only eight countries achieved ART coverage of more than 50% (Eswatini, Ethiopia, Malawi, Namibia, South Africa, Uganda, Zambia and Zimbabwe).

Improvements are still needed in the detection of active TB disease among HIV-positive people, the coverage of HIV testing among TB patients, and the enrolment of HIV-positive TB patients in ART. A recent example of efforts to address gaps in the cascade of care in India is provided in Box 4.6. An overview of progress and gaps in TB preventive treatment among people living with HIV is provided in Chapter 5.

4.2.3 Treatment coverage for MDR/RR-TB

Trends in the number of patients enrolled in MDR-TB treatment globally and in the 30 high MDR-TB countries since 2009 are shown in Fig. 4.12 and Fig. 4.13, respectively. The number of people enrolled in treatment globally was 139 114 in 2017, representing a more than fourfold increase since 2009 (when WHO first asked countries to report data) and up from 129 689 in 2016.

Between 2016 and 2017, there were notable increases in enrolments in the Democratic People's Republic of Korea (from 814 to 1732), Indonesia (from 1879 to 3042) and Angola (from 334 to 534), and modest increases in several other high MDR-TB burden countries. However, the number of enrolments fell in nine high MDR-TB burden countries (Fig. 4.13).

Globally, the 139 114 patients starting second-line MDR-TB treatment in 2017 represented 25% of the 558 000 estimated MDR/RR-TB incident cases in 2017 (Fig. 4.20). The highest levels of treatment coverage among high MDR-TB burden countries were in Kazakhstan (82%) and South Africa (73%). Ten countries accounted for about 75% of the gap between enrolments in MDR-TB treatment in 2017 and the estimated number of incident MDR/RR-TB cases in 2017; and China and India accounted for 40% of the total gap (Fig. 4.21). Treatment coverage will not improve globally unless there is an intensification of efforts in the countries with the largest burden, particularly China, India and Indonesia.

The number of cases starting MDR-TB treatment in 2017 was equivalent to 87% of the 160 684 MDR/RR-TB patients notified in 2017 (Fig. 4.12). The figure exceeded 90% in 17 high MDR-TB burden countries (Fig. 4.13) and in the WHO European Region and the WHO Region of the Americas; however, it was much lower in the WHO African Region and the WHO Western Pacific Region.² Enrolments represented no more than 60% of the number of notified MDR/RR-TB cases in two high MDR-TB burden countries in 2017: China (45%) and Indonesia (60%). These low percentages show that progress in detection is outstripping the capacity to provide treatment; they may also reflect weaknesses in data collection systems. In these settings, the risk of transmission of drug-resistant TB is high, and efforts are needed to rapidly close gaps in enrolment and notification.

In many countries, one of the barriers to adequate access to treatment of drug-resistant TB is that the network for the programmatic management of drug-resistant TB (PMDT) is too centralized and too reliant on hospital-based models of care. Greater decentralization and more use of outpatient models of care are needed.

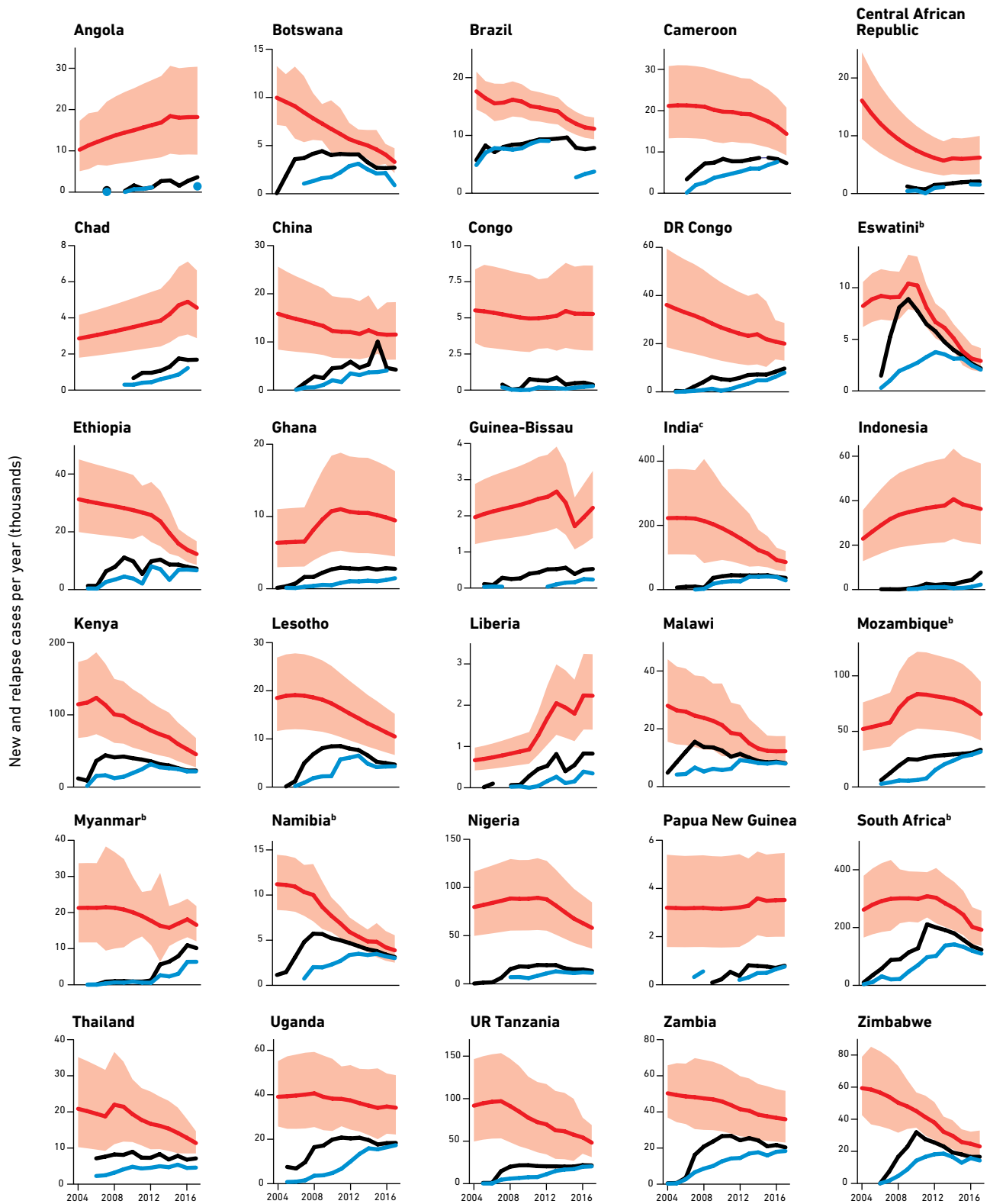
Globally, 8703 patients with XDR-TB were enrolled in treatment in 70 countries and territories, a small (2%) increase compared with 2016. In 20 of these countries, the number of XDR-TB cases enrolled in treatment was less than the number notified. Only one high MDR-TB burden country (Thailand) and three other countries (Algeria, Bhutan and Uganda) reported prescribing morphine to treat pain or terminal dyspnoea in patients for whom second-line TB treatment regimens did not work, suggesting that there are widespread unmet needs in terms of end-of-life care as well as inadequate data gathering on this issue.

¹ There may be discrepancies in data on provision of ART to HIV-positive TB patients that are reported by NTPs and national HIV programmes. These discrepancies have reduced in recent years and have mostly been resolved through follow-up and validation efforts.

² For data for WHO regions, see Annex 3.

FIG. 4.18

Number of new and relapse cases^a known to be HIV-positive (black) and number started on ART (blue) compared with estimated number of incident HIV-positive TB cases (red), 2004–2017, 30 high TB/HIV burden countries



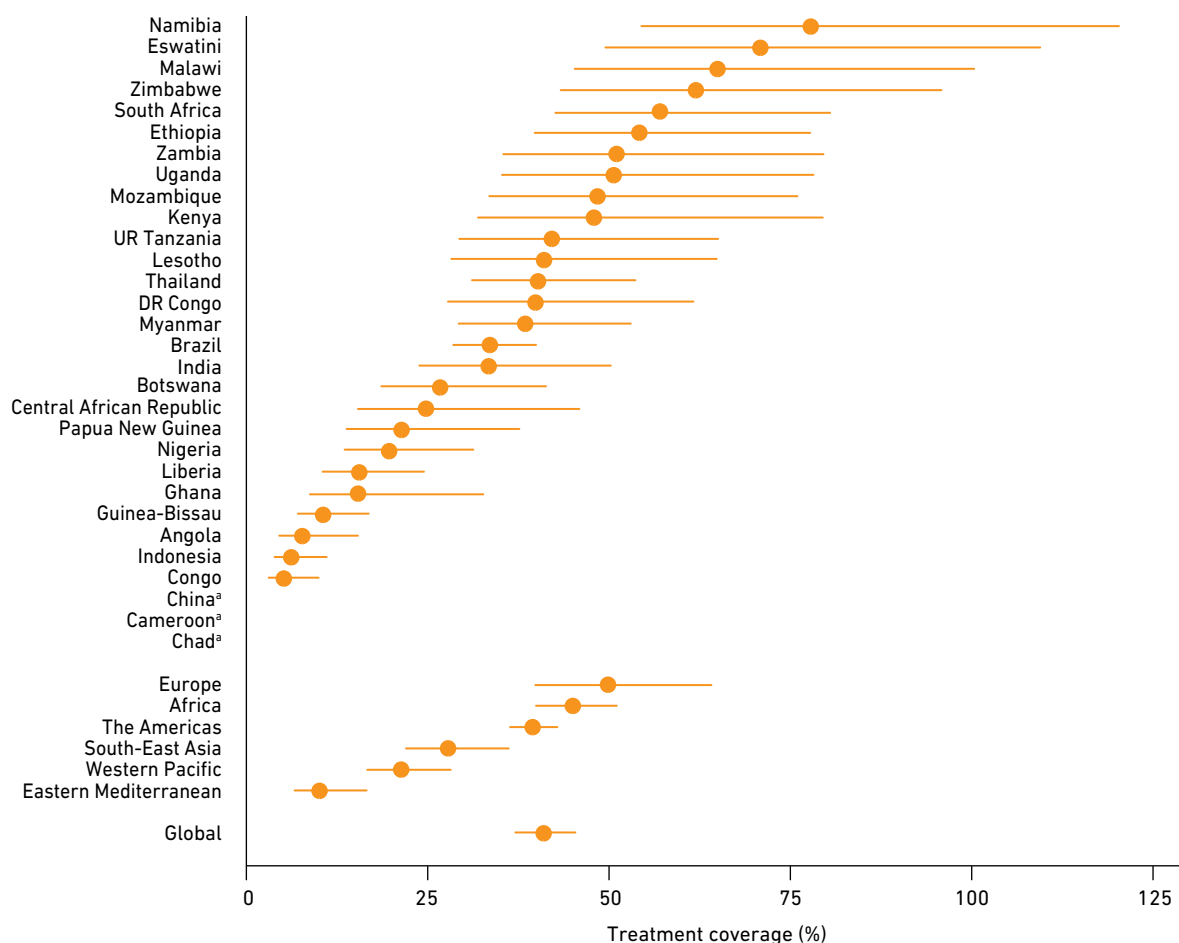
^a The calculation is for all cases in years prior to 2015.

^b Estimates of TB incidence for Eswatini, Mozambique, Myanmar, Namibia and South Africa will be reviewed after final results from their respective national TB prevalence surveys are available in 2019.

^c Estimates of TB incidence for India are interim in nature, pending results from the national TB prevalence survey planned for 2019/2020.

FIG. 4.19

Estimated ART treatment coverage for HIV-positive TB cases (HIV-positive TB patients on ART as a percentage of the estimated incidence of HIV-positive TB) in 2017, 30 high TB/HIV burden countries, WHO regions and globally



^a No data.

4.3 Treatment outcomes

This section summarizes the latest results of treatment for new and relapse cases of TB who started treatment on a first-line regimen in 2016 (including people with HIV-associated TB), and people detected with RR-TB, MDR-TB or XDR-TB who started a second-line MDR-TB regimen in 2015.¹

4.3.1 Treatment outcomes for new and relapse TB patients

Data on treatment outcomes for new and relapse cases of TB in 2016 are shown for the world, the six WHO regions and the 30 high TB burden countries in Fig. 4.22. The global trend 2012–2016 is shown in Fig. 4.23. Globally, the treatment success rate for the 5.9 million new and

relapse cases who were treated in the 2016 cohort was 82%. This is a reduction from 86% in 2013 and 83% in 2015; the main explanation is that in countries where notifications have increased (notably India), reporting of treatment outcomes has not kept pace with notifications. In India, treatment outcome data were not reported for 22% of cases notified in 2016. The absolute number of TB patients reported to have been successfully treated has risen substantially over the past 16 years, both globally and in all WHO regions (Fig. 4.24).

Among the six WHO regions, the highest treatment success rates in 2016 were in the WHO Eastern Mediterranean Region (92%) and the WHO Western Pacific Region (91%). The lowest rates (at 75%) were in the WHO South-East Asia Region (due to high proportions of unevaluated cases, especially in India), the WHO Region of the Americas (due to high levels of loss to follow-up and missing data) and the WHO European Region (due to high rates of treatment failure and death, influenced by the high frequency of MDR/RR-TB).

¹ For definitions of treatment outcomes, see World Health Organization. Definitions and reporting framework for tuberculosis – 2013 revision (updated December 2014) (WHO/HTM/TB/2013.2). Geneva: WHO; 2013 (www.who.int/iris/bitstream/10665/79199/1/9789241505345_eng.pdf, accessed 21 June 2018).

BOX 4.6

Cascade of TB case-finding among people attending HIV care in India, 2017

For the past year, the Revised National TB Control Programme (RNTCP) and National AIDS Control Organization (NACO) in India have been capturing data to evaluate TB case-finding activities among people attending ART centres.

According to data reported by the RNTCP for 2017, an estimated 58% (n = 49 000 [95% CI: 21 000–84 000]) of people with HIV-associated TB were not reported to have reached TB care. Reasons for missing people with TB include poor access to services, weaknesses in service delivery, gaps in recording and reporting, and limited engagement of the private sector.

Identifying the size and nature of these gaps requires data about each part of the cascade of care, from initial evaluation for TB signs and symptoms to the start of treatment. For HIV-associated TB, analysis should also be conducted on the HIV care cascade among TB patients. Depending on the degree of integration between a country's TB and HIV services and their respective monitoring and evaluation systems, the patient pathway for HIV-associated TB can be more complex to track, since there may be a need for referral from one programme to another. Close collaboration between the TB and HIV services is therefore critical for reducing loss to follow-up and preventable mortality. To strengthen integration, the Government of India has rolled out TB service delivery from all HIV clinics since December 2016.

Although reporting of HIV case-based data for the TB case-finding cascade has not yet been possible in India, the RNTCP and NACO have been able to report data based on clinic visits. People living with HIV made almost 11 million visits to ART centres in 2017. In 83% of these visits, the clinic attendees received an evaluation for TB. Of those who were evaluated, 6% had TB symptoms.

FIG. B4.6.1

Cascade of TB case-finding among people living with HIV (PLHIV) attending antiretroviral therapy (ART) clinics, India, 2017

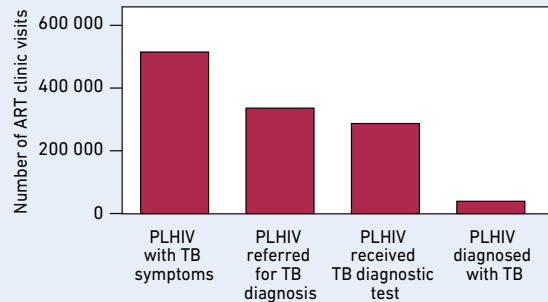


Fig. B4.6.1 depicts the next stages of the TB case-finding cascade. In 514 862 consultations, 65% of those attending ART centres in whom TB symptoms were identified were referred for TB diagnosis; of those, 85% underwent a TB diagnostic test, of whom 14% were found to have TB. These extensive nationwide efforts help to identify gaps in referral and diagnosis to prompt follow-up as necessary.

Capturing data for each step in the cascade can be challenging, unless a robust, integrated and preferably electronic, case-based recording and reporting system is in place that communicates the information across the programmes. In the absence of electronic recording and reporting systems, periodic patient record surveys of people attending HIV care are recommended, to evaluate the gaps in the TB case-finding continuum, which in turn helps to inform and improve programming, recording and reporting, and client follow-up.

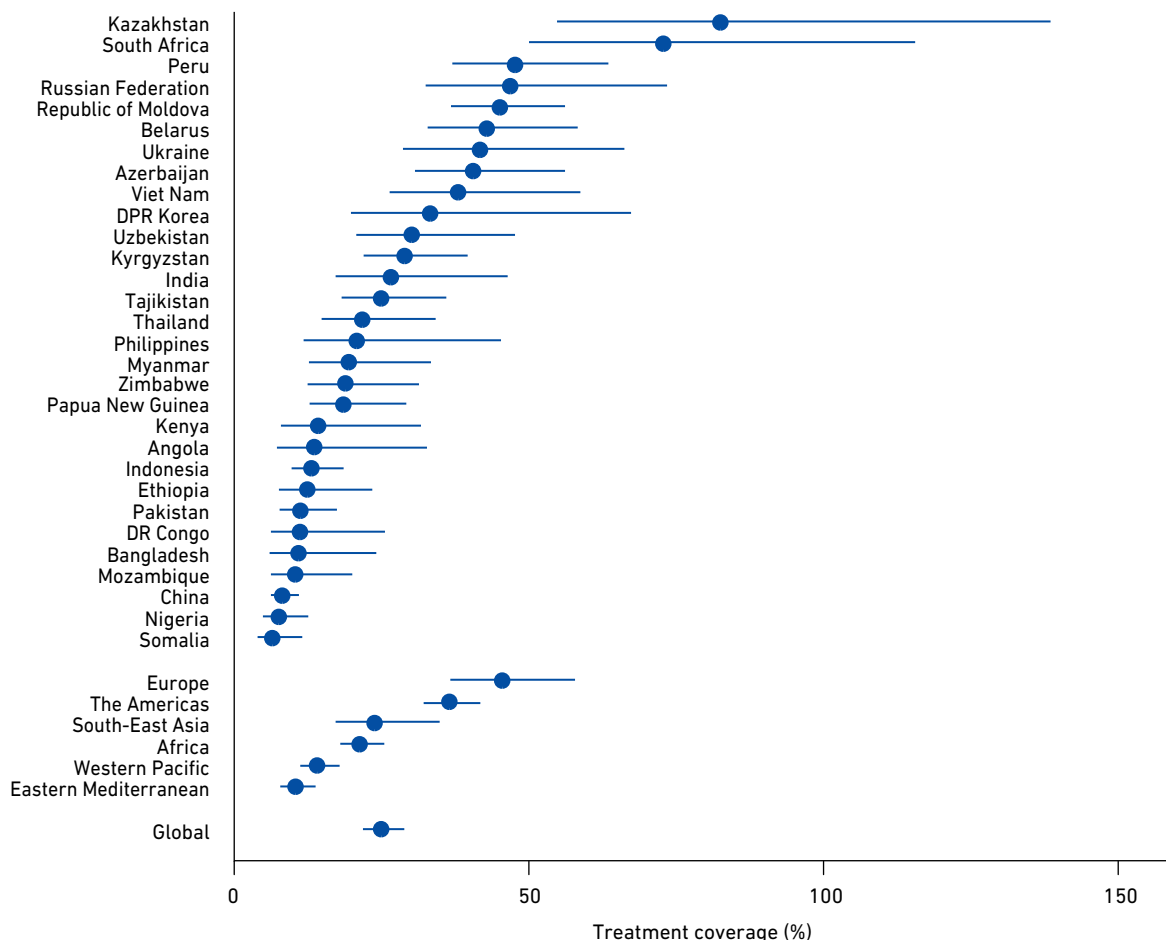
Only nine of the 30 high TB burden countries reached or exceeded a 90% treatment success rate. However, in several high TB burden countries, the completeness of outcome reporting was low. In five countries (Angola, Brazil, Central African Republic, Liberia and Papua New Guinea), loss to follow-up exceeded 10%, and in four countries (Angola, Congo, India and Papua New Guinea), the treatment outcome was not documented for more than 10% of cases.

4.3.2 Treatment outcomes for new and relapse TB patients coinfecting with HIV

A total of 124 countries reported treatment outcomes for the 2016 patient cohort disaggregated by HIV status; collectively, these countries accounted for 94% of the HIV-positive TB patients reported by NTPs in 2016. These 124 countries included 24 of the 30 high TB/HIV burden countries; no data were reported by Chad, Congo, the Democratic Republic of the Congo, Ethiopia, Liberia and Papua New Guinea (Fig. 4.25). Overall, the treatment success rate was 77%, an increase from 68% in 2012 (Fig. 4.23), although still worse than the level of 82% for all new and relapse TB patients.

FIG. 4.20

Estimated treatment coverage for MDR/RR-TB (patients started on treatment for MDR-TB as a percentage of the estimated incidence of MDR/RR-TB) in 2017, 30 high MDR-TB burden countries, WHO regions and globally



Globally, the proportion of HIV-positive TB patients who died during treatment was 11%, similar to previous years and about three times the level among all new and relapse cases (4%). In the WHO regions, the relative difference was smallest in the WHO African Region (9% versus 5%) and highest in the WHO Eastern Mediterranean Region (14% versus 2%). In the WHO Region of the Americas and the WHO European Region, the proportion of HIV-positive TB patients who died was 20% and 22%, respectively.

Reasons for comparatively poor outcomes for HIV-positive TB patients include late detection of HIV-associated TB and MDR-TB, and delays in starting ART or TB treatment. To reduce excessive TB mortality in people who are HIV-positive, WHO recommends routine HIV testing among presumptive and diagnosed TB cases, TB screening among people living with HIV, early ART, improved infection control and provision of TB preventive treatment. Options that could help to ensure earlier diagnosis and reduce mortality include strategic placement of WHO-recommended rapid molecular TB diagnostics such as Xpert MTB/RIF within HIV care settings, and

uptake of the lateral flow urine lipoarabinamannan assay (LF-LAM) for seriously ill people living with HIV.

4.3.3 Treatment outcomes for TB patients with MDR/RR-TB and XDR-TB

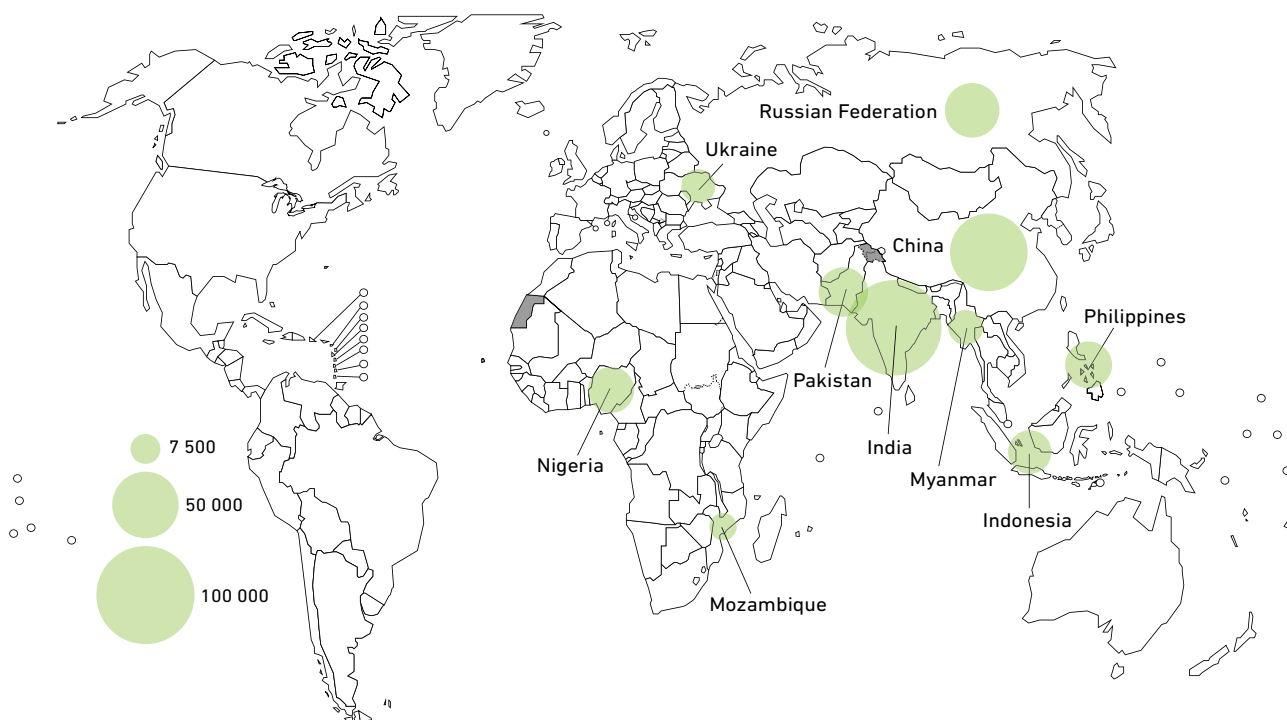
A total of 144 countries and territories reported treatment outcomes for people started on MDR-TB treatment in 2015.¹ The number of cases reported in annual cohorts has steadily increased over time, reaching 114 180 cases globally in the 2015 cohort. Overall, the proportion of MDR/RR-TB patients in the 2015 cohort who successfully completed treatment (i.e. cured or treatment completed) was 55%: in 8% the treatment failed, 15% died, 14% were lost to follow-up and for 7% there was no outcome information (Fig. 4.26).

Globally, treatment success has increased slightly in recent years (Fig. 4.23). In the 2015 cohort, the treatment success rate was highest in the WHO Eastern Mediterranean Region (62%) and was lowest in the WHO

¹ This is the latest year for which data on treatment outcomes for drug-resistant TB have been reported to WHO. There is a longer lag time for reporting of data due to the longer duration of treatment for drug-resistant TB.

FIG. 4.21

The ten countries with the largest gaps between the number of patients started on treatment for MDR-TB and the best estimates of MDR/RR-TB incidence, 2017^a



^a The ten countries ranked in order of the size of the gap between the number of patients started on MDR-TB treatment and the best estimate of MDR/RR-TB incidence in 2017 are India, China, Russian Federation, Pakistan, Nigeria, the Philippines, Indonesia, Ukraine, Myanmar and Mozambique.

South-East Asia Region (50%). In contrast, treatment failure was highest in the WHO European Region (12%), and the death rate was highest in the WHO African Region (20%). Loss to follow-up was highest in the WHO Region of the Americas (26%), while the WHO Western Pacific Region had the highest percentage of cases without outcome data (17%).

Among the 30 high MDR-TB burden countries, 15 had MDR/RR-TB cohorts in 2015 with more than 1000 cases; among these, only Kazakhstan and Myanmar reported treatment success of more than 75%. Treatment success was less than 50% in China, India, Indonesia, Mozambique, the Republic of Moldova and Zimbabwe. This was due to high rates of death and loss to follow-up in India (20% and 19%) and Indonesia (16% and 31%), high rates of loss to follow-up or missing data about treatment outcome in China (13% and 32%), a high death rate and missing data about treatment outcome in Mozambique (26% and 13%) and Zimbabwe (17% and 32%), and high rates of loss to follow-up and failure in the Republic of Moldova (20% and 18%).

Among 8399 patients started on treatment for XDR-TB in 2015, in 49 countries and territories for which outcomes were reported, 34% completed treatment successfully, 26% died, treatment failed for 19%, and 21% were lost to follow-up or their treatment outcome was not evaluated. India, the Russian Federation and Ukraine accounted for

74% of the 2015 XDR-TB cohort. Among seven countries with XDR-TB cohorts of more than 100 individuals, mortality was highest in India and Uzbekistan (42% and 41%).

Although improving in some countries, the treatment success rate for MDR/RR-TB globally remains unacceptably low. The wider use of more effective MDR-TB treatment regimens designed on the basis of the latest available evidence, and the use of more patient-centred models of care, are expected to help improve this situation. New guidance related to the treatment of drug-resistant TB was recently reviewed by a WHO Guidelines Development Group, and WHO will issue updated guidance later in 2018 (Box 4.7).

By the end of 2017, 62 countries, mostly in Africa and Asia, reported having used shorter MDR-TB regimens (Fig. 4.27). These regimens have been reported to achieve high treatment success rates (87–90%) in selected MDR/RR-TB patients. By the end of 2017, 68 countries reported having imported or started using bedaquiline, and 42 countries had used delamanid (Fig. 4.28 and Fig. 4.29). Most (79%) of the patients treated with bedaquiline were reported by two countries: the Russian Federation and South Africa.

With the introduction of new drugs and regimens, there is a need for active TB drug-safety monitoring and management (aDSM). aDSM is defined as the active and systematic clinical and laboratory assessment of

FIG. 4.22

Treatment outcomes for new and relapse TB cases in 2016, 30 high TB burden countries, WHO regions and globally

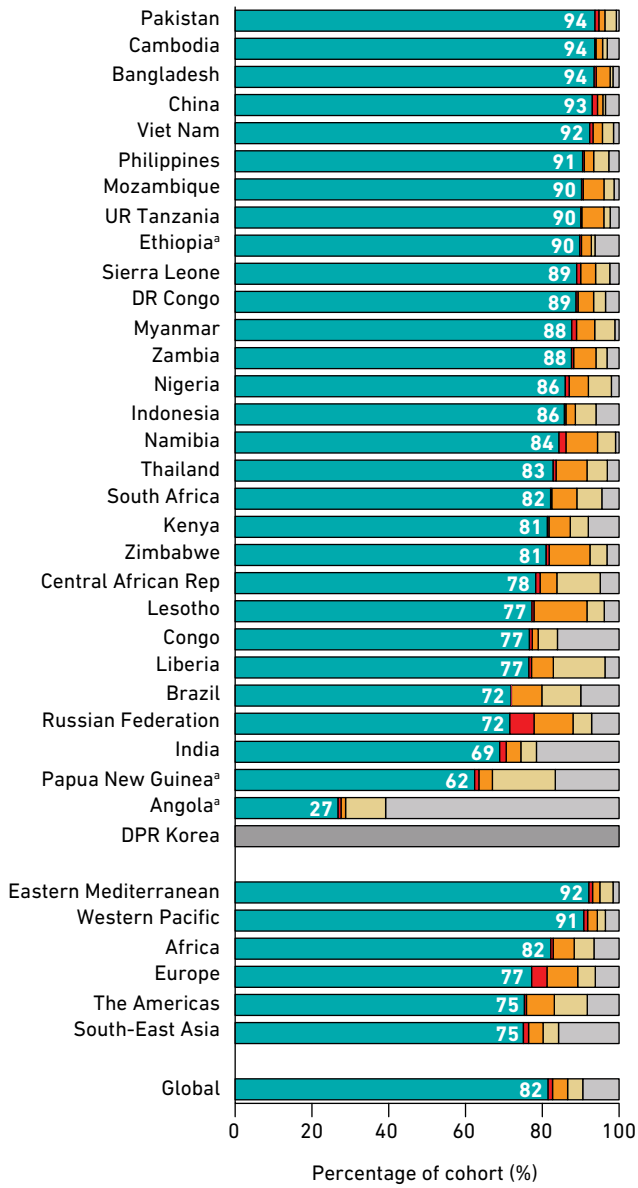
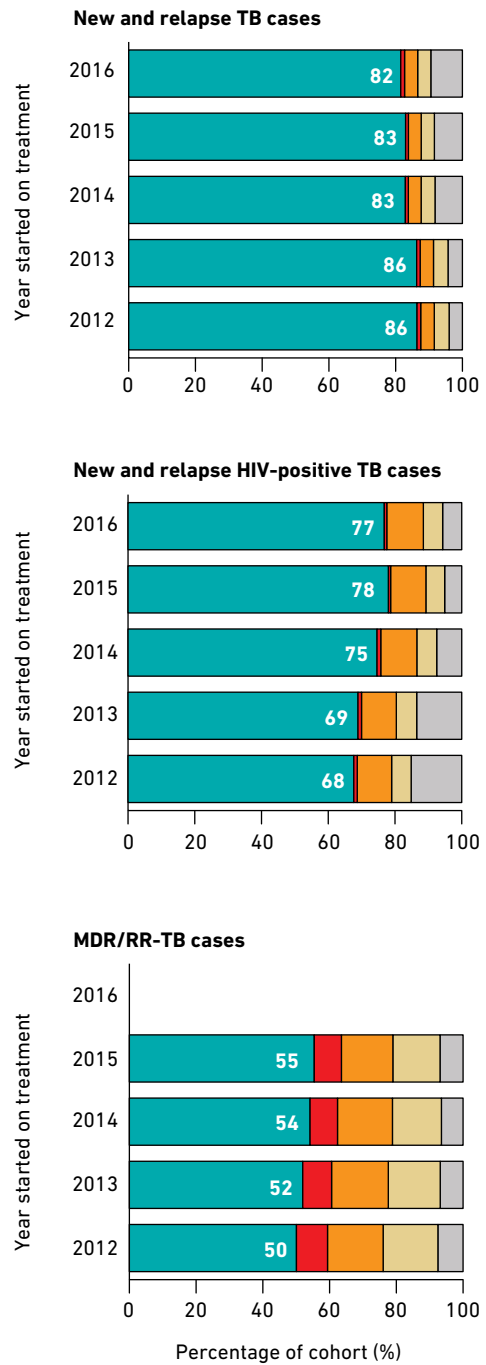


FIG. 4.23

Treatment outcomes for new and relapse TB cases, new and relapse HIV-positive TB cases, and MDR/RR-TB cases, 2012–2016 globally^a



■ Treatment success
 ■ Failure
 ■ Died
 ■ Lost to follow-up
 ■ Not evaluated
 ■ No data reported

^a Treatment outcomes are for new cases only.

^a MDR/RR-TB annual treatment cohorts are reported one year later than other TB cohorts.

BOX 4.7

WHO revises its guidance on the treatment of MDR/RR-TB

Since WHO last issued MDR/RR-TB treatment guidelines in 2016,^a new patient data have emerged. These include final results from the first-ever Phase III randomized controlled trials of delamanid and the 9-month shorter MDR-TB regimen.

In 2018, WHO convened a Guideline Development Group (GDG) to advise on the revision of its guidance. The scope of the new guidelines covered the use of the shorter MDR-TB regimen, the composition and duration of longer MDR-TB regimens, and the use of culture to monitor treatment response in these regimens. Before a meeting of the GDG in July 2018, WHO issued a public call for individual patient data from trials, as well as observational studies and programmatic cohorts. Systematic reviews and meta-analyses were undertaken using the Grading of Recommendations Assessment, Development and Evaluation (GRADE) approach, in accordance with WHO's requirements for the formulation of evidence-based health policy.

Based on the latest evidence, the GDG recommended significant changes to the design of longer MDR-TB treatment regimens. The listing of medicines to choose from as first priority include three that have shown the most advantageous balance between effectiveness and toxicity: later generation fluoroquinolones (levofloxacin or moxifloxacin), bedaquiline and linezolid. The regimen is completed with other agents, in order of relative potency. Unlike the previous algorithms, the systematic inclusion of injectable medicines is no longer required, and all-oral longer regimens are now an option for many patients, subject to individual need. Some of the older TB medicines, such as the injectable agents kanamycin and capreomycin, are no longer recommended for use.^b

As a result of these modifications, and as global access to the most effective agents in longer regimens improves, the role of the shorter MDR-TB treatment regimen in programmatic TB care is destined to change.

All patients on longer and shorter MDR-TB regimens should receive appropriate counselling and be informed of their options before enrolment on treatment. aDSM for all patients enrolled in treatment is essential.

The full guidelines will be released in late 2018, along with revised decision aids and practical advice (e.g. drug dosage schedules for adults and children) in an update of the WHO MDR-TB implementation manual.^c

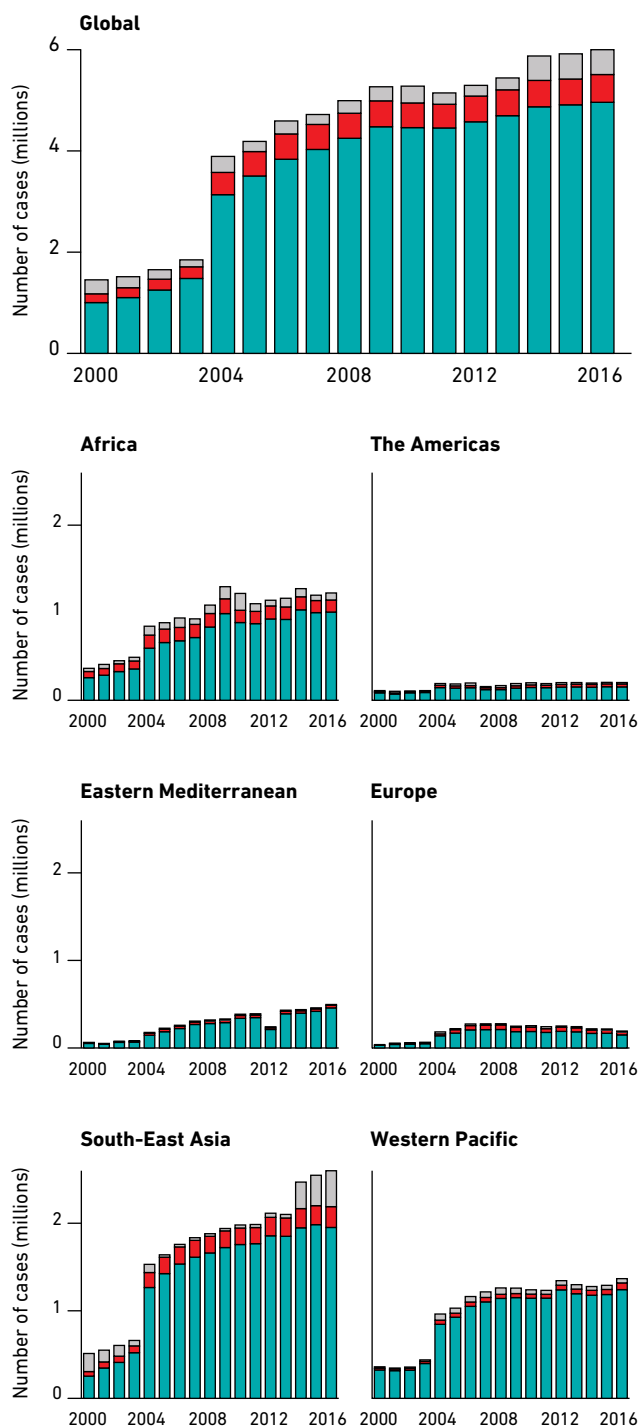
^a World Health Organization. WHO treatment guidelines for drug-resistant tuberculosis (2016 update) (WHO/HTM/TB/2016.04). Geneva: WHO; 2016 (<http://www.who.int/tb/areas-of-work/drug-resistant-tb/treatment/resources/en/>, accessed 3 July 2018).

^b World Health Organization. Rapid Communication: Key changes to treatment of multidrug- and rifampicin-resistant tuberculosis (MDR/RR-TB) (WHO/HTM/TB/20186.18). Geneva: WHO; 2018 (http://www.who.int/tb/publications/2018/WHO_RapidCommunicationMDRTB.pdf?ua=1, accessed 28 August 2018).

^c World Health Organization. Companion handbook to the WHO guidelines for the programmatic management of drug-resistant tuberculosis (WHO/HTM/TB/2014.11). Geneva, World Health Organization; 2015. Available from: http://apps.who.int/iris/bitstream/10665/130918/1/9789241548809_eng.pdf, accessed 15 August 2016).

FIG. 4.24

Treatment outcomes for new and relapse TB cases^a (absolute numbers), 2000–2016, globally and for WHO regions



Legend:
■ Treatment success
■ Failure/Died/Lost to follow-up
■ Not evaluated

^a Cohorts before 2012 included new cases only.

FIG. 4.25

Treatment outcomes for new and relapse HIV-positive TB cases in 2016, 30 high TB/HIV burden countries, WHO regions and globally

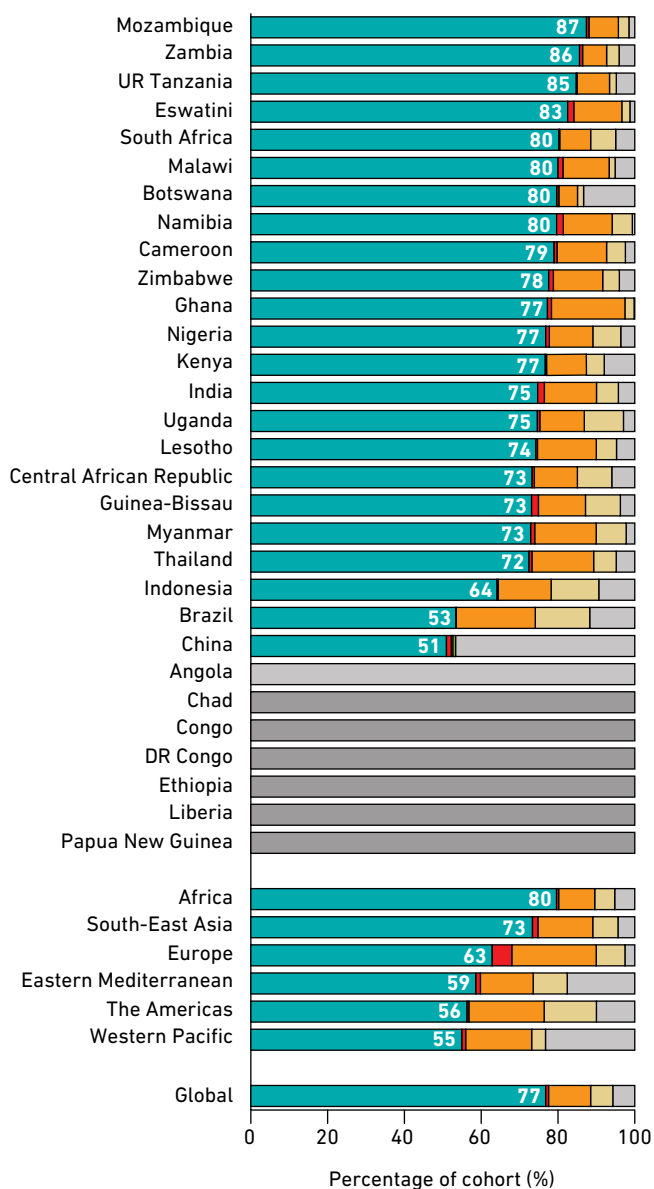
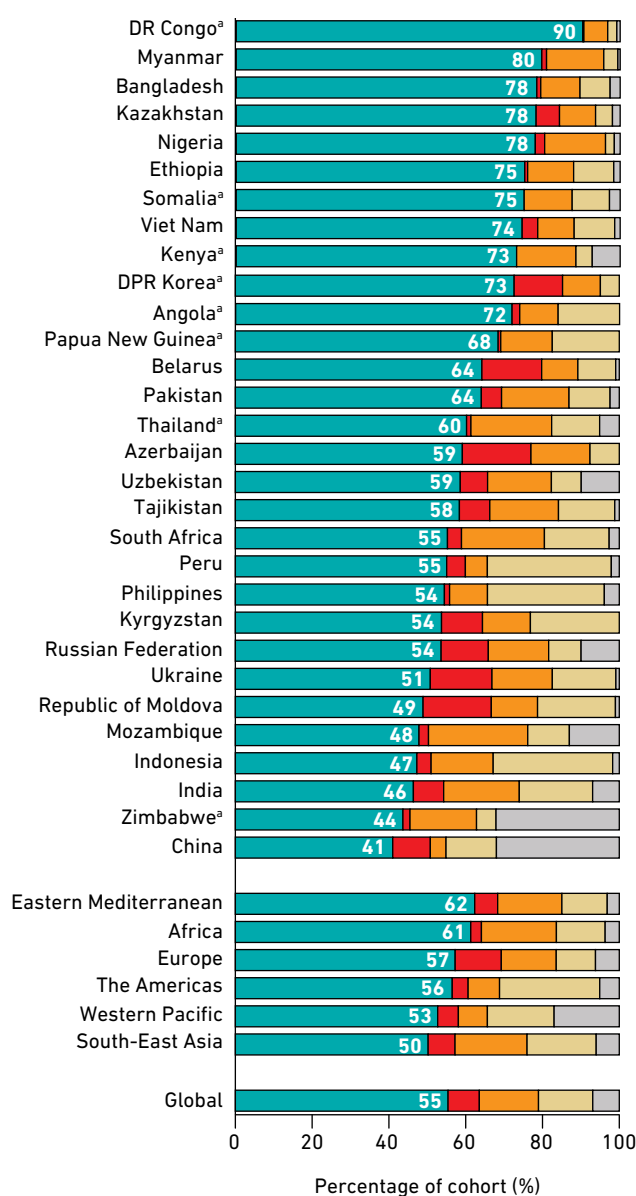


FIG. 4.26

Treatment outcomes for MDR/RR-TB cases started on treatment in 2015, 30 high MDR-TB burden countries, WHO regions and globally



^a These countries reported less than 500 MDR/RR-TB cases who started second line TB treatment in 2015.

patients on treatment with new TB drugs, novel MDR-TB regimens or XDR-TB regimens to detect, manage and report suspected or confirmed drug toxicities.¹ In 2017, 16 of the 30 high MDR-TB burden countries reported data on adverse events collected from their TB information systems.

¹ World Health Organization. Active tuberculosis drug-safety monitoring and management (aDSM): framework for implementation (WHO/HTM/TB/2015.28). Geneva: WHO; 2015 (http://apps.who.int/iris/bitstream/10665/204465/1/WHO_HTM_TB_2015.28_eng.pdf, accessed 15 August 2018).

FIG. 4.27

Countries that had used shorter MDR-TB treatment regimens by the end of 2017

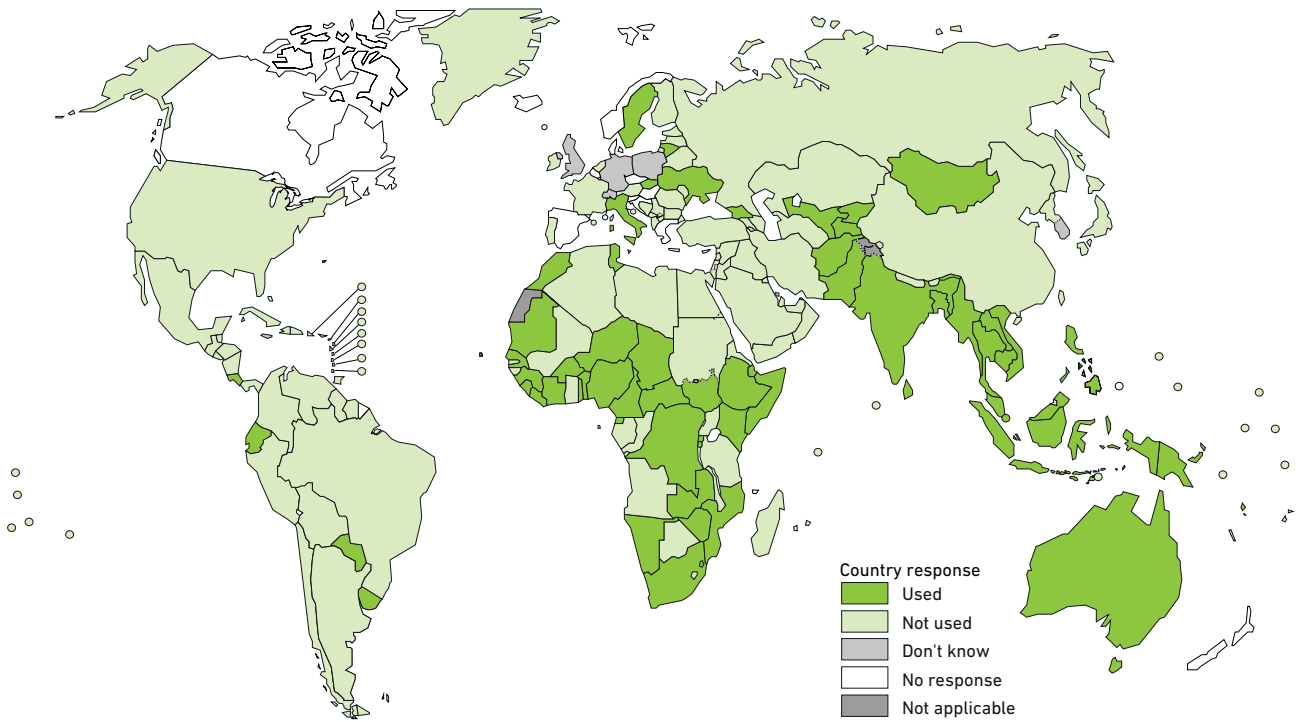


FIG. 4.28

Countries that had used bedaquiline for the treatment of M/XDR-TB as part of expanded access, compassionate use or under normal programmatic conditions by the end of 2017

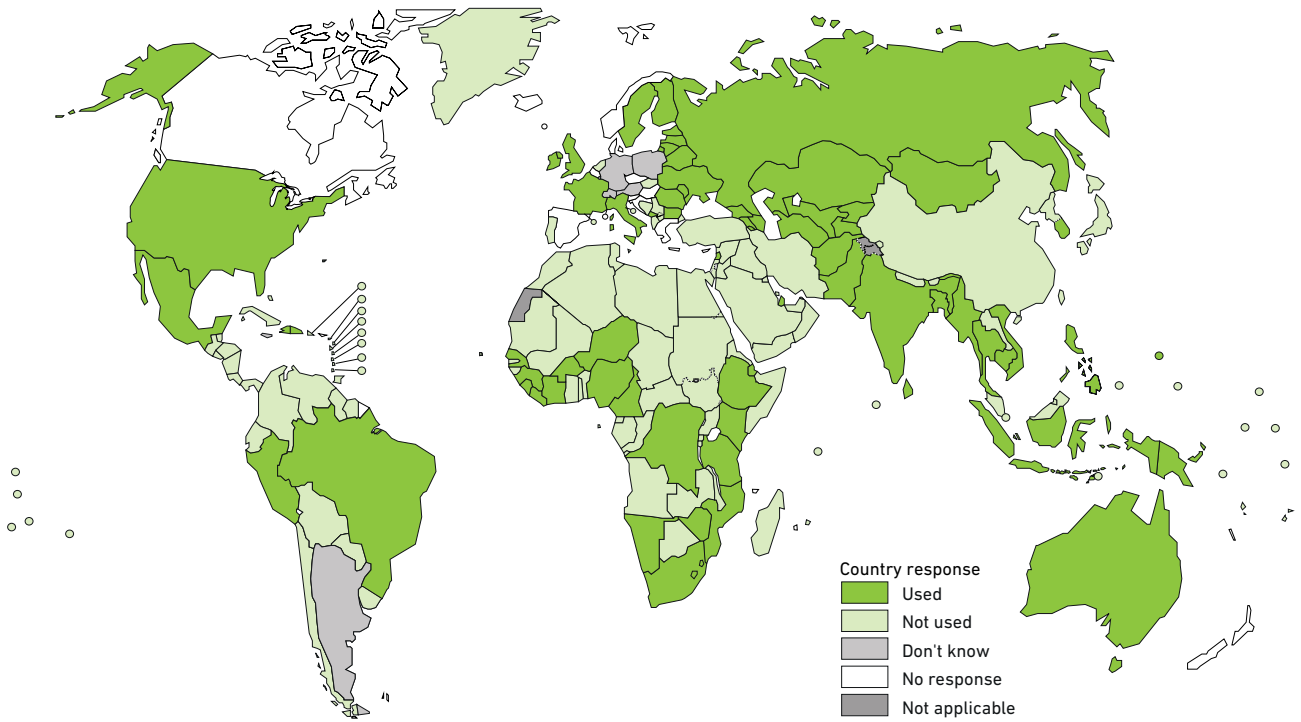
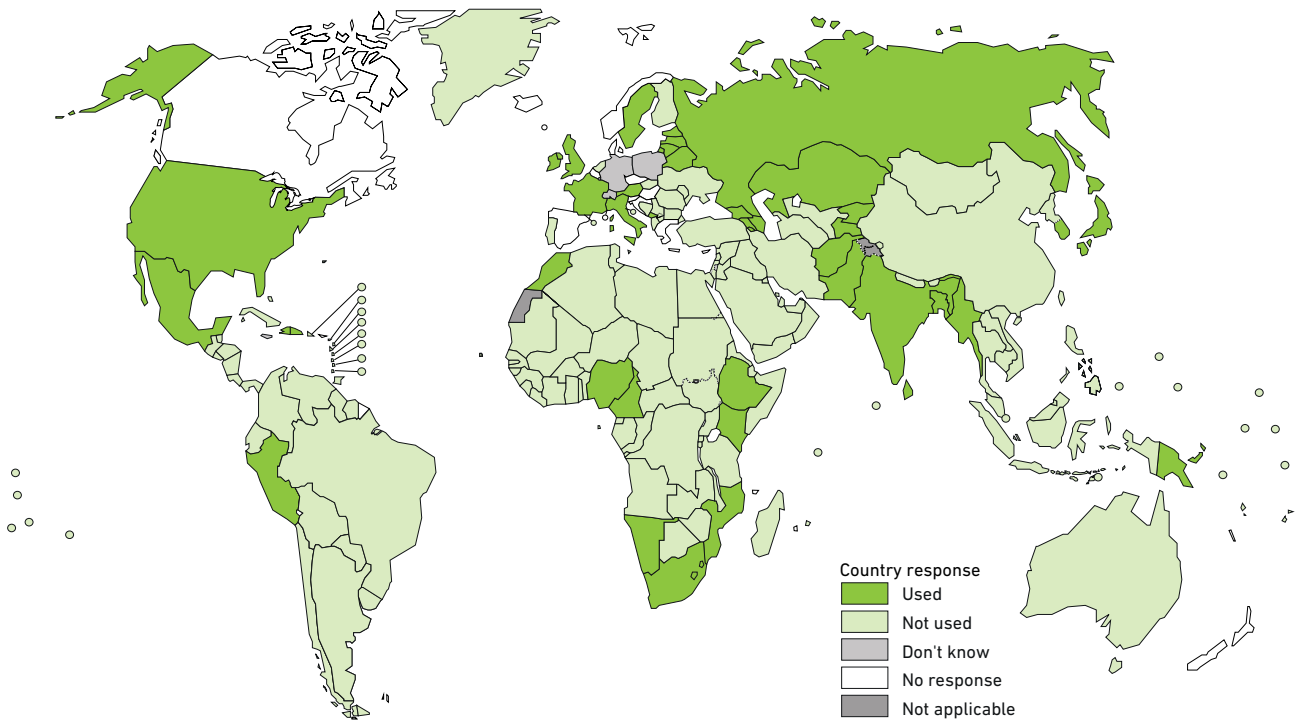


FIG. 4.29

Countries that had used delamanid for the treatment of M/XDR-TB as part of expanded access, compassionate use or under normal programmatic conditions by the end of 2017





A young TB patient waits with his mother to see a doctor at a clinic in New Delhi, India

Gary Hampton / WHO and Vital Strategies

Chapter 5. TB prevention services

KEY FACTS AND MESSAGES

Prevention of new infections of *Mycobacterium tuberculosis* and their progression to tuberculosis (TB) disease is critical to reduce the burden of disease and death caused by TB, and to achieve the End TB Strategy targets set for 2030 and 2035.

Current health interventions for TB prevention are treatment of latent TB infection (LTBI), prevention of transmission of *M. tuberculosis* through infection prevention and control, and vaccination of children with the bacille Calmette-Guérin (BCG) vaccine.

WHO guidelines published before 2018 identified four priority groups in which testing and treatment for LTBI was strongly recommended: people living with HIV (all countries), children aged under 5 years who are household contacts of bacteriologically confirmed pulmonary TB cases (all countries), people aged five years or more who are household contacts of pulmonary TB cases (upper middle-income and high-income countries with a low incidence of TB) and clinical risk groups (upper middle-income and high-income countries with a low incidence of TB). Updated guidance published in 2018 includes an additional recommendation to consider testing and treatment for people aged 5 years or more who are household contacts of bacteriologically confirmed pulmonary TB cases in countries with a high incidence of TB; it also expands the recommendation for clinical risk groups to all countries.

Globally in 2017, 67 countries reported initiating TB preventive treatment for a total of 958 559 people living with HIV; this included 640 201 people who were newly enrolled in HIV care, in 60 countries. South Africa accounted for 39% of the total.

Of the 30 high TB/HIV burden countries, 15 reported providing TB preventive treatment to people newly enrolled in HIV care in 2017, up from 11 in 2016. Coverage ranged from 1% in Eswatini (formerly Swaziland) to 53% in South Africa. Overall, in the 59 countries for which it could be calculated, coverage was 36%.

There were an estimated 1.3 million children aged under 5 years who were household contacts of bacteriologically confirmed pulmonary TB cases and thus eligible for treatment according to WHO recommendations. The number of children in this age group reported to have been started on TB preventive treatment increased to 292 182 in 2017, up 79% from 163 720 in 2016, and a more than three-fold increase from 87 242 in 2015, but was still only 23% of those estimated to be eligible.

Improvements to routine surveillance are needed to capture complete and reliable data on the provision of TB preventive treatment.

The ratio of the TB notification rate among health-care workers to the TB notification rate in the general adult population is a good indicator of the impact of TB infection prevention and control in health facilities. In 2017, a total of 9299 health-care workers from 65 countries were reported with TB; China accounted for 35% of these cases. In six countries (Brunei Darussalam, Colombia, Dominican Republic, Honduras, Paraguay and Zimbabwe), the number of TB cases per 100 000 health-care workers was more than double the notification rate in the general adult population.

BCG vaccination should be provided as part of national childhood immunization programmes according to a country's TB epidemiology. In 2017, 158 countries reported providing BCG vaccination as a standard part of these programmes, of which 120 reported coverage of at least 90%, up from 111 countries in 2016.

Prevention of new infections of *Mycobacterium tuberculosis* and their progression to tuberculosis (TB) disease is critical to reduce the burden of disease and death caused by TB, and to achieve the End TB Strategy targets set for 2030 and 2035.¹ The targets of an 80% reduction in TB incidence from the 2015 level by 2030, and of a 90% reduction by 2035, will require a historically unprecedented acceleration in the rate at which TB incidence falls after 2025 (Chapter 2). This accelerated rate (an average of 17% per year between 2025 and 2035) is possible only if the probability of progression from latent TB infection (LTBI) to active TB disease among the approximately 1.7 billion people already infected worldwide² is substantially reduced.³ Health-care interventions that could help to cut the risk of progression from LTBI to active TB disease include more effective drug treatments for LTBI, and development of a vaccine to prevent reactivation of LTBI in adults. Action on the broader determinants of TB could also cut the risk; these are discussed further in Chapter 7.

Currently, three major categories of health interventions are available for TB prevention:

- treatment of LTBI;
- prevention of transmission of *M. tuberculosis* through infection prevention and control; and
- vaccination of children with the bacille Calmette-Guérin (BCG) vaccine.

The three main sections of this chapter present and discuss progress in provision of these services. Particular attention is given to the 30 high TB burden countries and the 30 high TB/HIV burden countries (Chapter 2).

5.1 Treatment of latent TB infection

LTBI is defined as a state of persistent immune response to *M. tuberculosis* without clinically manifested evidence of active TB disease.

WHO guidelines published before 2018 identified four priority groups in which testing and treatment for LTBI was strongly recommended: people living with HIV (all countries); children aged under 5 years who are household contacts of bacteriologically confirmed pulmonary TB cases (all countries); people aged 5 years or more who are household contacts of pulmonary TB cases (upper middle-income and high-income countries with a low incidence of TB); and clinical risk groups (upper

middle-income and high-income countries with a low incidence of TB).^{4,5,6}

Updated guidance published in 2018 includes an additional recommendation to consider testing and treatment for people aged 5 years or more who are household contacts of bacteriologically confirmed pulmonary TB cases in countries with a high incidence of TB; it also expands the recommendation for clinical risk groups to all countries.⁷ Further details, including estimates of the number of people eligible for treatment based on the updated guidance, are provided in Box 5.1.

Data on the provision of TB preventive treatment for people living with HIV (specifically, those newly enrolled in HIV care⁸) have been collected annually by WHO for more than a decade, and for children aged under 5 years since 2016. In 2018, data for people aged 5 years or more were requested for the first time. The rest of this section presents and discusses the available data for these three priority groups.

Data for clinical risk groups – such as patients starting anti-tumour necrosis factor (TNF) therapy and those preparing for organ transplantation – are currently not requested as part of WHO's annual rounds of global TB data collection. Countries are nevertheless encouraged to monitor treatment initiation and completion, either routinely or through special surveys.

Routine collection of data about TB preventive treatment remains challenging. To facilitate faster, systematic and more complete data collection, WHO has developed a mobile phone application (app) which can be adapted at country level to record and report case-based data on TB preventive treatment.⁹

5.1.1 People living with HIV

Trends in the annual number of people living with HIV provided with TB preventive treatment 2005–2017 are shown in Fig. 5.1. Globally, the number was low until 2009; it then grew substantially to reach 932 663 in 2014, and has subsequently levelled off. The number in 2017

¹ World Health Organization. WHO End TB Strategy: global strategy and targets for tuberculosis prevention, care and control after 2015. Geneva: WHO; 2015 (http://www.who.int/tb/post2015_strategy/en/, accessed 30 July 2018).

² Houben RM, Dodd PJ. The global burden of latent tuberculosis infection: a re-estimation using mathematical modelling. *PLoS Med.* 2016;13(10):e1002152 (<https://www.ncbi.nlm.nih.gov/pubmed/27780211>, accessed 30 July 2018).

³ In an article published in 2000, the lifetime risk was estimated at 5–10%. See Vynnycky E, Fine PE. Lifetime risks, incubation period, and serial interval of tuberculosis. *Am J Epidemiol.* 2000; Aug 15;152(3):247–63.

⁴ World Health Organization. Guidelines for intensified tuberculosis case-finding and isoniazid preventive therapy for people living with HIV in resource-constrained settings. Geneva: WHO; 2011 (http://apps.who.int/iris/bitstream/10665/44472/1/9789241500708_eng.pdf, accessed 30 July 2018).

⁵ World Health Organization. Recommendations for investigating contacts of persons with infectious tuberculosis in low- and middle-income countries. Geneva: WHO; 2012 (http://www.who.int/tb/publications/2012/contact_investigation2012/en/, accessed 30 July 2018).

⁶ World Health Organization. Guidelines on the management of latent tuberculosis infection. Geneva: WHO; 2015 (http://www.who.int/tb/publications/ltbi_document_page/en/, accessed 30 July 2018).

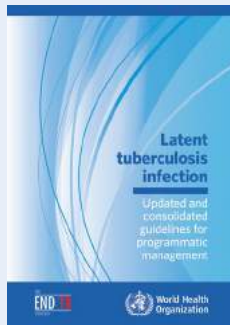
⁷ World Health Organization. Latent tuberculosis infection: updated and consolidated guidelines for programmatic management. Geneva: WHO; 2018 (<http://www.who.int/tb/publications/2018/latent-tuberculosis-infection/en/>, accessed 30 July 2018).

⁸ The indicator focuses on people newly enrolled in HIV care to help monitor trends in the programmatic management of LTBI. However, all people living with HIV should be regularly assessed for TB disease, and be offered preventive TB treatment if they are eligible for it.

⁹ http://www.who.int/tb/areas-of-work/preventive-care/ltbi/ltbi_app/en/

BOX 5.1

Updated WHO guidance on the programmatic management of latent TB infection issued in 2018, and associated estimates of the number of people eligible for TB preventive treatment from 2018–2022



WHO issued updated and consolidated guidelines for the programmatic management of LTBI in 2018.^a The document brings together guidance that was previously available in separate guidelines; it also includes updates to previous guidance. Recommendations in the guidelines are based on the probability

of progression to active TB disease in specific risk groups, the level and distribution of TB disease burden, the availability of resources, the likelihood of a broad public health impact, and the relative benefits and risks of treatment provision. The guiding principle is that individual benefits should outweigh risks.

The 2018 guidelines define three priority groups in which testing and treatment for LTBI is strongly recommended, in all countries:

- people living with HIV;
- infants and children aged under 5 years who are household contacts of bacteriologically confirmed pulmonary TB cases; and
- clinical risk groups, including patients initiating anti-TNF treatment, receiving dialysis, or preparing for organ or haematological transplantation, and those with silicosis.

In low-incidence settings, treatment for LTBI is also strongly recommended for HIV-negative people aged 5 years or more who are household contacts of bacteriologically confirmed pulmonary TB cases.

In high-incidence settings, the guidelines recommend considering testing and treatment for LTBI for HIV-negative people aged 5 years or more who are contacts of bacteriologically confirmed pulmonary TB patients, taking into account resource availability

and health system capacity. Compared with earlier guidance, this conditional recommendation substantially increases the potential number of people eligible for treatment.

Active TB disease should always be ruled out by screening for symptoms before prescribing preventive treatment. Chest radiography may also be considered for ruling out active TB disease among people living with HIV who are on ART, and among HIV-negative household contacts aged 5 years or more. The guidelines recommend testing for LTBI using either a tuberculin skin test or interferon-gamma release assay, in all countries. Lack of access to chest radiography, tuberculin skin testing, or interferon-gamma release assay should not be a barrier to initiation of treatment of LTBI.

The guidelines include four options for treatment of LTBI, including new shorter drug regimens:

- A weekly dose of rifapentine and isoniazid for 3 months.
- A daily dose of rifampicin plus isoniazid daily for 3 months.
- A daily dose of 3–4 months of rifampicin.
- A daily dose of isoniazid for at least six months.

Shorter regimens are expected to help patients to adhere to and complete treatment.

Based on the updated guidance, WHO estimates that at least 30 million people would be eligible for TB preventive treatment between 2018 and 2022. This represents an enormous increase compared with the numbers reported to have received TB preventive treatment in 2017 (Section 5.1).

^a World Health Organization. Latent tuberculosis infection: updated and consolidated guidelines for programmatic management. Geneva: WHO; 2018 (<http://www.who.int/tb/publications/2018/latent-tuberculosis-infection/en/>, accessed 30 July 2018).

was 958 559, based on data from 67 countries;¹ this included 640 201 people who were *newly* enrolled in HIV care, in 60 countries.²

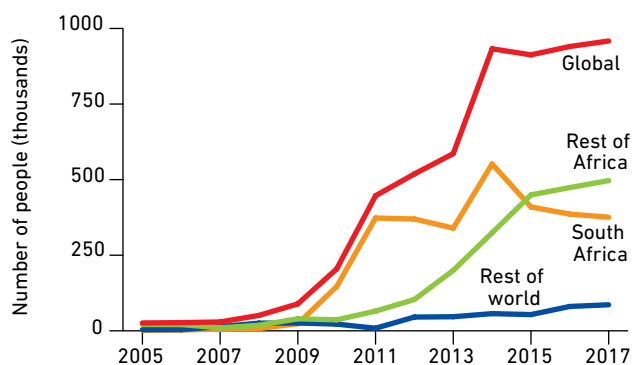
¹ These 67 countries accounted for 72% of the estimated number of TB cases among people living with HIV in 2017.

² Countries have regularly reported challenges in capturing data on TB preventive treatment among people newly enrolled in HIV care. In 2018, if countries could not report data for this group, they were asked to provide data on the number of people *currently* enrolled in HIV care who were started on TB preventive treatment.

As in previous years, South Africa accounted for the largest share (39%) of the global total. Large absolute increases in numbers compared with 2016 were reported in Nigeria (+29 000) and India (+10 000). For three of the 30 countries with a high burden of HIV-associated TB – Angola, Namibia and Papua New Guinea – data were reported for the first time in at least 4 years.

Despite progress in some countries, much remains to be done. For example, only 15 of the 30 high TB/HIV

FIG. 5.1
Provision of TB preventive treatment to people newly enrolled in HIV care,^a 2005–2017



^a For seven countries, data are for people currently enrolled in HIV care: Congo, Ecuador, Grenada, Kenya, Mozambique, Nepal and Ukraine.

burden countries reported some provision of TB preventive treatment to people living with HIV in 2017. Coverage among those newly enrolled in HIV care varied from 1% in Eswatini (formerly Swaziland) to 53% in South Africa (Table 5.1). Overall, in the 59 countries for which it could be calculated, coverage was 36%.

Gaps in the provision of TB preventive treatment to people living with HIV for selected high TB burden countries or high TB/HIV burden countries are illustrated in Fig. 5.2.

5.1.2 Children aged under 5 years who are household contacts of TB cases

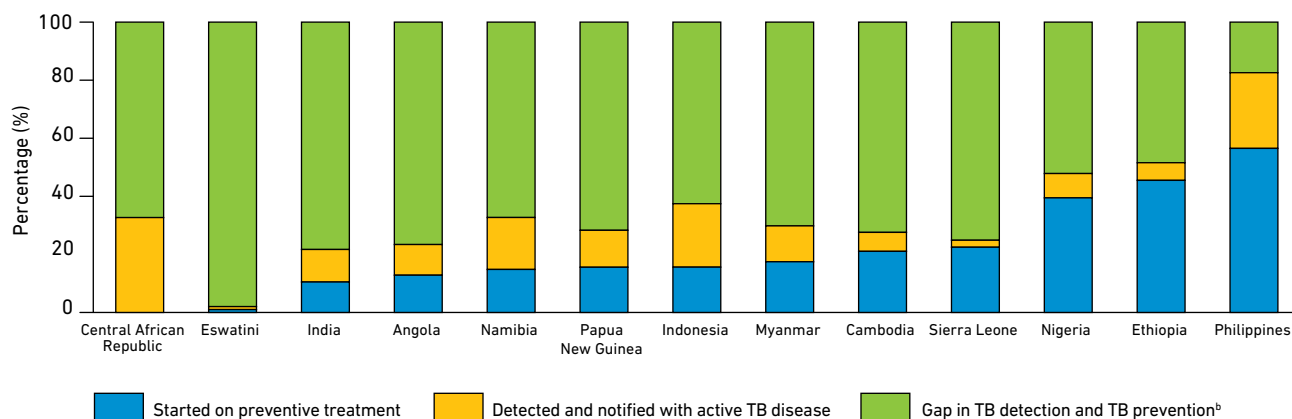
There were 189 countries that reported at least one notified bacteriologically confirmed pulmonary TB case

in 2017, of which 138 (73%) reported data about the number of household contacts aged under 5 years who were started on TB preventive treatment. Among these 138 countries, 124 reported that at least one child was started on preventive treatment in 2017 (up from 110 countries in 2016). This included 28 of the 38 high TB or high TB/HIV burden countries (Table 5.1) – an increase from 20 countries in 2016. Of these countries, data were reported to WHO for the first time by Botswana, Congo, Eswatini, Ethiopia, South Africa, Thailand and Zambia.

A total of 292 182 children aged under 5 years were reported to have been initiated on TB preventive treatment in 2017, an increase of 79% from 163 720 in 2016 and a more than three-fold increase from 87 242 in 2015. The largest numbers were reported by the WHO African Region (47% of the global total; 32 countries) and the WHO South-East Asia Region (25% of the global total; 9 countries). In the 28 high TB and TB/HIV burden countries that reported data, 199 630 children started TB preventive treatment (68% of the global total). At country level, India reported the largest number (38 745), followed by South Africa (32 104) (Table 5.1).

Globally, the 292 182 children started on TB preventive treatment in 2017 represented 23% of the approximately 1.3 million children estimated to be eligible for treatment. Higher levels of coverage were estimated for 20 countries in the WHO European Region (of which 13 reached coverage of ≥75%), followed by 23 countries in the WHO Region of the Americas (of which 13 reached coverage of ≥75%) and 16 countries in the Eastern Mediterranean Region (of which 9 reached coverage of ≥75%) (Fig 5.3). The 199 630 children started on treatment in the 28 high TB and TB/HIV burden countries that reported data represented 18% of the 1.1 million children estimated

FIG. 5.2
Gaps in TB prevention and TB detection for people who were newly enrolled in HIV care in 2017, selected countries^a



^a The selected countries are high TB or TB/HIV burden countries that reported on all three of the following: the number of people newly enrolled on HIV care; the number of TB cases detected among people newly enrolled on HIV care; and the number of people newly enrolled on HIV care who were started on TB preventive treatment. In high TB burden countries, testing for LTBI is not a requirement for initiation of TB preventive treatment, such that all those without active TB disease are eligible for TB preventive treatment.

^b The gap represents people living with HIV who should have undergone complete evaluation for TB disease or TB preventive treatment.

TABLE 5.1

TB preventive treatment for people living with HIV and children under 5 years of age who were household contacts of a bacteriologically confirmed pulmonary TB case, high TB or TB/HIV burden countries, 2017

	TREATMENT PEOPLE LIVING WITH HIV NEWLY ENROLLED IN CARE	PEOPLE LIVING WITH HIV NEWLY ENROLLED IN CARE STARTED ON TB PREVENTIVE		ESTIMATED NUMBER OF CHILD CONTACTS UNDER 5 YEARS OF AGE ELIGIBLE FOR TB PREVENTIVE TREATMENT ^a		CHILDREN UNDER 5 YEARS OF AGE STARTED ON TB PREVENTIVE TREATMENT		
		NUMBER	COVERAGE (%)	BEST ESTIMATE	UNCERTAINTY INTERVAL	NUMBER	COVERAGE ^b (%)	
							BEST ESTIMATE	UNCERTAINTY INTERVAL
Angola	22 137	2 842	13	24 800	(22 600–27 000)			
Bangladesh				54 400	(49 500–59 200)	11 394	21	(19–23)
Botswana				781	(711–850)	556	**	
Brazil				2 860	(1 850–3 860)			
Cambodia	3 582	754	21	5 540	(5 040–6 030)	2 410	44	(40–48)
Cameroon				10 300	(9 400–11 200)	89	0.86	(0.79–0.95)
Central African Republic	8 180	0	0	3 500	(3 190–3 810)			
Chad				4 520	(4 110–4 920)			
China				10 400	(6 710–14 000)			
Congo	1 380	272	*	2 410	(2 190–2 620)	98	4.1	(3.7–4.5)
DPR Korea				9 450	(8 610–10 300)	11 984	**	
DR Congo				84 400	(76 900–92 000)	13 045	15	(14–17)
Eswatini	23 271	232	1.0	1 200	(1 090–1 300)	89	7.4	(6.8–8.2)
Ethiopia	39 126	17 801	45	27 500	(25 100–30 000)	2 964	11	(9.9–12)
Ghana				3 710	(3 380–4 040)			
Guinea-Bissau				2 060	(1 880–2 250)	394	19	(18–21)
India	173 174	18 162	10	344 000	(314 000–375 000)	38 745	11	(10–12)
Indonesia	49 215	7 681	16	71 500	(65 200–77 900)	6 080	**	
Kenya	155 377	136 495	*	24 900	(22 700–27 100)	6 521	26	(24–29)
Lesotho				1 760	(1 610–1 920)			
Liberia				2 300	(2 100–2 510)	133	5.8	(5.3–6.3)
Malawi	168 972			4 530	(4 130–4 940)	3 068	**	
Mozambique	1 480 908	170 022	*	21 000	(19 100–22 800)	26 087	>100	
Myanmar	37 402	6 531	17	15 900	(14 500–17 300)	337	2.1	(1.9–2.3)
Namibia	21 213	3 143	15	3 150	(2 860–3 430)	1 069	34	(31–37)
Nigeria	232 822	91 873	39	52 500	(47 800–57 200)	10 258	**	
Pakistan				107 000	(97 100–116 000)			
Papua New Guinea	3 788	590	16	2 430	(2 210–2 640)			
Philippines	6 755	3 817	57	55 200	(50 200–60 100)	6 473	**	
Russian Federation	16 868	16 368	97	1 690	(1 100–2 290)	10 096	**	
Sierra Leone	23 693	5 323	22	7 690	(7 010–8 380)			
South Africa	710 083	375 650	53	40 800	(37 200–44 500)	32 104	79	(72–86)
Thailand				5 510	(5 020–6 000)	278	5.0	(4.6–5.5)
Uganda				20 700	(18 900–22 500)	1 734	8.4	(7.7–9.2)
UR Tanzania				21 000	(19 100–22 800)	7 303	35	(32–38)
VietNam	14 509	4 552	31	15 600	(14 200–17 000)	4 128	26	(24–29)
Zambia	234 586	42 961	18	10 600	(9 670–11 600)	404	3.8	(3.5–4.2)
Zimbabwe	164 963	17 618	**	7 610	(6 930–8 290)	1 789	24	(22–26)

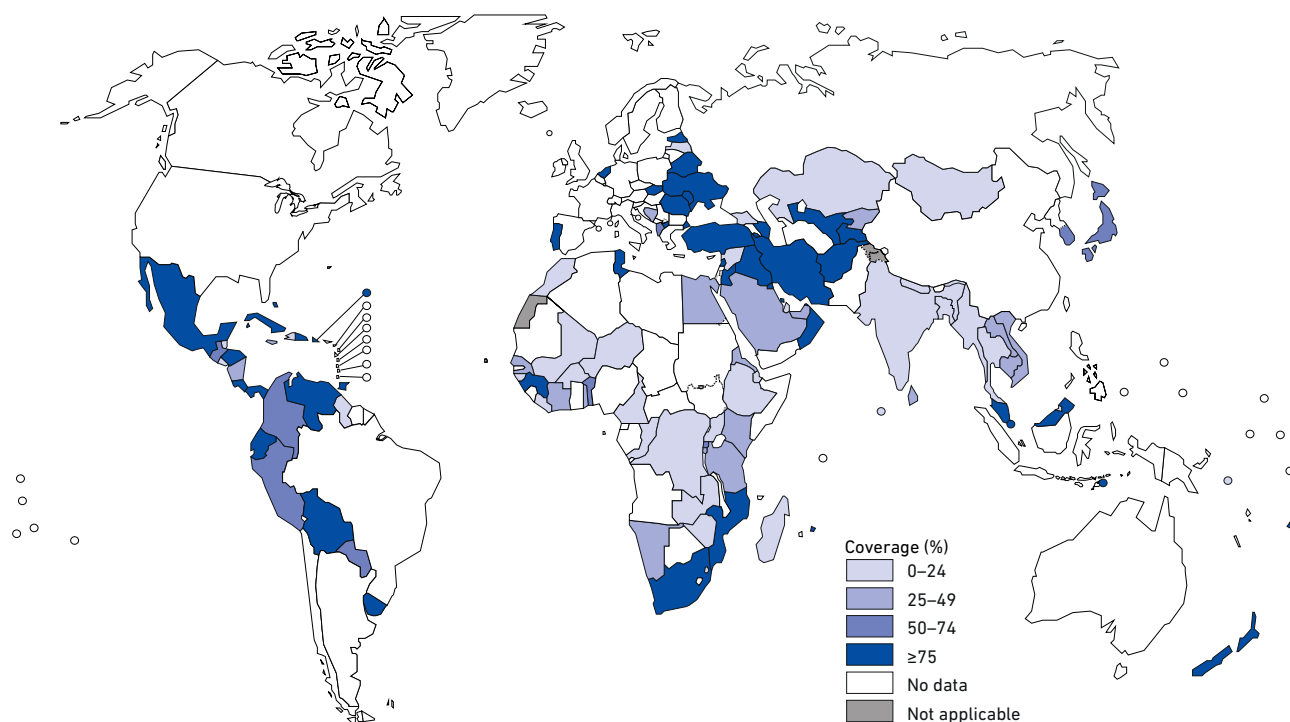
Blank cells indicate data not reported.

^a Estimates are shown to three significant figures.

^b Reasons for higher than expected coverage might be that the numerator did not exclude non-household contacts or children of five years and older.

* Coverage was not calculated because reported data on people living with HIV is for all enrolled in care, not just those newly enrolled in care.

** Coverage was not calculated because the numerator included contacts aged 5 years or older (Botswana, DPR Korea and Nigeria), those who were non-household contacts of TB cases (Indonesia and the Russian Federation), or those household contacts of clinically diagnosed TB cases (Malawi and the Philippines), or the number of PLHIV on TB preventive treatment was provided for the period July–December 2017 only (Zimbabwe).

FIG. 5.3**Coverage of TB preventive treatment among eligible children aged under 5 years,^a 2017**

^a Children aged <5 years who were household contacts of bacteriologically confirmed pulmonary TB cases. Estimated coverage was not calculated because the numerator included contacts aged 5 years or older (Botswana, DPR Korea and Nigeria), those who were non-household contacts of TB cases (Indonesia and the Russian Federation), or those household contacts of clinically diagnosed TB cases (Malawi and the Philippines).

to be eligible for treatment in the 38 high TB or TB/HIV burden countries. The highest coverage levels ($\geq 75\%$) in this group of countries were in Mozambique and South Africa (Fig 5.3 and Table 5.1).¹

5.1.3 Household contacts aged 5 years and older

In WHO's 2018 round of global TB data collection, data on provision of TB preventive treatment to household contacts aged 5 years or more were requested for the first time. Of the 189 countries that reported at least one notified bacteriologically confirmed pulmonary TB case in 2017, 78 (41%) reported data about the number of household contacts aged 5 years or more who were started on TB preventive treatment. Of the 78 countries, 70 reported that at least one contact aged 5 years or more was started on preventive treatment. This included eight countries that are in the lists of high TB burden, high multidrug-resistant TB (MDR-TB) or high TB/HIV burden countries: Azerbaijan, Kazakhstan, Kyrgyzstan, Republic of Moldova, Tajikistan, Thailand, Ukraine and Uzbekistan.

A total of 103 344 household contacts aged 5 years or more were reported to have been initiated on TB preventive treatment in 2017. The largest numbers were reported by the WHO European Region (56 335, 55% of the global total) and the WHO Eastern Mediterranean

Region (29 331, 28% of the global total). At country level, Afghanistan reported the largest number (22 939), followed by Ukraine (15 811) and Turkey (11 940).

5.2 TB infection prevention and control

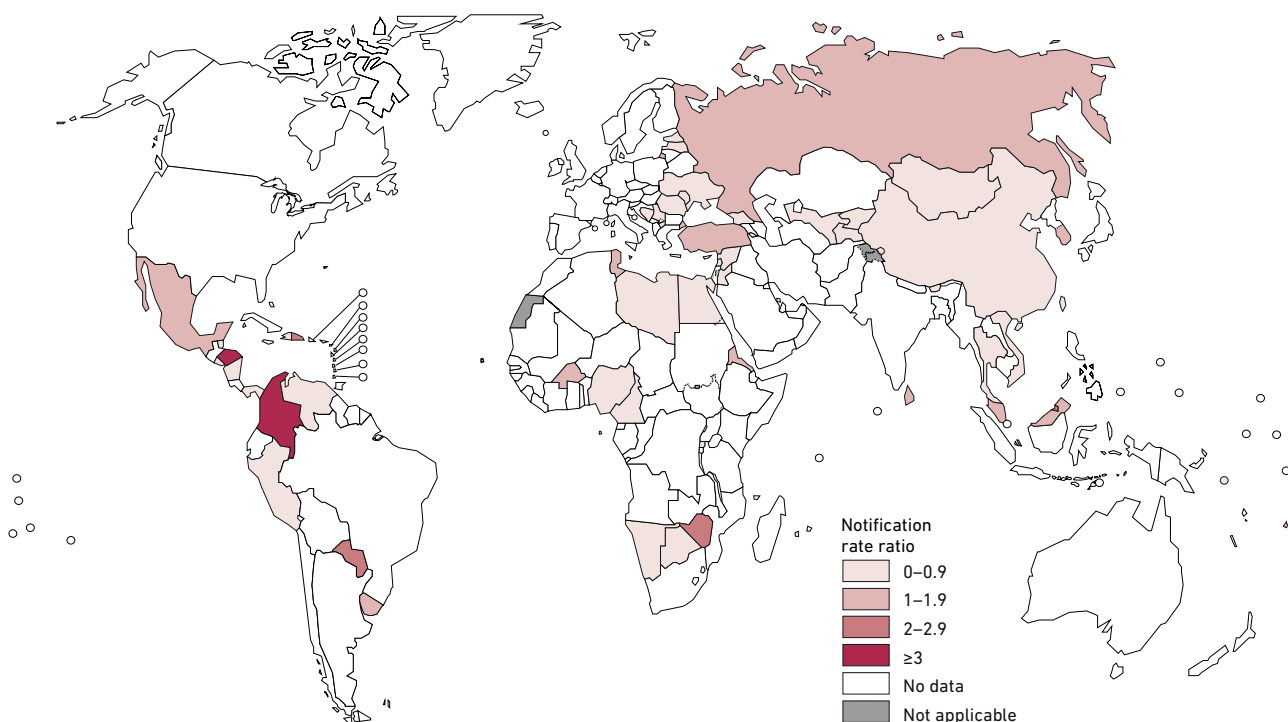
TB infection prevention and control is one of the components of Pillar 2 of the End TB Strategy (Chapter 2); it is also one of the collaborative TB/HIV activities that falls under Pillar 1. The risk of TB transmission is high in health-care and other congregate settings. This puts health-care workers at greater risk of TB infection and disease, and nosocomial outbreaks of MDR-TB and extensively drug-resistant TB (XDR-TB) among people living with HIV have been documented in the literature.^{2,3}

TB infection prevention and control should be part of national infection prevention and control policy. In health-care facilities and congregate settings, a comprehensive set of infection control measures – comprising administrative, environmental and personal protection measures

¹ Annex 1 explains how to access data reported by other countries.

² Gandhi NR, Weissman D, Moodley P, Ramathal M, Elson I, Kreiswirth BN et al. Nosocomial transmission of extensively drug-resistant tuberculosis in a rural hospital in South Africa. *J Infect Dis.* 2013;207(1):9–17.

³ Moro ML, Gori A, Errante I, Infuso A, Franzetti F, Sodano L et al. An outbreak of multidrug-resistant tuberculosis involving HIV-infected patients of two hospitals in Milan, Italy. *AIDS.* 1998;12(9):1095–102.

FIG. 5.4**Notification rate ratio of TB among health-care workers compared with the adult population (15–65 years), 2017^a**

^a Data from 3 countries were excluded where the number of health-care workers reported was less than 1000.

– should be implemented.¹ Periodic assessment of TB infection prevention and control in health-care facilities is essential to ensure that appropriate measures are in place.²

The risk of TB among health-care workers relative to the risk in the general adult population is one of the indicators recommended by WHO for measuring the impact of TB infection prevention and control activities in health-care facilities.³ If effective TB infection control measures are in place, the relative risk of TB in health-care workers compared with the general adult population should be close to one.

In 2017, 9299 TB cases among health-care workers were reported from 65 countries; China accounted for 35% of these cases and Brazil for 11%. The notification rate among health-care workers could be calculated for 58 of the 65 countries; it ranged from zero to 685 cases

per 100 000 health-care workers, with the highest rate observed in Mozambique.

The notification rate among the general adult population in each country was calculated based on the number of notified TB cases in adults and the estimated size of the adult population from the United Nations population division (2017 revision),⁴ restricted to those aged 15–64 for comparability with the health workforce. The ratios of the TB notification rate among health-care workers to the rate in the general adult population are shown in Fig. 5.4. The ratio was above two in four countries (Brunei Darussalam, Dominican Republic, Paraguay and Zimbabwe) and above three in Colombia and Honduras. The ratio was below one in four high TB burden countries: China, Namibia, Nigeria and Thailand.

5.3 TB vaccination

There is a clear need for a vaccine that is more effective than the BCG vaccine, in particular to reduce the risk of infection with *M. tuberculosis* and the risk of progression from infection to active TB disease in adults. Although there are 12 new candidates in the pipeline, it is unlikely that a new TB vaccine will be available in the immediate future (Chapter 8). To promote the development of

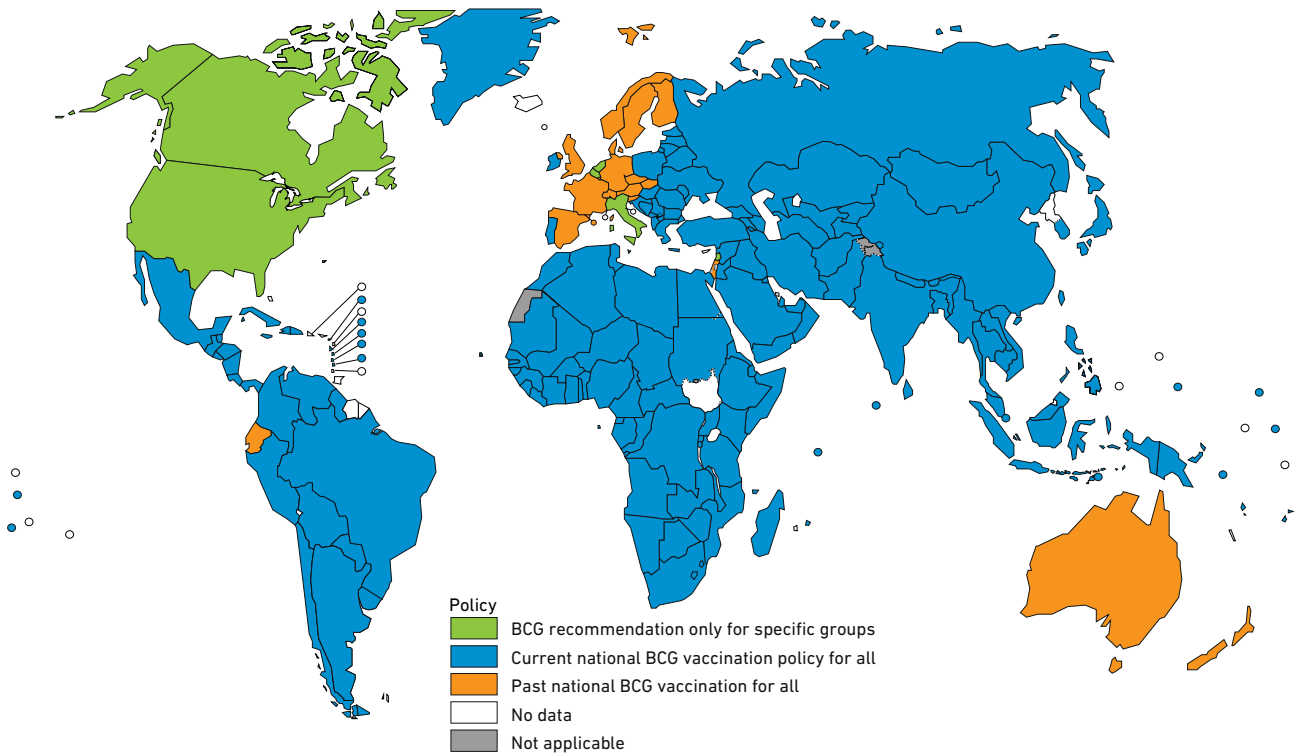
¹ World Health Organization. WHO policy on TB infection control in health-care facilities. Geneva: WHO; 2009 (http://www.who.int/tb/publications/2009/infection_control/en/, accessed 30 July 2018).

² World Health Organization. Checklist for periodic evaluation of TB infection control in health-care facilities. Geneva: WHO; 2015 (http://www.who.int/tb/areas-of-work/preventive-care/checklist_for_periodic_evaluation_of_tb_infection_control_in_health_facilities.pdf, accessed 30 July 2018).

³ World Health Organization. A guide to monitoring and evaluation for collaborative TB/HIV activities: 2015 revision (WHO/HTM/TB/2015.02). Geneva: WHO; 2015 (<http://www.who.int/tb/publications/monitoring-evaluation-collaborative-tb-hiv/en/>, accessed 30 July 2018).

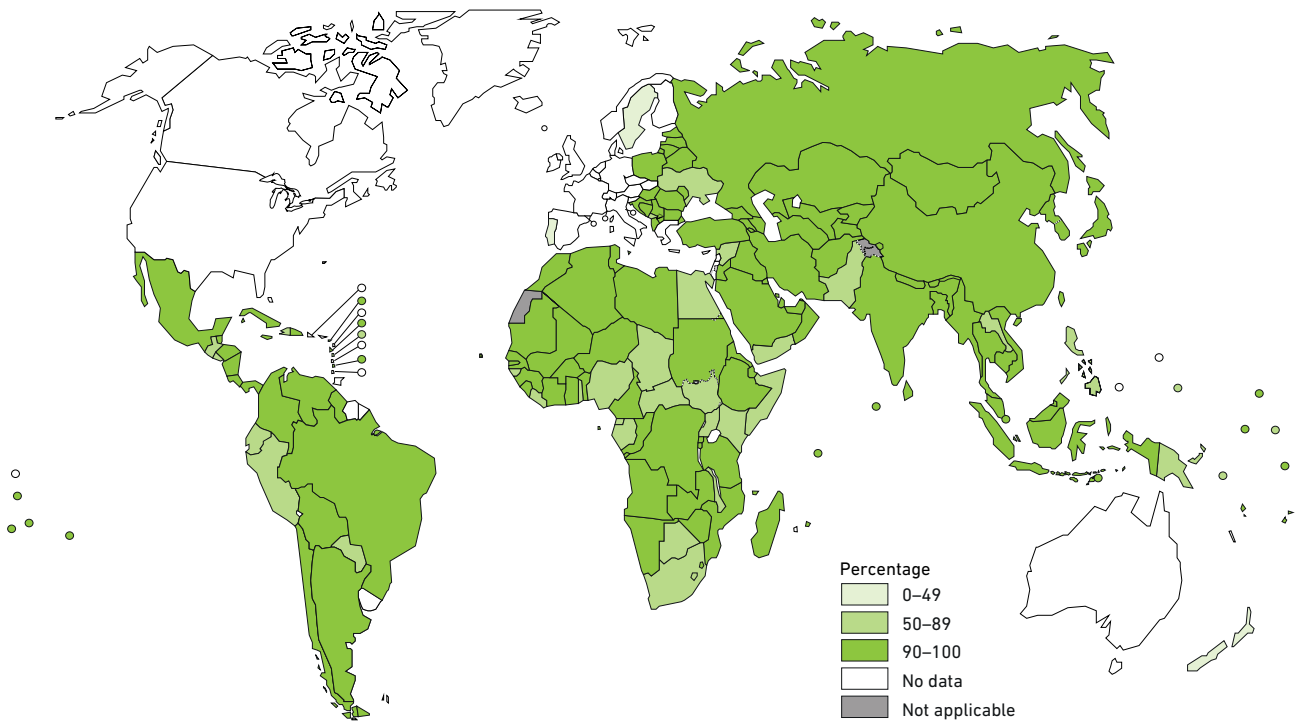
⁴ <https://esa.un.org/unpd/wpp/>

FIG. 5.5
BCG vaccination policy by country



Source: The BCG World Atlas 2nd Edition, <http://www.bcgatlas.org/>, accessed 19 July 2018.

FIG. 5.6
Coverage of BCG vaccination, 2017



The target population of BCG coverage varies depending on national policy, but is typically for the number of live births in the year of reporting.
 Source: http://apps.who.int/immunization_monitoring/globalsummary/timeseries/tscveragebcg.html, accessed 30 July 2018.

vaccines with optimal effectiveness and suitability in low- and middle-income countries, in 2018 the WHO Preferred Product Characteristics for New Tuberculosis Vaccines was developed through a multi-stakeholder process led by WHO's Initiative for Vaccine Research.¹

BCG vaccination has been shown to prevent disseminated disease; this category includes TB meningitis and miliary TB, which are associated with high mortality in infants and young children. Currently, WHO recommends that, in countries with a high TB burden, a single dose of the BCG vaccine should be provided to all infants as soon as possible after birth as part of childhood immunization programmes. In countries with low TB incidence rates, provision of the BCG vaccine may be limited to neonates and infants in recognized high-risk groups, or to older children who are skin-test negative for TB infection.

Fig. 5.5 summarizes national policies on BCG vaccination.² Among 180 countries for which data were collected, 154 recommended universal BCG vaccination; 20 reported having had a national BCG policy for everyone in the past, and the remaining six countries had policies of selective vaccination for at-risk individuals in high-risk groups.

The latest data on BCG coverage³ (for 2017) are shown in **Fig. 5.6**. In the 158 countries for which data are available, 120 reported coverage of at least 90%. Among the 30 high TB burden countries, coverage ranged from 53% in Nigeria to 99% in Bangladesh, Brazil, China, Mozambique, Thailand, the United Republic of Tanzania and Zambia; 21 reported coverage of at least 90%, up from 12 in 2016. In addition to Nigeria, coverage was below 80% in five other high TB burden countries: Central African Republic, Kenya, Lesotho, Papua New Guinea and South Africa.

¹ World Health Organization. WHO preferred product characteristics for new tuberculosis vaccine. Geneva: WHO; 2018 (http://www.who.int/immunization/documents/who_ivb_18.06/en/, accessed 30 July 2018).

² The BCG world atlas, 2nd edition, a database of global BCG vaccination policies and practices, 2nd edition. 2017 (<http://www.bcgatlas.org/>, accessed 19 July 2018).

³ World Health Organization. Reported estimates of BCG coverage. (http://apps.who.int/immunization_monitoring/globalsummary/timeseries/tscveragebcg.html, accessed 19 July 2018).



Volunteers sign in patients at a TB clinic in Dar es Salaam, United Republic of Tanzania
INTERFOTO / Alamy Stock Photo

Chapter 6. Financing for TB prevention, diagnosis and treatment

KEY FACTS AND MESSAGES

Financing for tuberculosis (TB) prevention, diagnosis and treatment continues to fall short of the amount needed and commitments from both domestic and international donor sources need to be stepped up.

The Stop TB Partnership's *Global Plan to End TB, 2016–2020*^a (the Global Plan) estimates that, in low- and middle-income countries, US\$ 10.4 billion is required in 2018, increasing to US\$ 12.3 billion in 2020. Based on data reported to WHO by 119 low- and middle-income countries with 97% of the world's notified TB cases, US\$ 6.9 billion is available for TB prevention, diagnosis and treatment in 2018, representing a shortfall of US\$ 3.5 billion.

The Global Plan estimates that an additional US\$ 2 billion per year is needed for TB research and development.

Of the total of US\$ 6.9 billion available in 119 low- and middle-income countries in 2018, US\$ 4.8 billion is for drug-susceptible TB and US\$ 2.0 billion is for MDR-TB; the remainder is for TB/HIV interventions or miscellaneous items.

Of the total of US\$ 6.9 billion available in 119 low- and middle-income countries in 2018, US\$ 6.0 billion (86%) is from domestic sources. However, this aggregate figure is strongly influenced by the BRICS group of countries (Brazil, the Russian Federation, India, China and South Africa).

BRICS accounted for 54% of the available funding for TB in 2018 (and 46% of the world's TB cases), with 96% (range 91–100%) of the group's funding coming from domestic sources. In other countries with a high TB burden, international donor funding remains crucial; for example, it accounted for 39% of the funding available in the 25 high TB burden countries outside BRICS (which have 40% of the world's notified TB cases) and for 57% of the funding in low-income countries.

International donor funding reported by national TB programmes (NTPs) dropped to US\$ 0.9 billion in 2018, falling far short of the US\$ 2.6 billion annual requirement estimated in the Global Plan. The single largest source (74% of the total) of funding reported by NTPs is the Global Fund to Fight AIDS, Tuberculosis and Malaria (the Global Fund).

International donor funding documented in the Organisation for Economic Co-operation and Development (OECD) creditor reporting system includes funding that flows through NTPs as well as other recipients. The total amount recorded in 2016 (the latest year for which data are available) was US\$ 0.9 billion, of which 69% was from the Global Fund (the Fund's contribution averaged 66% from 2006 to 2016). This was much less than funding for HIV (US\$ 6.8 billion) and malaria (US\$ 1.9 billion). To provide some context for these amounts, the latest estimates (for 2016) of the burden of disease in terms of disability-adjusted life years (DALYs) lost due to illness and death are 58 million for HIV/AIDS, 56 million for malaria and 44 million for TB.

The median cost per patient treated in 2017 was US\$ 1224 for drug-susceptible TB and US\$ 7141 for MDR-TB. The latter figure was lower than in previous years, following expanded use of a lower cost shortened treatment regimen that was first recommended by WHO in 2016.

Health financing data from national health accounts provide important insights into the current status of progress towards universal health coverage, as discussed in [Chapter 7](#).

^a The Global Plan to End TB, 2016–2020. Geneva: Stop TB Partnership; 2015 (<http://www.stoptb.org/global/plan/>, accessed 21 June 2018).

Progress in tuberculosis (TB) prevention, diagnosis and treatment requires adequate funding sustained over many years. WHO began annual monitoring of funding for TB in 2002, and has published findings in global TB reports and peer-reviewed publications.¹

This chapter has four main sections. It starts with a summary of the most up-to-date estimates of the financial resources required to achieve the 2020 milestones of the End TB Strategy (Section 6.1). It then presents and discusses trends in funding for TB prevention, diagnosis and treatment by category of expenditure and source of funding for the period 2006–2018, both globally and for major country groupings (Section 6.2). More detailed country-specific data for 2018 are also presented for the 30 high TB burden countries. The third section analyses funding gaps reported by national TB programmes (NTPs) to WHO, with breakdowns by category of expenditure and country group (Section 6.3). The final section provides the latest estimates (for 2017) of the unit costs of treatment for drug-susceptible TB and multidrug-resistant TB (MDR-TB) (Section 6.4).

As highlighted in the previous three editions of the *Global tuberculosis report*,^{2,3,4} analysis of health financing data can provide important insights into progress towards universal health coverage (UHC), which is necessary to achieve the End TB Strategy milestones set for 2020 and 2025 (Chapter 2). Measurement of costs faced by TB patients and their households is also required to assess progress towards one of the three high-level indicators of the End TB Strategy; that is, the percentage of TB patients and their households who face catastrophic costs as a result of TB disease. The 2020 milestone of zero set for this indicator requires progress towards UHC and social protection (included under Pillar 2 of the End TB Strategy). These two topics – analysis of health financing data, and measurement of costs faced by TB patients and their households – are discussed in Chapter 7.

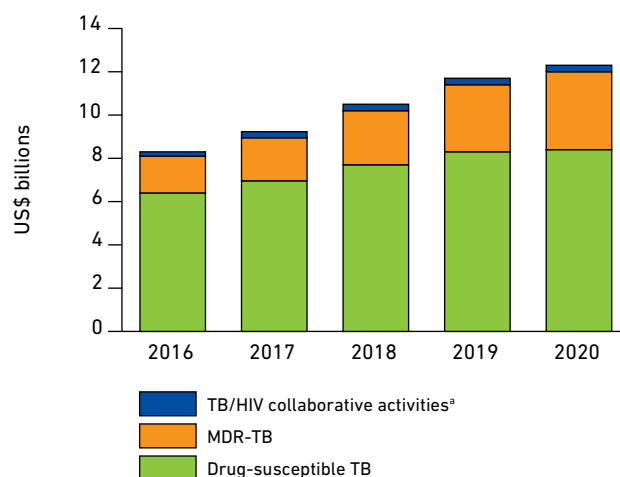
Further country-specific data on TB financing can be found in finance profiles that are available online.⁵

6.1 Estimates of funding required to achieve the 2020 milestones of the End TB Strategy

The 2020 milestones of the End TB Strategy are a 35% reduction in TB deaths compared with deaths in 2015, a 20% reduction in the TB incidence rate compared with 2015, and that no TB patients and their households face catastrophic costs as a consequence of TB disease (Chapter 2). Estimates of the funding required to achieve these milestones have been set out in the Stop TB Partnership's *Global Plan to End TB, 2016–2020* (the Global Plan).⁶ Worldwide, the total amount required for implementation of TB prevention, diagnostic and treatment interventions is US\$ 58 billion for the period 2016–2020, rising from US\$ 9.5 billion for the year 2016 to US\$ 14 billion for the year 2020.² An additional US\$ 9.0 billion is needed for global TB research and development in the same period.⁷

Of the US\$ 58 billion required over the 5 years from 2016 to 2020 (excluding research and development), an estimated US\$ 52 billion is required in low- and middle-income countries, growing from US\$ 8.3 billion in 2016 to US\$ 12 billion in 2020 (Fig. 6.1). In 2018, an estimated total of US\$ 10.4 billion is required: US\$ 7.7 billion (74%) for diagnosis and treatment of drug-susceptible TB, US\$ 2.5 billion for drug-resistant TB⁸ and the remainder for TB/

FIG. 6.1
Estimates of funding required for TB prevention, diagnosis and treatment in low- and middle-income countries in the Global Plan to End TB 2016–2020.



^a Funding estimates for TB/HIV exclude the cost of antiretroviral therapy (ART) for TB patients living with HIV. Such costs are included in global estimates of the funding required for HIV, published by UNAIDS.

Source: Stop TB Partnership Global Plan to End TB 2016–2020.

¹ The most recent publication is Floyd K, Fitzpatrick C, Pantoja A, Raviglione M. Domestic and donor financing for tuberculosis care and control in low-income and middle-income countries: an analysis of trends, 2002–11, and requirements to meet 2015 targets. *Lancet Glob Health*. 2013;1(2):e105–15 (<http://www.ncbi.nlm.nih.gov/pubmed/25104145>, accessed 11 July 2018).

² World Health Organization. *Global tuberculosis report 2015* (WHO/HTM/TB/2015.22). Geneva: WHO; 2015 (http://apps.who.int/iris/bitstream/10665/191102/1/9789241565059_eng.pdf, accessed 21 June 2018).

³ World Health Organization. *Global tuberculosis report 2016* (WHO/HTM/TB/2016.13). Geneva: WHO; 2016 (<http://apps.who.int/iris/bitstream/10665/250441/1/9789241565394-eng.pdf>, accessed 11 July 2018).

⁴ World Health Organization. *Global tuberculosis report 2017* (WHO/HTM/TB/2017.23). Geneva: WHO; 2017 (<http://apps.who.int/iris/bitstream/handle/10665/259366/9789241565516-eng.pdf>, accessed 21 June 2018).

⁵ www.who.int/tb/data

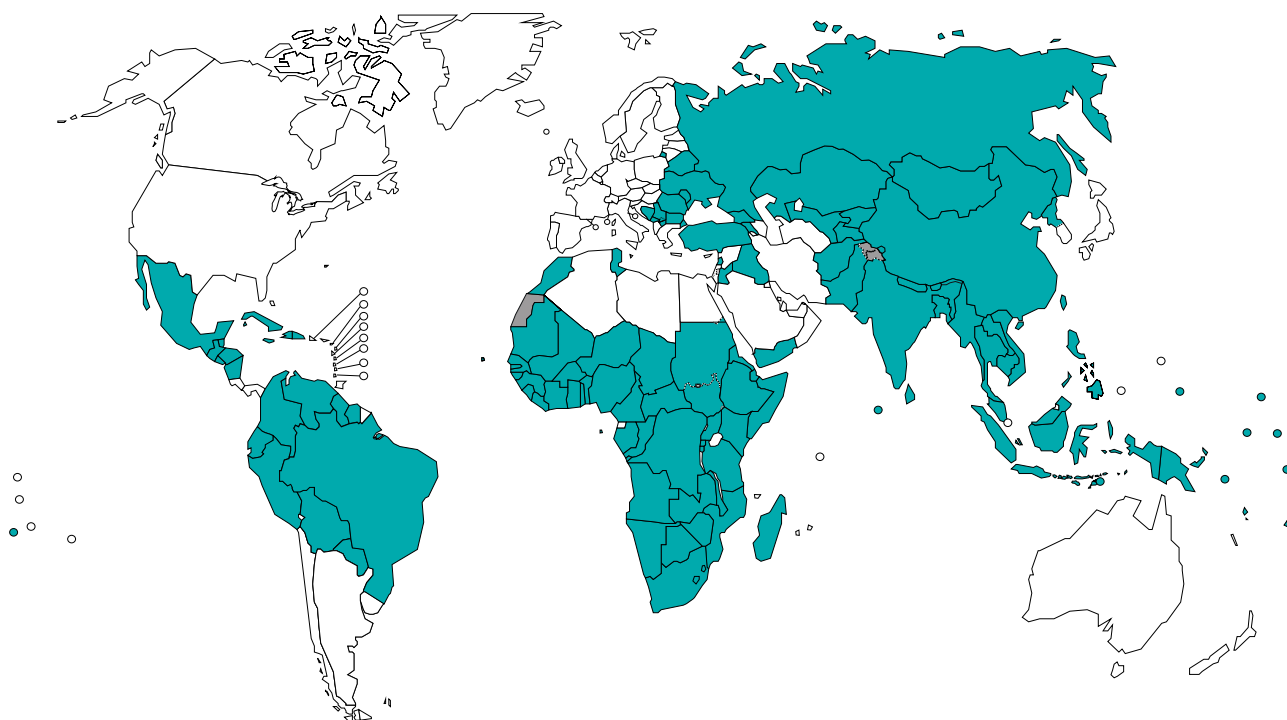
⁶ The Global Plan to End TB, 2016–2020. Geneva: Stop TB Partnership; 2015 (<http://www.stoptb.org/global/plan/>, accessed 21 June 2018).

⁷ Funding for TB research and development, which is monitored by the Treatment Action Group, is discussed further in Chapter 8.

⁸ The burden of drug-resistant TB (in terms of cases per year) is not projected to increase between 2016 and 2020. Increased funding is required to close detection and treatment gaps (see also Chapter 4).

FIG. 6.2

The 119 low- and middle-income countries included in analyses of TB financing, 2006–2018^a



^a Countries were included in trend analyses if at least three years of high-quality finance data were available in the period 2006–2018. Low-income (33/34), lower-middle income (44/46), and upper-middle income (42/56) countries representing 14%, 61% and 22% of 2017 notified cases, respectively, were included. The following 17 low- and middle-income countries were excluded: Albania, Azerbaijan, Costa Rica, Dominica, Algeria, Egypt, Equatorial Guinea, Grenada, Iran, Libya, St Lucia, Mauritius, West Bank and Gaza Strip, Syrian Arab Republic, Turkmenistan, St Vincent and Grenadines, Samoa.

HIV interventions. The amount for TB/HIV interventions is comparatively small because it does not include the funding needed for antiretroviral therapy for HIV-positive TB patients; this figure is instead included in estimates of funding required for HIV, published by the Joint United Nations Programme on HIV/AIDS (UNAIDS).¹ In 2018, in the context of preparations for the UN high-level meeting on TB, Global Plan projections of funding requirements were extended to 2022, suggesting that at least US\$ 13 billion will be needed in 2022.

In the Global Plan, estimates of the funding that could be mobilized from domestic and international donor sources were restricted to countries eligible to apply to the Global Fund to Fight AIDS, Tuberculosis and Malaria (the Global Fund).² For eligible countries, the funding required over 5 years amounted to US\$ 29 billion. Of this total, it was estimated that about US\$ 16 billion could be mobilized from domestic sources, and that the remainder (an average of US\$ 2.6 billion per year) would need to be provided by international donors.

The Global Plan did not attempt to assess the broader

investments required to increase the overall coverage and quality of health-care services or the funding required to remove financial barriers to accessing care. Such investments are needed for many essential preventive, treatment and care interventions, not only those related to TB. Progress on these fronts is critical, as reflected in Pillar 2 of the End TB Strategy. The costings in the Global Plan can thus be seen as the financial resources required for Pillars 1 and 3 of the End TB Strategy. Recent estimates of the funding required in low- and middle-income countries to make progress towards UHC by 2030 are presented and discussed in [Chapter 7](#).

6.2 TB funding, overall and by category of expenditure and source of funding, 2006–2018

Data reported by NTPs to WHO since 2006 were used to analyse funding trends for 2006–2018 in 119 low- and middle-income countries ([Fig. 6.2](#)). These countries accounted for 97% of the global number of TB cases notified in 2017. The methods used to compile, review, validate and analyse financial data are summarized in [Box 6.1](#).

In these 119 low- and middle-income countries, funding for TB prevention, diagnosis and treatment reached US\$ 6.9 billion in 2018, an increase from US\$ 6.2 billion

¹ World Health Organization. Global tuberculosis report 2016 (WHO/HTM/TB/2016.13). Geneva: WHO; 2016 (http://www.who.int/tb/publications/global_report/archive/en/, accessed 11 July 2018).

² Countries not eligible to apply to the Global Fund include Brazil, China, the Russian Federation and about half of the other 52 countries classified as upper-middle-income.

BOX 6.1

Methods used to compile, review, validate and analyse financial data reported to WHO

Overview

WHO began monitoring government and international donor financing for TB in 2002. All data are stored in the WHO global TB database. The standard methods used to compile, review, validate and analyse these data have been described in detail elsewhere;^{a,b} this box provides a summary.

Each year, WHO asks NTPs in all low- and middle-income countries to report:

- the funding they estimate will be needed for TB prevention, diagnosis and treatment in their current fiscal year, by category of expenditure and source of funding; and
- expenditures for the most recently completed fiscal year, also by category of expenditure and source of funding.

In the 2018 round of global TB data collection, the fiscal years were 2018 (for funding needs) and 2017 (for expenditures). Categories of expenditure used to report TB budget and expenditure data have been kept consistent as far as possible, to enable monitoring of trends.

Categories used for reporting of budgets and expenditures from 2002 to 2018

The categories used for annual reporting of funding needs (current fiscal year) and expenditures (last fiscal year) by NTPs in low- and middle-income countries are summarized below.

1. Drug-susceptible TB

- Laboratory infrastructure, equipment and supplies.
- NTP staff at central and subnational levels (e.g. NTP managers and provincial or district TB coordinators).
- First-line drugs.
- Programme costs; for example, management and supervision activities, training, policy development, meetings, purchase of office equipment and vehicles, recording and reporting of notifications and treatment outcomes, advocacy and communication, public-private mix activities and community engagement.
- Operational research, including surveys.
- Patient support – this category was first introduced as a stand-alone category in 2015, to reflect the emphasis on financial and social protection in the End TB Strategy.

2. MDR-TB

- Second-line drugs.
- Programme costs specifically related to MDR-TB.

3. TB/HIV

- Collaborative TB/HIV activities, including TB preventive treatment for people newly enrolled in HIV care. This category excludes any budget items that are financed by HIV programmes, such as antiretroviral therapy for TB patients living with HIV.

An “other” category is used to capture miscellaneous items that do not fit in one of the categories listed above.

Sources of funding

Low- and middle-income countries report on the breakdown of the total amount of available or committed funding by source, using four standard categories. These categories are domestic funding excluding loans; external loans, also considered domestic funding; the Global Fund; and grant financing from sources other than the Global Fund.

High-income countries

As in previous years, in 2018, all high-income countries were asked to report their funding requirements and expenditures in total, without any breakdown by category of expenditure or source of funding. However, of the 73 high-income countries, only seven reported expenditures and only 23 reported the amount of funding needed in 2018. Total TB funding trends for 2006–2018 are available for eight of the 73 high-income countries; however, those countries are not featured in this chapter given its focus on low- and middle-income countries.

Average cost of drugs per patient (since 2014)

Since 2014, an extra question about the average cost of drugs per patient treated has been asked, to allow reviewers to better assess the validity of budgets reported for first-line and second-line drugs, and to identify whether reported budgets include funding for buffer stocks.

Use of general health services (2002–2018)

Annually since 2002, all countries (irrespective of income level) have been asked to report on the use of inpatient and outpatient care for treatment of people with drug-susceptible TB and MDR-TB on a per-patient basis (i.e. the average number of days spent in hospital, and the average number of outpatient visits to a health facility). These data can be based on actual use of services (preferable where available), or on the expected use of services based on the typical approach used to deliver treatment (which may be defined in national policy documents and protocols). These data on health service use are then combined with other data to estimate the financial resources used for TB treatment that are not reflected in NTP-reported budgets and expenditures (further details are provided below).

Data validation by WHO's Global TB Programme

The core methods used to review and validate data have remained consistent since 2002. They include the following:

- routine checks for plausibility and consistency, including validation checks that are built into the online reporting system – examples of validation checks are checks for implausibly large year-to-year changes (e.g. in total reported funding by source and by category of expenditure), or implausibly high or low values of funding for drugs relative to the number of TB patients (that differ substantially from prices quoted by the Global TB Drug Facility);
- discussions with country respondents to resolve queries; and
- triangulation with other data sources – these include estimates of unit costs from independent economic

evaluations (Box 6.3)^c and data extracted by the Global Fund from funding applications submitted to them (comprehensive budgets for national strategic plans for TB are an essential requirement for funding applications to the Global Fund); further details about the comparisons with other data sources are available from WHO upon request.

In review and validation of data, particular attention has always been given to the high TB burden countries.

In 2018, specific efforts to improve the quality of financial data reported to WHO included discussions with NTP staff during a workshop on TB budgeting; and individual and customized follow-up with in-country staff involved in the development of national strategic plans and reporting of financial data.

Except for the Russian Federation, China, South Africa in 2018 and Viet Nam in 2017, TB funding reported by NTPs in 2017 and 2018 did not include the financial costs associated with the inpatient and outpatient care required during TB treatment. Since many detailed costing studies in numerous countries show that these costs can account for a large share of the cost of treating someone with TB, WHO analyses of TB financing have always included estimates of the funding required for both inpatient and outpatient care. These costs have been estimated from a provider perspective only, and do not include the costs faced by TB patients and their households. Increasing attention is now being given to costs faced by TB patients and their households, as discussed in [Chapter 7](#).

Estimates of the costs of inpatient and outpatient care for patients with drug-susceptible TB or MDR-TB

To estimate the funding used to provide inpatient and outpatient care for TB patients, WHO multiplies the number of outpatient visits and days of inpatient care per patient (reported by NTPs each year, as explained above) by the cost per bed day and per clinic visit available from the WHO CHOosing Interventions that are Cost-Effective (WHO-CHOICE) database,^d and then by the reported number of TB patients notified or projected to be notified. These estimates are done separately for drug-susceptible TB and MDR-TB. In 2018, costs per bed day and per clinic visit were estimated using the CHOICE regression model and the latest data available from the World Bank. For two countries (India and Thailand), WHO-CHOICE estimates were replaced with estimates of unit costs obtained directly from national health account data, or from recent studies and discussions with experts supporting the costing of national strategic plans.

Where possible, estimates are compared with hospital and clinic expenditure data for drug-susceptible TB and MDR-TB that are being tracked through the System of Health Accounts (SHA).^e In 2018, SHA data were available for 27 countries for 1 or 2 years, including six high burden countries (Cambodia, the Democratic Republic of the Congo, Namibia, the Philippines, Sierra Leone and the United Republic of Tanzania).^f After review, the SHA data were used in preference to estimates based on the combination of reported use and WHO-CHOICE unit cost estimates. The WHO Health Governance and Financing Department has initiated a process to assess the validity of the latest results from the new SHA, including disease-specific results.

Expanded implementation of SHA and validation against existing disease-specific tracking systems may facilitate more comprehensive reporting of domestic funding for TB. In particular, it may facilitate reporting of the contributions from subnational administrative levels that are not always known or compiled at the national level. Although much of this contribution is probably for delivery of inpatient and outpatient care (which is included in current WHO estimates of domestic funding for TB, as explained above), reporting of funding from these levels (including TB-specific budgets) is a particular challenge in large countries with decentralized systems for TB treatment (e.g. Indonesia, Nigeria and South Africa).

Estimates of the cost per patient treated for drug-susceptible TB and MDR-TB

Since 2014, WHO has been reporting estimates of the costs per patient treated for drug-susceptible TB and MDR-TB. Costs are calculated separately for drug-susceptible TB and MDR-TB. In each case, the numerator is the total estimated cost of treatment, which has two main parts – the national expenditures reported by the NTP, and the costs associated with the use of health services for TB patients – with data validated as described above. Categories of expenditure considered as costs for MDR-TB were second-line drugs and all other inputs used or activities implemented for the programmatic management of MDR-TB. All other categories (except collaborative TB/HIV activities) were assumed to be for drug-susceptible TB. An exception was made for the Russian Federation, for which expenditures for staff and infrastructure were allocated by WHO to drug-susceptible TB (33%) and MDR-TB (67%), based on the proportion of bed days used for these two categories of patients.

Unit costs were then calculated as the sum of 2017 NTP expenditures and total costs for use of inpatient and outpatient care, divided by the reported number of patients treated. Again, this calculation was carried out separately for drug-susceptible TB and MDR-TB.

- ^a Floyd K, Pantoja A, Dye C. Financing tuberculosis control: the role of a global financial monitoring system. *Bull World Health Organ.* 2007;85(5):334–40 (<http://www.ncbi.nlm.nih.gov/pubmed/17639216>, accessed 12 July 2018).
- ^b Floyd K, Fitzpatrick C, Pantoja A, Raviglione M. Domestic and donor financing for tuberculosis care and control in low-income and middle-income countries: an analysis of trends, 2002–11, and requirements to meet 2015 targets. *Lancet Glob Health.* 2013;1(2):e105–15 (<http://www.ncbi.nlm.nih.gov/pubmed/25104145>, accessed 11 July 2018).
- ^c Global Health Cost Consortium unit cost study data repository, accessed 12 July 2018 (<http://ghcosting.org/pages/data/ucsr/app/index>).
- ^d Cost effectiveness and strategic planning (WHO-CHOICE): health service delivery costs. Geneva: World Health Organization; 2008 (http://www.who.int/choice/cost-effectiveness/inputs/health_service/en/, accessed 12 July 2018).
- ^e OECD/Eurostat/WHO. A system of health accounts. OECD Publishing; 2011 (<http://www.who.int/health-accounts/methodology/sha2011.pdf>, accessed 12 July 2018).
- ^f Health accounts. Geneva: World Health Organization; data shared by WHO Health Governance and Financing Department.

FIG. 6.3
Funding for TB prevention, diagnosis and treatment in total and by category of expenditure, 2006–2018, 119 countries with 97% of reported cases

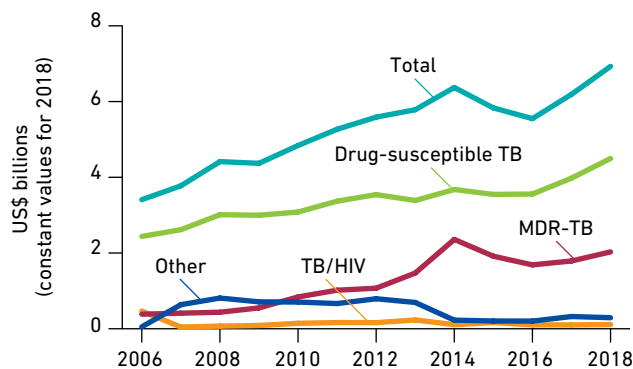
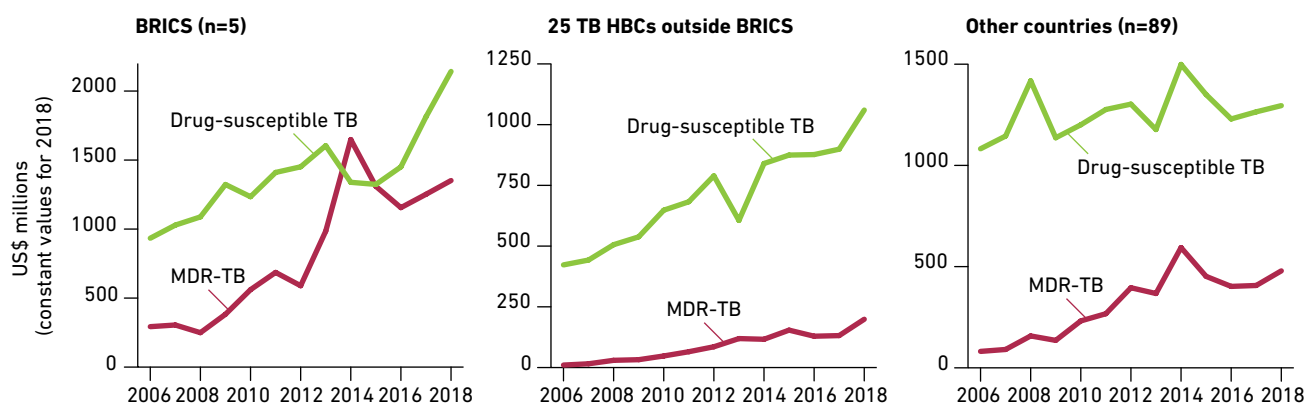


FIG. 6.4
Funding for drug-susceptible TB and MDR-TB, 2006–2018, by country group^a



^a BRICS accounted for 46% of the total number of TB cases notified globally in 2017. The 25 high TB burden countries outside BRICS accounted for 40%. The remaining countries (n=89) included in financing analyses accounted for 11% of the TB cases notified globally in 2017.

in 2017 and more than double the US\$ 3.4 billion that was available in 2006 (Fig. 6.3; all figures are in constant 2018 US dollars). Despite this growth in funding, amounts continue to fall short of what is needed. Moreover, the shortfall compared with the Global Plan estimate (Section 6.1) has widened compared with 2016 and 2017, increasing from US\$ 2.3 billion in 2017 to US\$ 3.5 billion in 2018 (US\$ 6.9 billion available compared with an estimated requirement of US\$ 10.4 billion).

Of the total of US\$ 6.9 billion available in 2018, US\$ 4.8 billion (69%) is for the diagnosis and treatment of drug-susceptible TB. This is US\$ 2.9 billion less than the requirement of US\$ 7.7 billion estimated in the Global Plan.

Funding for MDR-TB reached US\$ 2.0 billion in 2018; the annual amount increased steadily from 2006 to 2014, declined from 2014 to 2016, and has been increasing since 2016 (Fig. 6.3). This aggregate trend reflects the pattern in the BRICS group of countries (Brazil, Russian Federation, India, China and South Africa) (Fig. 6.4),

where a downturn after 2014 is explained by decreases in funding in the Russian Federation (2014–2016) and in South Africa (2013–2014 and 2017–2018). Funding for MDR-TB in other countries decreased from 2014 to 2017 and then increased in 2018 (Fig. 6.4).

Although the gap between the funding available in 2018 and the requirement of US\$ 2.5 billion in 2018 estimated in the Global Plan is relatively small (US\$ 0.5 billion), it has widened over the past 2 years. This is consistent with the need to increase the coverage of diagnosis and treatment for MDR-TB, which currently falls far short of Global Plan targets both overall and in most countries with a high burden of MDR-TB (for details, see Chapter 4). It is also evident from the Global Plan estimates of funding required in future years: the annual funding required for MDR-TB will reach US\$ 3.6 billion

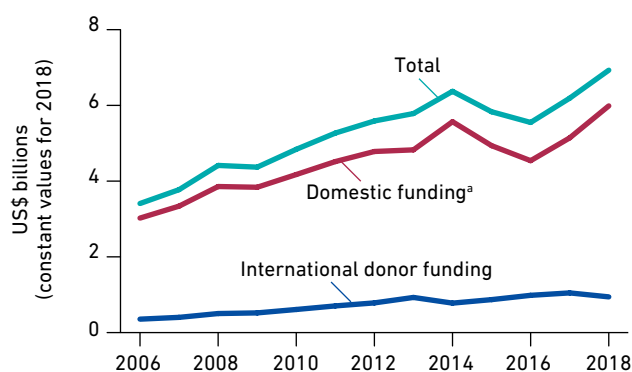
in 2020, substantially higher than the amount of US\$ 2.0 billion available in 2018.

Overall, most funding during the period 2006–2017 has been provided from domestic sources, and this remains the case in 2018 (Fig. 6.5).¹ In 2018, US\$ 6.0 billion (86%) of the total funding of US\$ 6.9 billion for TB is from domestic sources. However, aggregated figures for the 119 low- and middle-income countries are strongly influenced by BRICS, and they conceal substantial variation among countries in the share of funding from domestic and international sources (Fig. 6.6).

BRICS accounted for 54% of the available funding for

¹ Domestic funding includes both funding for TB-specific budgets, and funding for inpatient and outpatient care (usually funded through more general budget lines), as also explained in Box 6.1. In Fig. 6.5 and Fig. 6.6, it is assumed that funding for inpatient and outpatient care is provided domestically rather than by international donors. This is justified on the basis that most (99%) of the funding estimated to be used for inpatient and outpatient care for TB patients is accounted for by middle-income countries, where international donor funding for such components of care is unlikely (such support is more likely to occur in low-income countries, via general budget support to the health sector).

FIG. 6.5
Funding for TB prevention, diagnosis and treatment by funding source, 2006–2018, 119 countries with 97% of reported cases



^a Domestic funding includes TB-specific budgets and the estimated resources used for inpatient and outpatient care (see [Box 6.1](#)). 93% of the funding of US\$ 2.3 billion for inpatient and outpatient care for 2018 is accounted for by middle-income countries; such countries do not typically receive international donor funding for inpatient and outpatient care services.

TB in 2018 (and 46% of the world's notified TB cases), with 96% (range 91–100%) of their funding coming from domestic sources. As highlighted in the 2017 *Global tuberculosis report*,¹ in India there has been a particularly striking increase in the TB-specific budget and domestic funding for this budget since 2016 ([Fig. 6.7](#)). Between 2016 and 2018, domestic funding for the national budget for TB in India more than quadrupled, from US\$ 110 million in 2016 to US\$ 458 million in 2018.

Domestic funding also dominates in seven of the nine (not mutually exclusive) country groups shown in [Fig. 6.6](#), ranging from 60% in Africa to 97% in upper-middle-income countries. In contrast, international donor funding continues to exceed funding from domestic sources in low-income countries (57% of the total in 2018). In the 25 high TB burden countries excluding BRICS,² the share of funding from domestic sources increased to 61% in 2018, up from 55% in 2017. In this group, the countries with notable increases in funding from domestic sources between 2017 and 2018 were Cambodia, Indonesia, Lesotho, Papua New Guinea and Philippines.³

International donor funding reported by NTPs to WHO amounted to US\$ 0.9 billion in 2018, a slight decline compared with 2017. Of this amount, most (74%) was provided by the Global Fund.

The importance of international donor funding in high TB burden countries is particularly evident when consid-

ering only the TB-specific budgets included in national strategic plans for TB ([Fig. 6.8](#), [Table 6.1](#) and [Table 6.2](#)). In 17 of the 30 high TB burden countries, more than 50% of available funding for the TB-specific budgets included in national strategic plans for TB is from international donors in 2018.

Both [Fig. 6.7](#) and [Fig. 6.8](#) illustrate the potential to increase domestic funding in some high TB burden countries.⁴ In the group of nine low-income high TB burden countries, the proportion of the TB budget funded from domestic sources in 2018 ranges from 0% in Zimbabwe to 14% in Central African Republic. In the group of 15 lower-middle-income high TB burden countries, the proportion ranges from 3% in Pakistan to 79% in India. In the group of six upper-middle-income countries, the proportion ranges from 85% in Brazil to 100% in the Russian Federation.

Funding reported by NTPs to WHO does not capture all international donor funding for TB.⁵ For this reason, a complementary analysis based on donor reports to the Organisation for Economic Co-operation and Development (OECD) is provided in [Box 6.2](#).⁶

6.3 Funding gaps reported by national TB programmes, 2006–2018

Many NTPs reported a shortfall in the funding required for full implementation of their national strategic plans ([Fig. 6.8](#), [Fig. 6.9](#) and [Table 6.2](#)). Funding gaps (i.e. the difference between assessments by NTPs of funding needs for TB prevention, diagnosis and treatment, and the actual amount of funds mobilized) have increased, and in 2018 they amounted to a reported total of US\$ 1.2 billion. This is almost a third of the gap of US\$ 3.5 billion that exists between the US\$ 10.4 billion estimated to be needed in low- and middle-income countries in 2018 according to the Global Plan ([Section 6.1](#)) and the US\$ 6.9 billion available in 2018 ([Section 6.2](#)). The difference can be explained by the fact that, in many countries, national strategic plans for TB are less ambitious than the targets set in the Global Plan ([Section 6.1](#)).

Of the US\$ 1.2 billion funding gap reported by NTPs in 2018, US\$ 0.99 billion (80%) is for drug-susceptible TB and US\$ 0.24 billion (20%) is for MDR-TB. Relative to total funding needs, the funding gap is larger for drug-susceptible TB than for MDR-TB.

Lower-middle-income countries account for the largest reported funding gap (US\$ 826 million) in 2018

¹ World Health Organization. *Global tuberculosis report 2017* (WHO/HTM/TB/2017.23). Geneva: WHO; 2017 (<http://apps.who.int/iris/bitstream/handle/10665/259366/9789241565516-eng.pdf>, accessed 21 June 2018).

² The list of 30 high TB burden countries being used by WHO during the period 2016–2020 is explained in [Chapter 2](#). The countries are those listed in [Fig. 6.8](#), [Table 6.1](#) and [Table 6.2](#).

³ These figures exclude domestic funding for inpatient and outpatient care during treatment. For further details for these countries, see the country profiles in [Annex 2](#).

⁴ Sustained and increased financing was one of the four topics of the Moscow Declaration to End TB; First WHO Global Ministerial Conference on Ending TB in the Sustainable Development Era: A Multisectoral Response, November 2017. Geneva: WHO and the Ministry of Health of the Russian Federation; 2017 (http://www.who.int/tb/Moscow_Declaration_MinisterialConference_TB/en/, accessed 21 June 2018).

⁵ Donor funding is also provided to entities other than NTPs, including international and national organizations, both governmental and nongovernmental.

⁶ Out-of-pocket expenditures are also not included in the financing data reported by NTPs. These are discussed in more detail in [Chapter 7](#).

TABLE 6.1

Reported budget in national strategic plans for TB, by intervention area and estimated cost of inpatient and outpatient care for drug-susceptible (DS-TB) and MDR-TB, 30 high TB burden countries, 2018 (current US\$ millions)

	TOTAL BUDGET IN NATIONAL STRATEGIC PLAN FOR TB	DS-TB	MDR-TB	TB/HIV	INPATIENT AND OUTPATIENT CARE (DS-TB)	INPATIENT AND OUTPATIENT CARE (MDR-TB)	ESTIMATED TOTAL RESOURCES REQUIRED FOR TB CARE
Angola	46	40	5.1	1.0	20	7.9	73
Bangladesh	66	64	2.7	<0.1	6.2	2.4	75
Brazil	57	49	6.1	2.0	23	1.5	81
Cambodia	37	34	2.4	0.9	28	0.5	65
Central African Republic	1.9	1.5	0.3	<0.1	0.6	<0.1	2.5
China ^a	609	492	117	0	—	—	609
Congo	1.5	1.4	0.1	0	0.3	<0.1	1.8
DPR Korea	84	78	5.6	0	46	4.7	134
DR Congo	74	56	13	5.1	13	4.8	92
Ethiopia	93	74	12	7.4	12	1.4	106
India	580	390	175	15	662	145	1 386
Indonesia	294	270	20	4.9	47	21	362
Kenya	42	36	2.9	3.0	7.3	1.3	51
Lesotho	3.3	1.9	1.4	0	0.2	0	3.6
Liberia	6.4	6.4	0	0	0.2	0.3	6.9
Mozambique	30	17	7.9	5.7	4.9	0.4	36
Myanmar	58	50	5.4	2.0	1.8	0.4	60
Namibia	49	48	1.2	0.2	4.9	9.0	63
Nigeria	409	285	109	15	6.0	16	431
Pakistan	118	82	36	0.2	7.2	0.4	126
Papua New Guinea	28	18	3.3	7.1	0.9	1.7	31
Philippines	160	142	17	0.8	116	18	294
Russian Federation ^{a,b}	1 435	530	904	1.2	—	—	1 435
Sierra Leone	16	12	2.8	1.1	18	0.8	34
South Africa ^a	271	227	34	9.2	—	—	271
Thailand ^c	26	20	5.5	<0.1	10	0.2	36
UR Tanzania	64	50	8.7	5.0	3.3	1.0	68
Viet Nam	70	55	13	1.8	24	2.8	96
Zambia	38	29	4.7	3.6	1.9	1.2	41
Zimbabwe	30	26	1.0	2.5	1.8	0.4	32
30 high TB burden countries	4 795	3 184	1 517	95	1 063	243	6 101

Subtotals do not always add to the total due to rounding.

— indicates values that cannot be calculated.

^a No amounts for the additional resources required for inpatient and outpatient care are shown for China, the Russian Federation and South Africa because the NTP budgets reported by those countries include all budgets for inpatient and outpatient care.

^b In the Russian Federation, the staff and infrastructure reported for TB care and control were allocated to DS-TB (33%) and MDR-TB (67%) by WHO based on the proportion of beddays used by DS-TB and MDR-TB patients.

^c In 2018, the budget reported by Thailand was for the central level only.

TABLE 6.2

Reported budget in national strategic plans for TB, available funding for this budget from domestic and international donor sources and funding gap, 30 high TB burden countries, 2018 (current US\$ millions)

	TOTAL BUDGET IN NATIONAL STRATEGIC PLAN FOR TB	DOMESTIC FUNDING (A)	INTERNATIONAL DONOR FUNDING (B)	SHARE OF AVAILABLE FUNDING (A+B) PROVIDED FROM DOMESTIC SOURCES (%)	SHARE OF AVAILABLE FUNDING (A+B) PROVIDED BY INTERNATIONAL DONORS (%)	FUNDING GAP ^b
Angola	46	3.3	5.7	37	63	37
Bangladesh	66	9.5	37	21	79	20
Brazil	57	48	0	100	0	8.7
Cambodia	37	4.6	8.7	34	66	24
Central African Republic	1.9	0.3	1.0	21	79	0.6
China	609	528	13	98	2.4	68
Congo	1.5	1.1	0.4	72	28	0
DPR Korea	84	5.7	0.9	86	14	77
DR Congo	74	1.5	36	4.0	96	37
Ethiopia	93	10	30	25	75	52
India	580	458	121	79	21	0
Indonesia	294	102	48	68	32	145
Kenya	42	14	13	52	48	15
Lesotho	3.3	1.0	1.8	36	64	0.5
Liberia	6.4	0.2	6.2	2.4	98	0
Mozambique	30	1.5	29	4.9	95	0
Myanmar	58	2.1	30	6.7	93	26
Namibia	49	28	1.4	95	5	20
Nigeria	409	32	65	33	67	312
Pakistan	118	3.6	64	5.4	95	51
Papua New Guinea	28	14	6.0	70	30	8.1
Philippines	160	59	34	63	37	66
Russian Federation	1 435	1 435	0	100	0	0
Sierra Leone	16	0.3	12	2.2	98	3.5
South Africa	271	247	22	92	8.0	1.6
Thailand ^a	26	22	3.5	87	13	0
UR Tanzania	64	1.8	29	5.7	94	33
Viet Nam	70	7.9	19	29	71	42
Zambia	38	11	20	35	65	7.5
Zimbabwe	30	<0.1	13	0.3	100	17
30 high TB burden countries	4 795	3 052	670	82	18	1 073

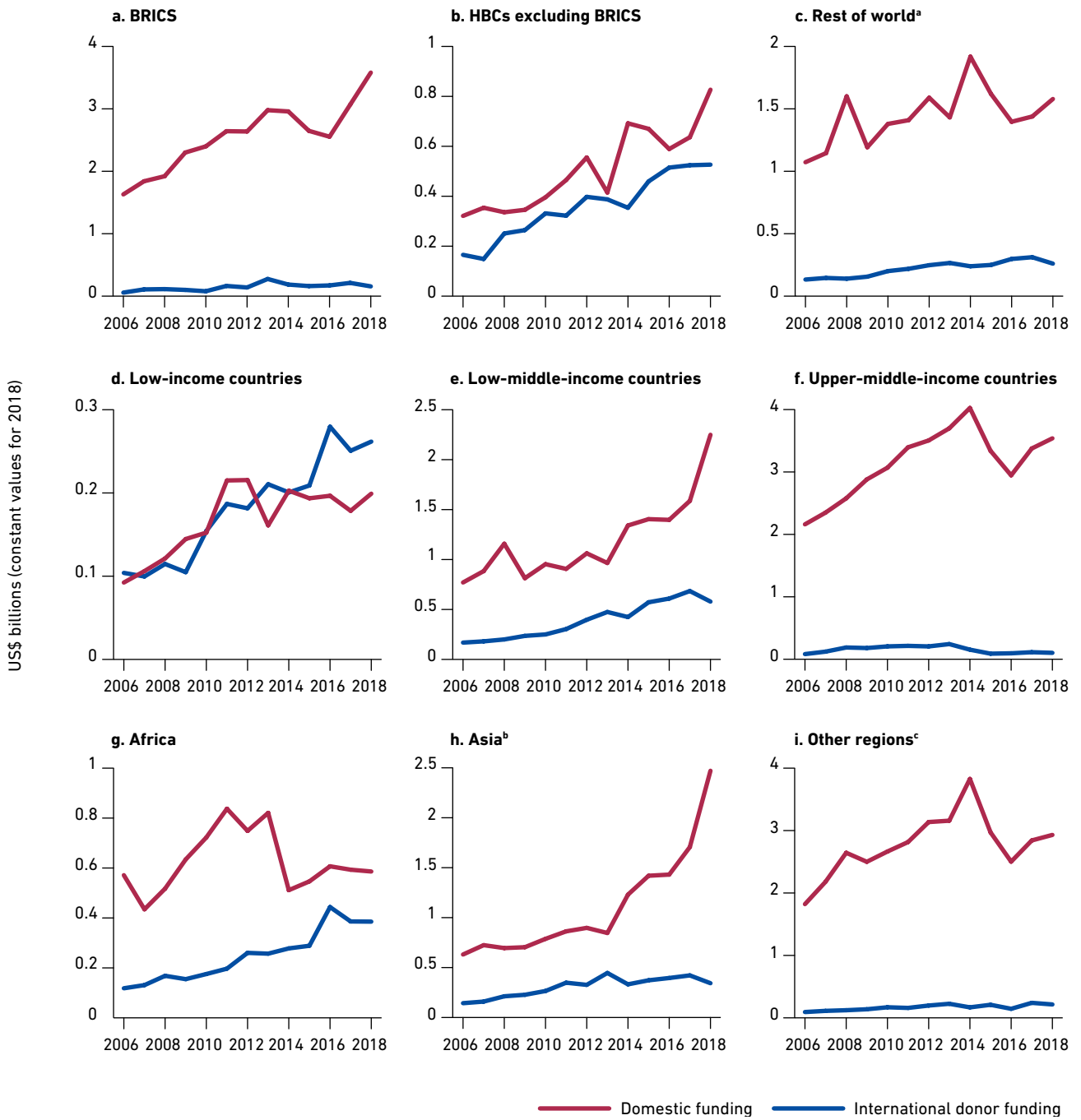
Subtotals do not always add to the total due to rounding.

^a In 2018, the budget reported by Thailand was for the central level only.

^b The funding gap reflects the anticipated gap for the year at the time a country reported data to WHO in the 2018 round of global TB data collection.

FIG. 6.6

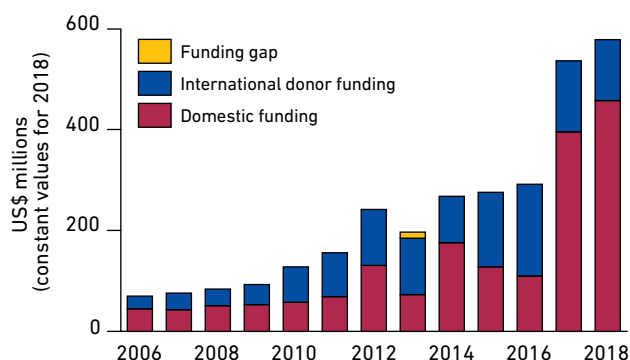
Funding for TB prevention, diagnosis and treatment from domestic sources and international donors, 2006–2018, 9 country groups



^a Rest of world includes 89 countries that are not in the list of 30 high TB burden countries.

^b Asia includes the WHO regions of South-East Asia and the Western Pacific.

^c Other regions consist of three WHO regions: the Eastern Mediterranean Region, the European Region, and the Region of the Americas.

FIG. 6.7**National budget for TB and sources of funding in India, 2006–2018**

(Fig. 6.9). Compared with 2017, funding gaps increased in all country groupings.

In 2018, the largest funding gaps among low-income countries were for high TB burden countries: the Democratic People's Republic of Korea (US\$ 77 million), Ethiopia (US\$ 52 million), the Democratic Republic of the Congo (US\$ 37 million) and the United Republic of Tanzania (US\$ 33 million) (Table 6.2). Funding gaps increased for upper-middle-income countries in 2018 (US\$ 154 million). This increase is mostly explained by large funding gaps reported by China (US\$ 67 million) and Guatemala (US\$ 16 million).

Half of the total reported funding gap in 2018 is accounted for by countries in the WHO African Region (US\$ 600 million), with Nigeria accounting for half of the region's gap (US\$ 312 million), followed by Ethiopia (US\$ 52 million), Angola (US\$ 37 million) and the Democratic Republic of the Congo (US\$ 37 million). Most of the remaining gap was reported by countries in the WHO South-East Asia Region, primarily Indonesia (US\$ 145 million) and the Democratic People's Republic of Korea (US\$ 77 million), and by countries in the WHO Western Pacific Region, primarily China (US\$ 68 million) and the Philippines (US\$ 66 million) (Fig. 6.9).

6.4 Unit costs of treatment for drug-susceptible TB and multidrug-resistant TB, 2017

The cost per patient treated in 2017 for drug-susceptible TB and MDR-TB was estimated for 113 countries and 85 countries, respectively.¹ All 30 countries in the lists of high TB burden countries and high MDR-TB burden countries were included in the analyses.² Unit cost estimates are shown in Fig. 6.10 and Fig. 6.11, and analytical methods are summarized in Box 6.1.

¹ Analysis for drug-susceptible TB was limited to countries that notified at least 100 TB cases in 2017. For MDR-TB, estimates were restricted to countries that reported at least 20 patients on second-line treatment for MDR-TB in 2017.

² For further details about both lists, see Chapter 2.

A recent global initiative to improve the availability and quality of unit cost data for TB is summarized in Box 6.3.

6.4.1 Drug-susceptible TB

The median cost per patient treated for drug-susceptible TB in 2017 was US\$ 1224 (Fig. 6.10).³ In general, about 68% of this cost was accounted for by reported NTP expenditures, with the remainder being costs for inpatient and outpatient care. There was a positive relationship between the cost per patient treated and the gross domestic product (GDP) per capita, and a negative relationship with the size of the patient caseload (indicating economies of scale, e.g. in China and India). In 29 of the 30 high TB burden countries included in the analysis, the cost per patient treated for drug-susceptible TB was less than the GDP per capita; the exceptions were Sierra Leone and Namibia.

The cost per patient treated was typically higher in the 17 countries included from the WHO European Region. Countries in Eastern Europe and Central Asia (EECA) have relatively high costs due to extensive use of hospitalization for patients in the intensive phase of treatment, with hospital admissions averaging 56 days per person in 2017. High programme costs relative to a smaller pool of patients also help to explain comparatively high per-patient costs in some countries (e.g. in Bosnia and Herzegovina, and Macedonia). However, it is also evident that some EECA countries have markedly reduced the use of hospitalization and have changed the model of care for patients with drug-susceptible TB. From 2014 to 2017, 15 of the 17 EECA countries reduced the number of bed days per patient treated for drug-susceptible TB (see Box 6.1 for estimation method); the exceptions were Ukraine (12% increase) and Kyrgyzstan (10% increase).

6.4.2 Multidrug-resistant TB

For MDR-TB, the median cost per patient treated was US\$ 7141 in 2017 (Fig. 6.11). As with drug-susceptible TB, the cost per patient treated was positively correlated with GDP per capita. New shortened regimens of 9–12 months cost about US\$ 1000 per person. These regimens have been recommended since 2016 by WHO for patients (other than pregnant women) with rifampicin-resistant or MDR pulmonary TB who do not have resistance to second-line drugs.⁴ The uptake of such regimens has contributed to a decrease in the unit cost of treatment for MDR-TB in 2017.

³ Median values are cited rather than means because of extreme values for a few countries.

⁴ For further details about this recommendation, see Chapter 1.

BOX 6.2

International donor funding for TB prevention, diagnosis and treatment, based on donor reports to the OECD

Not all international donor funding that is provided for TB prevention, diagnosis and treatment is channelled through NTPs. The creditor reporting system (CRS) of the OECD is the most comprehensive source of information about international donor funding. Funding data (both commitments and disbursements) are provided by 31 multilateral donor organizations, the 26 countries that are members of the OECD's Development Assistance Committee, and a further two non-committee members (Kuwait and the United Arab Emirates).

Disbursement data include both direct transfers to countries and the provision of goods and services, such as in-kind transfers or technical assistance. Data on gross disbursements^a for TB (code 12263: Tuberculosis control) received by non-OECD countries during 2006–2016 were analysed. Funding for TB that flows from one OECD member to an institution or government within the OECD, such as grants from the United States (US) National Institutes for Health flowing to the United Kingdom, is not captured in the CRS. Also, government contributions to multilateral organizations are not attributed to the government of origin, only to the multilateral organization.^b

Fig. B6.2.1 shows trends in international donor funding between 2006 and 2016, from four major sources as a total and by four major regions of the world. The total from all sources in 2016 was US\$ 871 million, a big increase from

US\$ 221 million in 2006. In 2016, 69% of international donor funding was provided by the Global Fund. The second largest contributor was the US government (21%, US\$ 179 million).^c Given that about one third of the contributions to the Global Fund are from the US government, about 44% of international donor funding for TB globally originated from the US government in 2016.

From 2006 to 2016, the Global Fund was consistently the largest provider of international donor funding (with the share averaging 66% in this period). There was a marked drop between 2013 (US\$ 710 million) and 2014 (US\$ 398 million), a period in which a new funding model was introduced. This was followed by a recovery to US\$ 483 million in 2015 and to US\$ 603 million in 2016. Disbursements from the US government steadily increased from 2006 to 2014, peaking at US\$ 253 million in 2014 before declining to US\$ 179 million in 2016.^c The regional panels show that Africa and Asia receive the vast majority of international donor funding.

Fig. B6.2.2 shows the proportion and amounts of funding from 2006 to 2016 from individual countries to non-OECD countries, including their estimated funding for TB via contributions to the Global Fund. Over this period, 48% of this funding came from the United States of America. The next largest individual country contributors were France (11%), the United Kingdom (9%), Germany (6%), Japan (6%) and the Netherlands (3%).

FIG. B6.2.1

International donor funding for TB prevention, diagnosis and treatment by source, globally and by region, 2006–2016

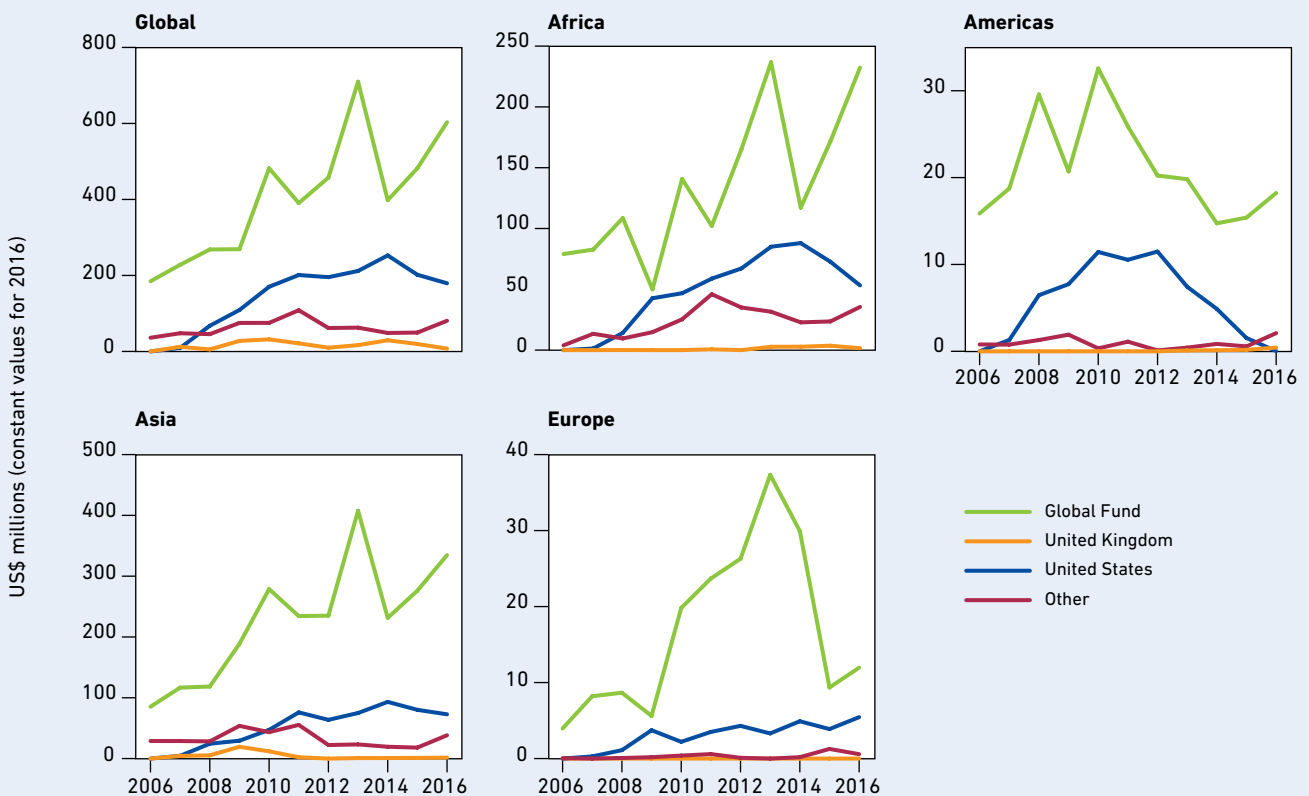
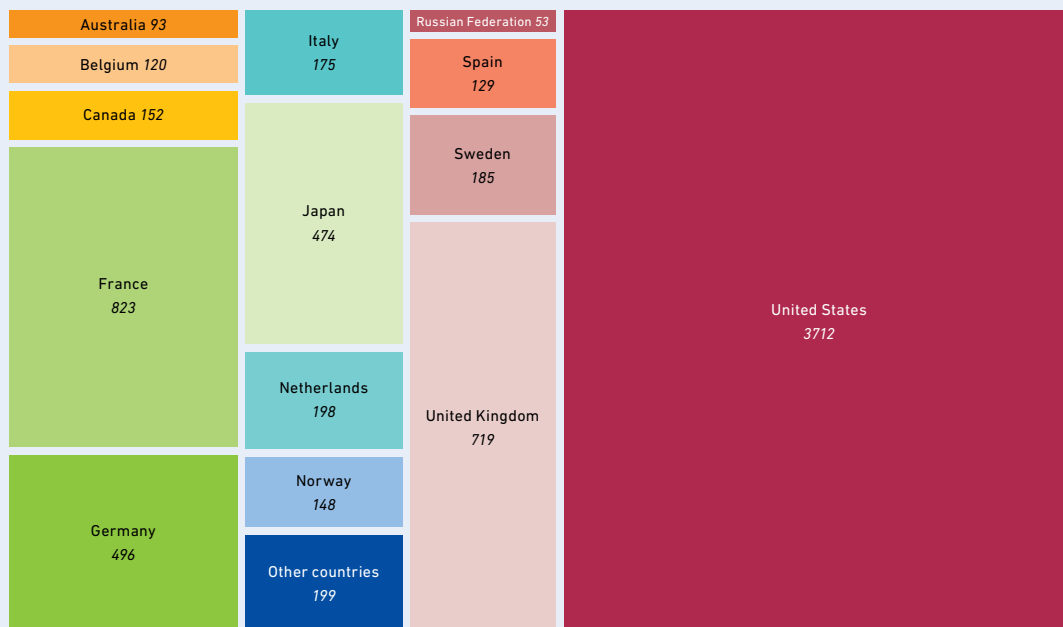


FIG. B6.2.2

International donor funding (in 2016 US\$ millions) for TB prevention, diagnosis and treatment from individual countries,^a 2006–2016



^a Funding amounts include bilateral funding as well as estimated funding for TB via contributions to the Global Fund (estimated based on the assumption that 18% of contributions are for TB). Data are shown for countries that provided at least US\$ 50 million in the time period 2006–2016. All other countries are combined in the “Other countries” box.

FIG. B6.2.3

International donor funding for TB, HIV and malaria, 2006–2016

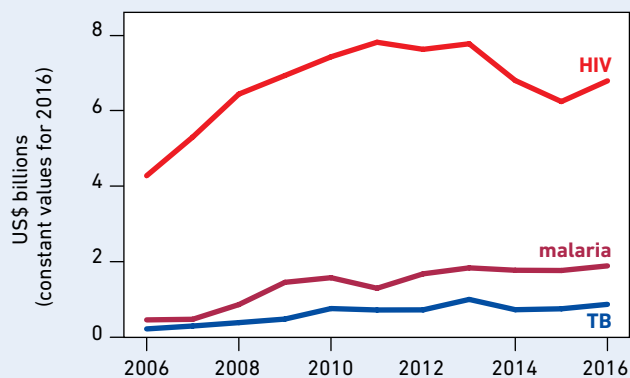
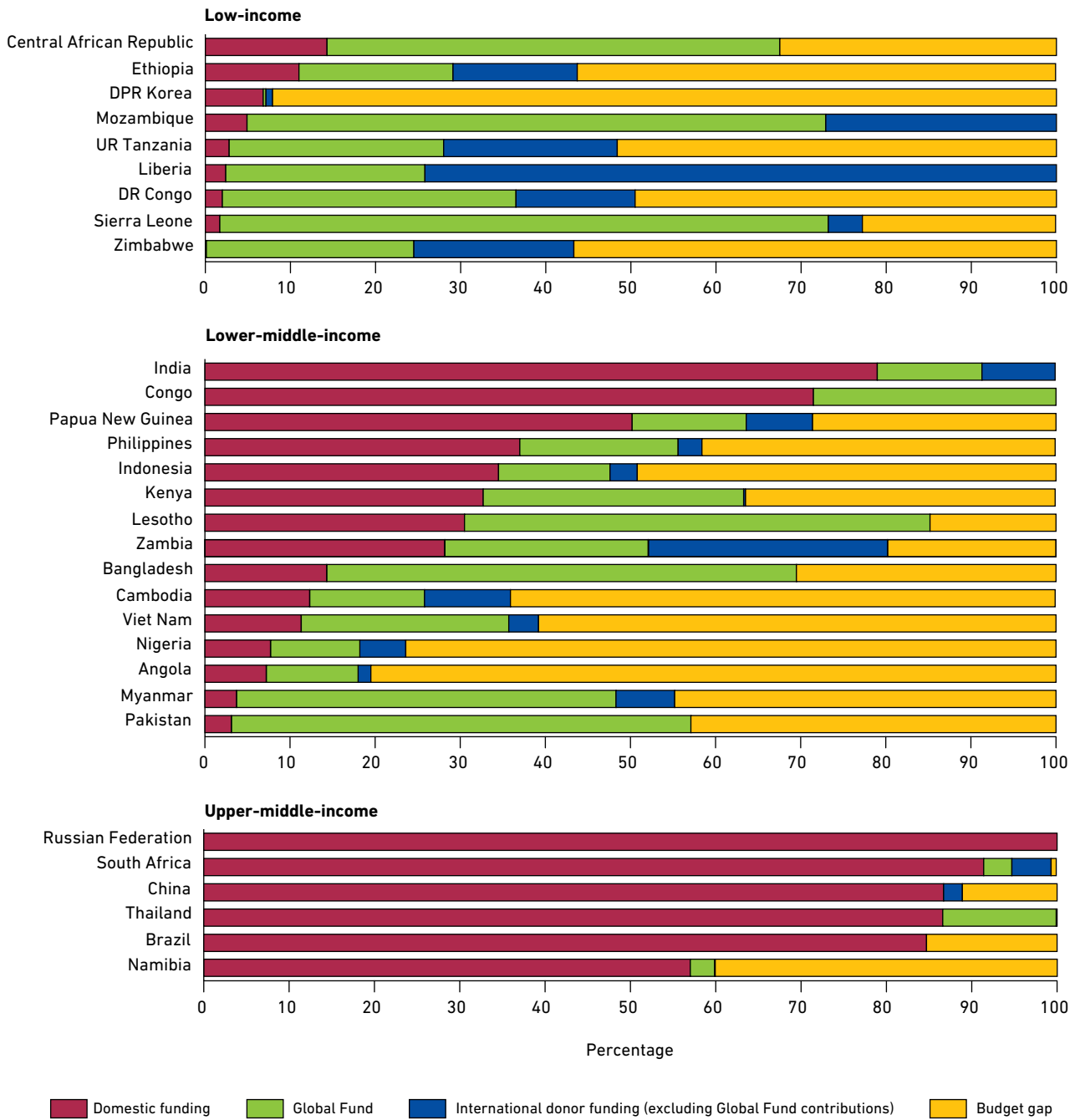


Fig. B6.2.3 shows that international funding for TB is less than half that for malaria and about one eighth that for HIV. To provide some context for these amounts, the disability-adjusted life years (DALYs) lost due to illness and death for these three diseases in 2017 were 58 million for HIV/AIDS, 56 million for malaria and 44 million for TB.^d This translates into US\$ 119 of international donor financing per DALY lost for HIV, US\$ 34 for malaria, and US\$ 20 for TB.

^a As opposed to commitments, which may not materialize.
^b An important example is funding from the Global Fund to non-OECD countries, which is attributed to the Global Fund and not to the governments or other entities that contribute to the Global Fund.
^c Disbursements from the US government captured in the OECD database are lower than official allocations.
^d Source: <http://ghdx.healthdata.org/gbd-results-tool>, accessed 8 July 2018.

FIG. 6.8

Sources of funding and funding gaps for the TB-specific budgets included in national strategic plans for TB in 2018, 30 high TB burden countries



BOX 6.3

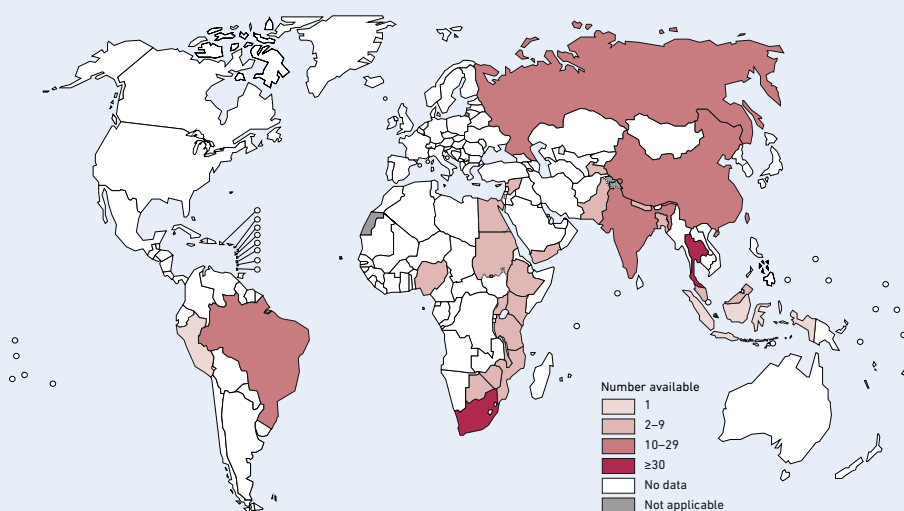
The Unit Cost Study Repository of the Global Health Cost Consortium: a new resource to estimate the cost of TB and HIV interventions

Planning and financing for TB prevention, diagnosis and treatment requires reliable data on the costs of TB services. Reporting of TB expenditures has improved substantially in recent years,^a and can be combined with data on the costs of specific TB services to inform TB budgeting and improve the efficiency of TB programmes. However, cost studies are time-consuming, and funding is required to implement them.

It has been recognized for several years that there is a global scarcity of cost data for TB services,^b and that available data were also becoming out of date. Examples of gaps in data availability include the cost of implementing new diagnostics, shortened treatment regimens for MDR-TB, innovative ways of providing TB services (e.g. public–private mix) and interventions to improve the quality of TB care. Many high TB burden countries do not have data on the unit cost of each of the TB interventions being provided.

FIG. B6.3.1

Cost per patient treated for drug-susceptible TB or MDR-TB: current availability of unit cost data from independent costing studies, 1990–2017



Source: Unit Cost Study Repository, <https://ghcosting.org/pages/data/ucsr/app/index>, accessed on 19 July 2018.

The Global Health Cost Consortium (GHCC) was established in 2016, with funding from the Bill & Melinda Gates Foundation, to help to address these gaps. The aim of the GHCC is to improve the resources available to estimate the costs of both TB and HIV interventions.^c To date, it has focused on two things: standards and methods for costing, and the development of a repository of unit cost data.

The *Reference case for estimating the global health services and interventions* provides a set of standardized principles and methods that can assist with both collecting and evaluating cost data.^d It also provides detailed costing guidelines and tools for costing TB services.

The Unit Cost Study Repository (UCSR) is a centralized source of costing data for TB and HIV interventions that is easily accessible to policy analysts, country officials and implementing organizations via an online application for desktop or mobile devices. It provides standardized unit cost data at the intervention and service output levels, with disaggregation of the unit cost available by input (e.g. personnel and capital) and for different categories of activity. An assessment of the quality of each estimate is included, based on the reference case publication.

As of July 2018, a total of 174 TB studies were included in the UCSR, based on a systematic review of costing studies implemented during the period 1990–2017 (Fig. B.6.3.1). Of these, 67 studies reported only costs incurred by patients, 60 reported provider costs, and 47 reported costs from a societal perspective (i.e. both patient and provider costs were

considered). The interventions that were costed in each study varied. As collection of new data takes place and data producers share their data with the GHCC, the UCSR will be enriched and, ultimately, this resource will improve the cost data available for use by the TB community.

UCSR data will be used by WHO's Global TB Programme to triangulate its estimates of the cost per patient treated for drug-susceptible TB and MDR-TB (Fig. 6.10 and Fig. 6.11); in particular, the unit cost estimates for inpatient stays and outpatient visits. Also, the Global TB Programme may use the UCSR data in preference to the data sources that are currently relied upon.

^a Floyd K, Fitzpatrick C, Pantoja A, Raviglione M. Domestic and donor financing for tuberculosis care and control in low-income and middle-income countries: an analysis of trends, 2002–11, and requirements to meet 2015 targets. *Lancet Glob Health*. 2013;1(2):e105–15 (<http://www.ncbi.nlm.nih.gov/pubmed/25104145>, accessed 11 July 2018).

^b Laurence YV, Griffiths UK, Vassall A. Costs to health services and the patient of treating tuberculosis: a systematic literature review. *Pharmacoeconomics*. 2015;33(9):939–55 (<http://www.ncbi.nlm.nih.gov/pubmed/25939501>, accessed 12 July 2018).

^c Global Health Cost Consortium 2016 website, available from <https://ghcosting.org/>.

^d Vassall A, Sweeney S, Kahn J, Gomez GB, Bollinger L, Marseille E et al. Reference case for estimating the costs of global health services and interventions. Global Health Cost Consortium; 2017 (https://ghcosting.org/pages/standards/reference_case, accessed 12 July 2018).

FIG. 6.9

Reported funding gaps for TB by income group and by WHO region, 2006–2018

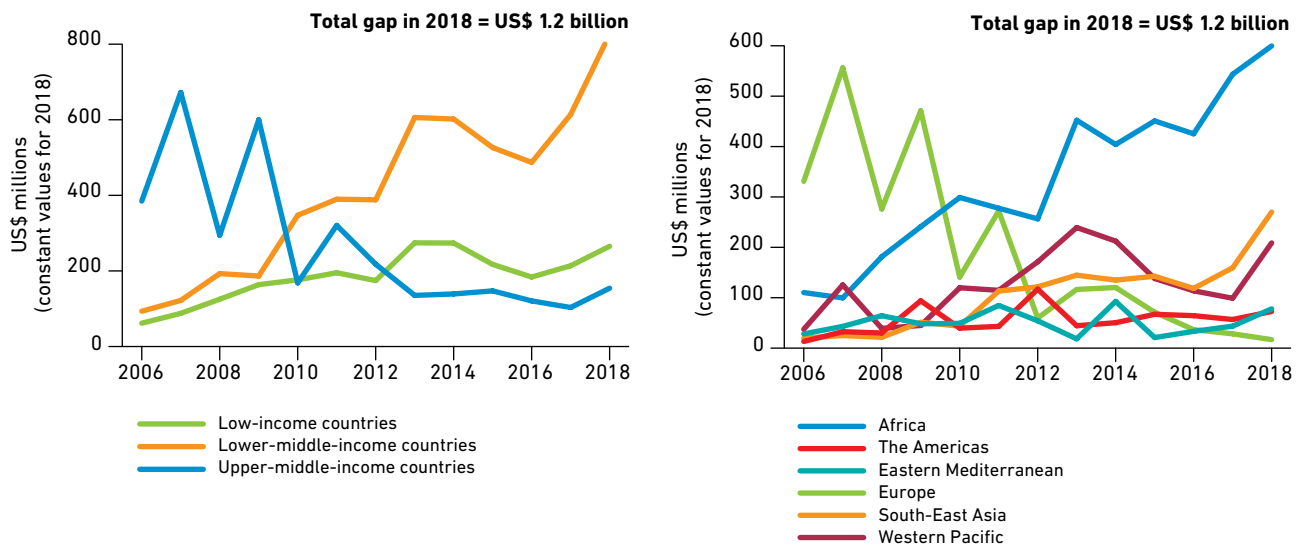
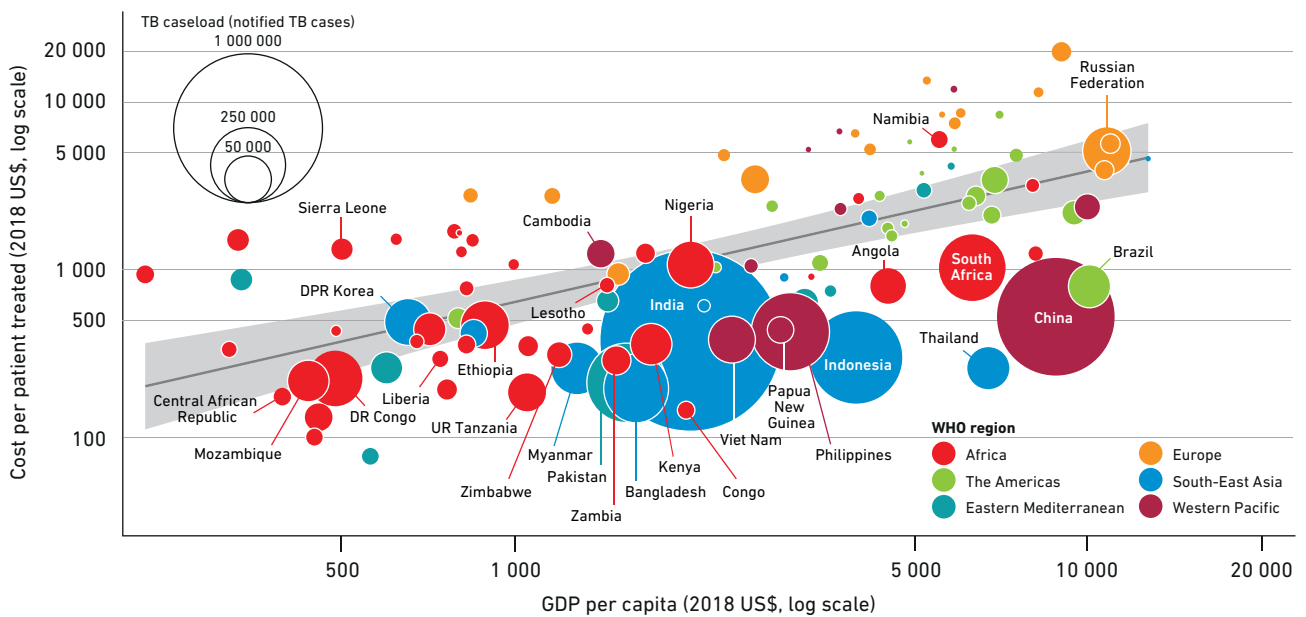


FIG. 6.10

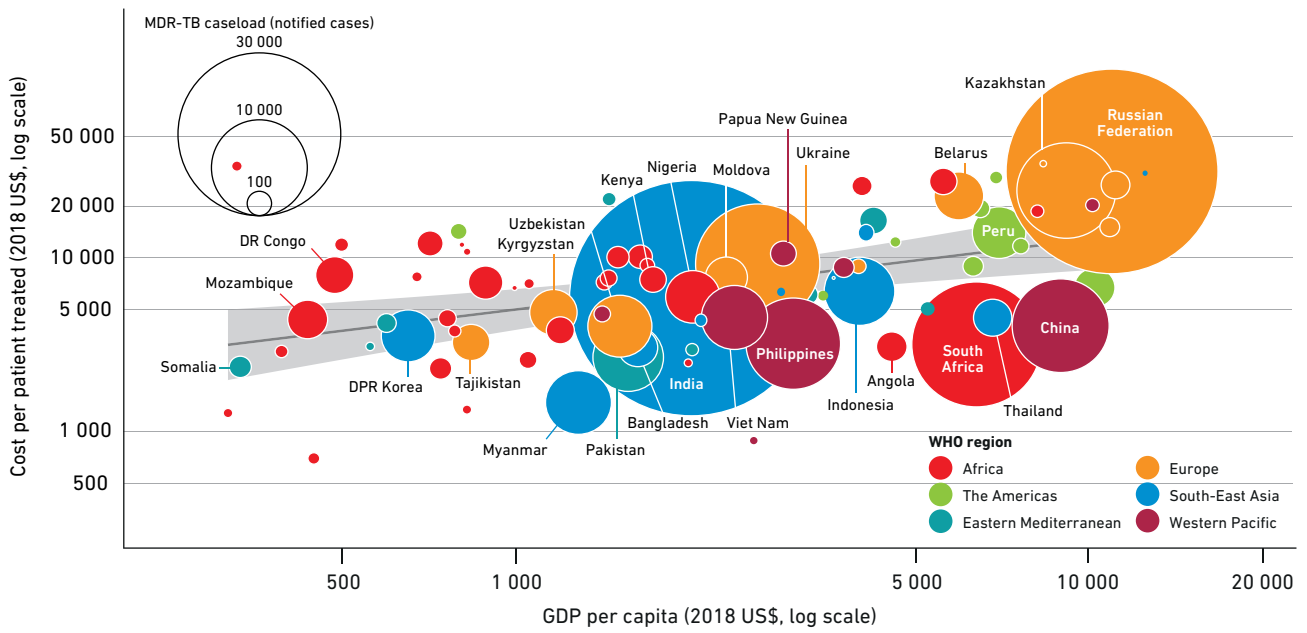
Estimated cost per patient treated for drug-susceptible TB in 113 countries, 2017^a



^a Limited to countries with at least 100 patients on first-line treatment in 2017.

FIG. 6.11

Estimated cost per patient treated for MDR-TB in 85 countries, 2017^a



^a Limited to countries with at least 20 patients on second-line treatment in 2017.



The Paraisópolis favela in São Paulo, Brazil, and the neighbouring district of Morumbi
Tuca Vieira

Chapter 7. Universal health coverage, social protection and social determinants

KEY FACTS AND MESSAGES

Achieving the tuberculosis (TB) targets and milestones of the End TB Strategy and the TB target set in the Sustainable Development Goals (SDGs) requires provision of TB care and prevention within the broader context of universal health coverage (UHC), and multisectoral action to address the social and economic determinants and consequences of TB.

UHC means that everyone – irrespective of their living standards or their particular disease or health needs – receives the health services they need, and that using health services does not cause financial hardship. SDG target 3.8 is to achieve UHC by 2030.

The SDGs include two indicators for monitoring progress towards UHC. The first is the coverage of essential health services, measured as an index (with values from 0 to 100). The second is the proportion of households in the general population that face catastrophic health expenditures, defined as household expenditures on health that exceed a certain percentage of total household expenditure or income; the denominator includes many households that did not use health services or had only very minor contact with the health system.

The latest estimates published by WHO and the World Bank suggest that at least half of the world's population lacks access to essential health services and that annually around 800 million people – almost 10% of the world's population – experience catastrophic expenditures on health. All high TB burden countries still face significant challenges to ensure both aspects of UHC.

In 2017, WHO published estimates of the resources needed to make progress towards UHC and reach other SDG-related health targets by 2030. These suggested that most middle-income countries could mobilize the required resources domestically, but that this is unlikely in low-income countries.

Social protection for people with TB and their households is part of Pillar 2 of the End TB Strategy, and social protection more broadly is an indicator under SDG 1. TB-specific social support is in place in some countries: examples include cash transfers, food packages and assistance with costs related to transport. There are also examples of TB patients being linked to more general social protection schemes.

Since 2016, considerable progress has been made in measuring costs faced by TB patients and their households. In national facility-based surveys completed to date, best estimates of the percentage of TB patients and their households facing direct and indirect costs that exceeded 20% of their annual household income ranged from 27% to 71%. Survey results should be used to help develop approaches to financing, service delivery and social protection that will reduce these costs. Examples of countries where this has been done in 2018 are Ghana and Kenya.

Many new cases of TB are attributable to undernourishment, HIV infection, smoking, diabetes and alcohol use. A recent modelling study also shows how poverty is an important underlying driver of national TB epidemics, and that eliminating extreme poverty and providing social protection (both targets under SDG 1) could substantially reduce TB incidence.

The End TB Strategy and the Sustainable Development Goals (SDGs) include a common aim: to end the global tuberculosis (TB) epidemic. Specific targets set in the End TB Strategy include a 90% reduction in TB deaths and an 80% reduction in the TB incidence rate (new cases per 100 000 population per year) by 2030 compared with 2015; more immediate milestones for 2020 are reductions of 35% and 20%, respectively. As highlighted elsewhere in this report, in particular in **Chapter 2**, achieving these targets requires provision of TB care and prevention within the broader context of universal health coverage (UHC), multisectoral action to address the social and economic determinants and consequences of TB, and a technological breakthrough by 2025 so that incidence can fall much faster than it has done historically.

This chapter has four major sections. **Section 7.1** provides an overview of the status of global progress towards UHC, and a summary of recent WHO estimates of the resources required for progress towards UHC and other SDG health targets during the period 2016–2030. **Section 7.2** discusses the role of social protection in improving care and support to TB patients and their households, and what is known about the current status of such protection in high TB burden countries. **Section 7.3** presents the status of progress in implementing national surveys of costs faced by TB patients and their households as of July 2018, and survey results from eight countries. It highlights the implications of such costs, including for approaches to TB service delivery, financing and social protection. **Section 7.4** covers the broader determinants of TB, including analysis of 14 indicators (under seven SDGs) that are associated with TB incidence.

7.1 Global progress towards UHC

UHC means that everyone – irrespective of their living standards – receives the health services they need, and that using health services does not cause financial hardship.¹

The SDG targets are for 2030, and SDG Target 3.8 is defined as “Achieve UHC, including financial risk protection, access to quality essential health-care services and access to safe, effective, quality and affordable essential medicines and vaccines” (**Chapter 2**).

Two SDG indicators have been defined to monitor progress towards SDG Target 3.8. The first (Indicator 3.8.1) is the coverage of essential health services; this is a composite index (with values from 0 to 100) based on 16 tracer indicators (one of which is TB treatment²). The

¹ World Health Organization/World Bank. Tracking universal health coverage: 2017 global monitoring report. Geneva: WHO; 2017 (<http://apps.who.int/iris/bitstream/handle/10665/259817/9789241513555-eng.pdf>, accessed 21 June 2018). See pp xii.

² The indicator used in the WHO/World Bank report is “effective TB treatment coverage”. It was calculated as treatment coverage multiplied by the treatment success rate, to capture a “quality” dimension of care. This differs from the definition of “treatment coverage” in the list of priority End TB Strategy indicators (see **Chapter 2**).

second (Indicator 3.8.2) is the “proportion of the population with large household expenditures on health as a share of total household expenditure or income”. Large is defined using two thresholds: 10% and 25%. When these thresholds are surpassed, expenditures are classified as “catastrophic”. Since this measure is population-based, the denominator includes many people who did not use health services or had only very minor contact with the health system.

The WHO/World Bank report on tracking progress towards UHC, published in December 2017, was the first to publish data on the two SDG indicators for UHC.^{3,4} Key findings from the report are summarized here.

7.1.1 Coverage of essential health services

The value of the UHC service coverage index in 2015, by country, is shown in **Fig. 7.1**. There was a great deal of variation among countries, with values ranging from a low of 22 in Somalia to 80 or above, and a median value of 65. The highest values were in high-income countries in Asia, Europe and North America. The lowest values were predominantly in countries in the WHO African Region; other countries with values below 40 were Afghanistan and Yemen. Overall, at least half of the world’s population lacked access to essential health services in 2015.

7.1.2 Financial protection

The estimated proportion of the general population facing catastrophic health spending, using the cut-off of more than 10% of household expenditure or income, is shown in **Fig. 7.2**. Compared with the service coverage index, there is more geographical variability in this indicator, including within regions.

Countries with the highest levels of catastrophic spending on health ($\geq 15\%$ of the general population facing catastrophic spending on health) include the countries that rank first (India) and second (China) in terms of their total number of TB cases, most countries in Latin America and several countries in the WHO African Region.

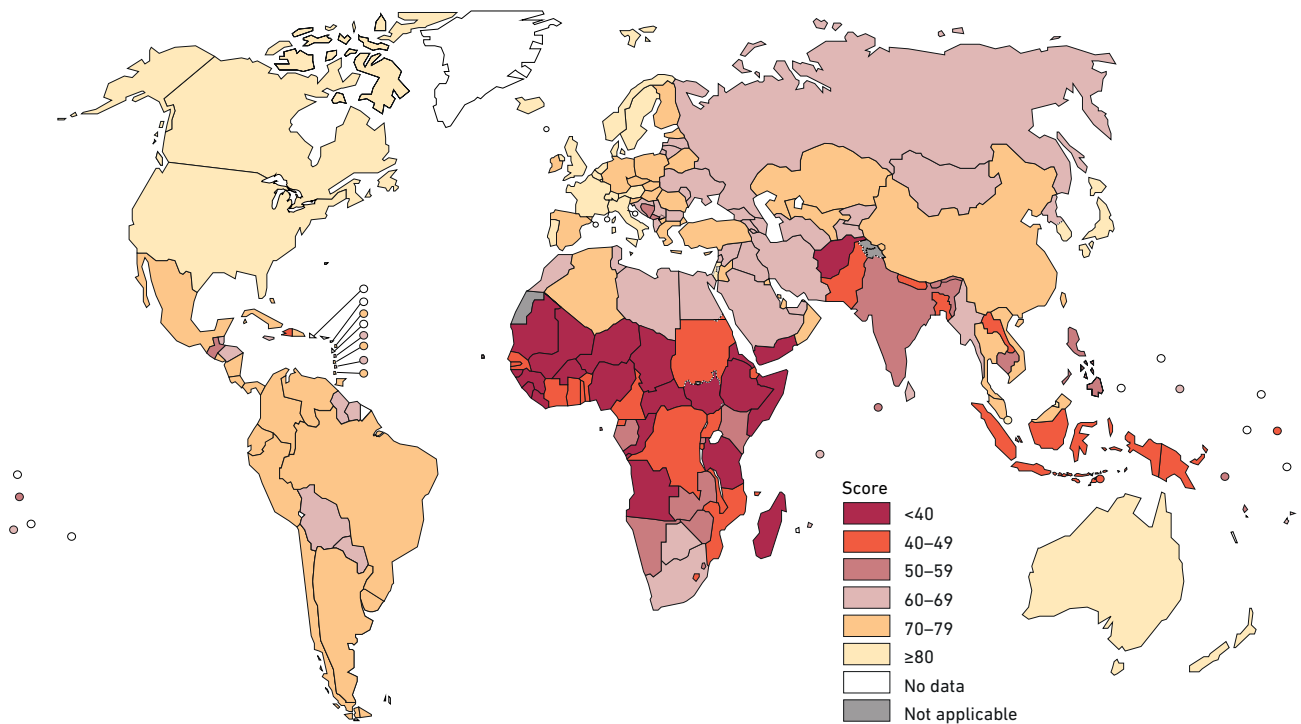
Countries with the lowest levels (0–3%) include a mix of high-income, middle-income and low-income countries. It is important to highlight that some countries may have low levels of measured spending on health because people do not access health care at all, or because capacity to spend household resources on health is very low. One example is Mozambique, for which the value of the service coverage index was 42 while the estimated proportion of households facing catastrophic expenditures on health was 1.2% (based on data for 2008).

Globally, WHO and the World Bank estimated that, in

³ World Health Organization/World Bank. Tracking universal health coverage: 2017 global monitoring report. Geneva: WHO; 2017 (<http://apps.who.int/iris/bitstream/handle/10665/259817/9789241513555-eng.pdf>, accessed 21 June 2018).

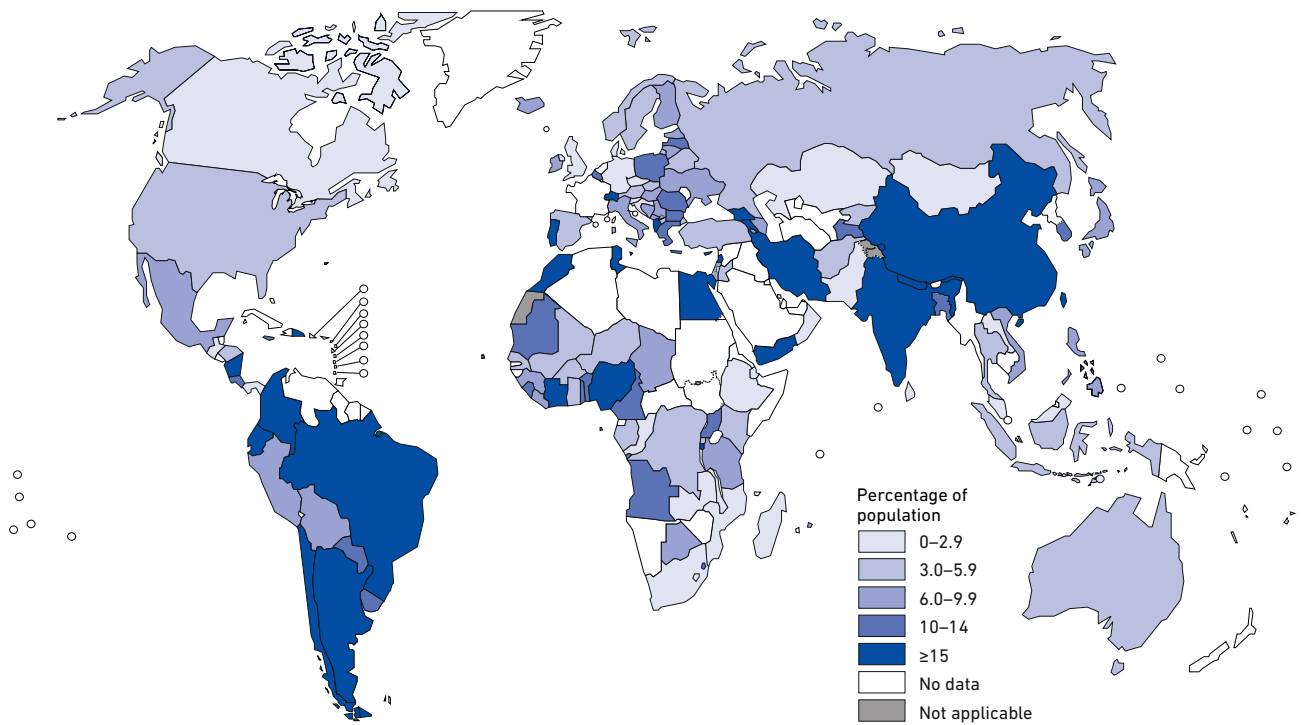
⁴ For Indicator 3.8.2, estimates in the report are based primarily on the cut-off of 10% of total household income.

FIG. 7.1
UHC service coverage index by country, 2015



Source: WHO Universal Health Coverage data portal.

FIG. 7.2
Percentage of the general population facing catastrophic health expenditures,^a latest available year of data^b



^a Defined as ≥10% of total household income or household consumption.

^b The latest available year ranges from 1993 to 2015, consistent with the WHO/World Bank report on tracking universal health coverage published in 2017. (<http://apps.who.int/iris/bitstream/handle/10665/259817/9789241513555-eng.pdf>, accessed 21 June 2018). Data can be obtained from the WHO Global Health Observatory: <http://apps.who.int/gho/portal/uhc-fp-cabinet-wrapper-v2.jsp?id=1020201> (accessed 15 August 2018).

2010, a total of 808 million people (12% of the world's population) incurred catastrophic expenditures on health.

7.1.3 Status of the two UHC indicators in the 30 high TB burden countries

The status of the two SDG indicators for UHC (3.8.1 and 3.8.2) in the 30 high TB burden countries, stratified by income group,¹ is shown in **Table 7.1**. There is a general trend for the service coverage index to improve with income, but the level of catastrophic health spending varies considerably within each income band. The countries with the highest service coverage index (above 70) are (in descending order) Brazil, China, Thailand and Viet Nam. The countries with the worst (highest) values for levels of catastrophic spending on health were middle-income countries (for example Brazil, India and Nigeria), while those with the lowest values (<2%) included both low-income countries and several middle-income countries. As explained above, some countries may have low values for the percentage of the population facing catastrophic expenditures on health because people do not access health care at all, or because capacity to spend household resources on health is very low.

7.1.4 UHC financing prospects, 2016–2030

In 2017, WHO published estimates of the resources needed during the period 2016–2030 to make progress towards UHC and to reach other SDG-related health targets, and compared these with projected total health expenditures in the same time period. Referred to in shorthand as the *WHO SDG Health Price Tag*,² the estimates are for 67 low- and middle-income countries that account for 75% of the world's population, and they focus on the additional (or incremental) resources needed compared with levels in 2014. Two scenarios were considered for resource needs (termed “ambitious” and “progress”), and two scenarios (referred to as “moderate” and “optimistic”) were also considered for total health expenditures.

Key findings included the following:

- In the “ambitious” scenario for resource needs (based on achievement of the 2030 SDG targets), the additional investment (compared with 2014) required per year grew from US\$ 134 billion in 2016 to US\$ 371 billion (equivalent to an extra US\$ 58 per person) in 2030.
- Most of the increased investment required (75% of the total) was for expanding and strengthening the health workforce and health services infrastructure

(including buildings and medical equipment) to reach recommended benchmarks. The remainder was for specific priorities, including TB. The largest share of investments needed for specific diseases or programmes was accounted for by noncommunicable diseases.

- Overall, health expenditure (in both the “moderate” and “optimistic” scenarios) was projected to be sufficient to cover “ambitious” scenario investment needs in middle-income countries. However, given uneven capacity to mobilize additional resources, some countries were expected to face gaps, especially in the first few years. In the period 2026–2030, it was predicted that about five out of the 39 middle-income countries included in the analysis would face funding gaps.
- Overall, projected health expenditures were not sufficient to cover investment needs in low-income countries, even in the “optimistic” scenario for health expenditures and the “progress” scenario for resource needs.
- Improved revenue generation and management of public expenditures, and increased public health budgets were needed, in both low- and middle-income countries.

Further analyses based on the *WHO SDG Health Price Tag*, focusing on comparisons of total (as opposed to incremental) investment needs with projections of total health expenditures, are shown in **Fig. 7.3** and **Fig. 7.4**. These illustrate the same key messages in terms of the extent to which the resources needed to achieve UHC and other SDG-related health targets can be mobilized in low- and middle-income countries. In addition, they indicate that, by 2030, total average health spending would need to increase to US\$ 112 per capita in low-income countries, US\$ 146 per capita in lower-middle-income countries and US\$ 536 per capita in upper-middle-income countries (**Fig. 7.4**). Total funding needs as a percentage of gross domestic product (GDP) would rise from 5.6% in 2014 to 7.5% by 2030 across all 67 countries (data not shown), with a much greater increase needed in low-income countries (from an average of 6% in 2014 to around 12% by 2024).

7.2 Social protection in the context of the TB response

Social protection refers to an integrated set of policies and programmes that provide support to those in need. Major elements include:

- social assistance programmes;
- social insurance schemes; and
- labour market interventions, for example to facilitate access to employment and development of skills.

Social protection floors are a basic minimum set of policies and programmes that provide a safety net in the

¹ Income groups are based on the classification of 2018 for consistency with **Chapter 6** of this report. However, estimates of the incidence of catastrophic health spending in the WHO/World Bank report, on monitoring UHC, are for earlier years.

² Stenberg K, Hanssen O, Tan-Torres Edejer T et al. Financing transformative health systems towards achievement of the health Sustainable Development Goals: a model for projected resource needs in 67 low-income and middle-income countries. *Lancet Glob Health*. 2017;5(9):e875–e877 (<https://www.ncbi.nlm.nih.gov/pubmed/28728918>, accessed 11 July 2018).

TABLE 7.1
UHC service coverage indicator (SDG 3.8.1) and percentage of general population facing catastrophic health expenditures (SDG 3.8.2),^a 30 high TB burden countries, stratified by income group^b

COUNTRY	SERVICE COVERAGE INDEX	CATASTROPHIC HEALTH EXPENDITURE ^c
LOW INCOME		
Ethiopia	39	0.82
Mozambique	42	1.2
DR Congo	40	5.8
Liberia	34	7.9
UR Tanzania	39	9.9
Sierra Leone	36	10
Zimbabwe	55	—
Central African Republic	33	—
DPR Korea	68	—
LOWER-MIDDLE INCOME		
Zambia	56	0.29
Pakistan	40	1.0
Lesotho	45	1.8
Congo	38	2.0
Indonesia	49	3.6
Kenya	57	5.8
Philippines	58	6.3
Viet Nam	73	9.8
Angola	36	12
Bangladesh	46	14
India	56	17
Nigeria	39	25
Myanmar	60	—
Cambodia	55	—
Papua New Guinea	41	—
UPPER-MIDDLE INCOME		
South Africa	67	1.4
Thailand	75	3.4
Russian Federation	63	4.9
China	76	18
Brazil	77	26
Namibia	59	—

— Data were not available.

^a Data of UHC service coverage are from 2015, and data of catastrophic health expenditures cover the period 2002–2015.

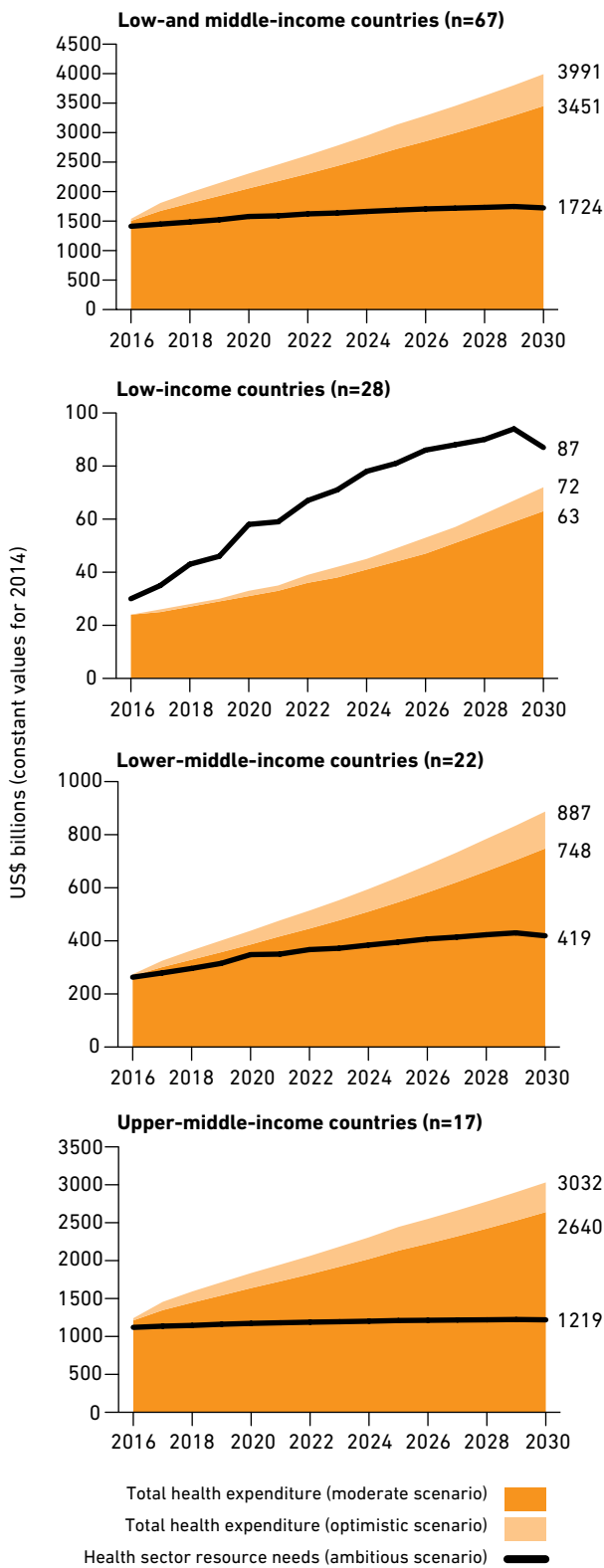
^b Countries are listed within each income group (as per 2018 World Bank classification) according to the level of catastrophic health expenditure (from lowest to highest).

^c Defined as ≥10% of total household income or household consumption.

The latest available year for data on “catastrophic health expenditures” ranges from 2008 to 2015 for 16 of the 30 countries; this time period is consistent with the years of data used for joint analysis of service coverage and catastrophic expenditures in the WHO/World Bank report on UHC monitoring (<http://apps.who.int/iris/bitstream/handle/10665/259817/9789241513555-eng.pdf>, accessed 21 June 2018). For 7 countries, however, the values shown in the table are for earlier years: China (2007), DR Congo (2004), Ethiopia (2004), Kenya (2005), Lesotho (2002), Liberia (2007) and Sierra Leone (2003). No data are available for Cambodia, Central African Republic, DPR Korea, Myanmar, Namibia, Papua New Guinea and Zimbabwe. Data can be obtained from the WHO Global Health Observatory: <http://apps.who.int/gho/portal/uhc-fp-cabinet-wrapper-v2.jsp?id=1020201> (accessed 15 August 2018).

FIG. 7.3

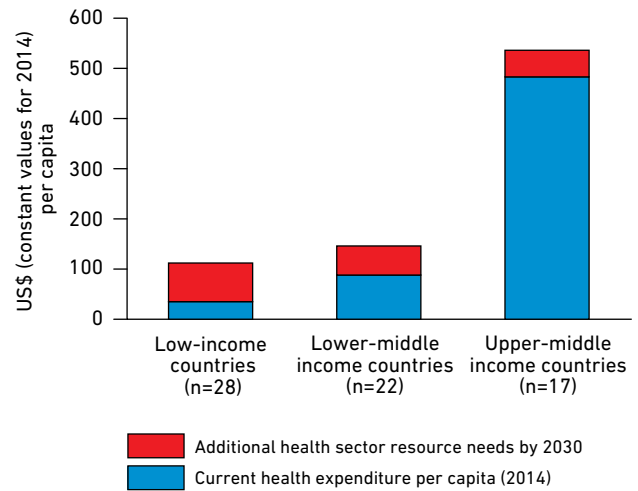
Funding needs to progressively expand services towards UHC and reach other SDG health targets in 67 low- and middle-income countries compared with projected total health expenditures, 2016–2030



Source: Data from Stenberg K, Hanssen O, Tan-Torres Edejer T et al. Financing transformative health systems towards achievement of the health Sustainable Development Goals: a model for projected resource needs in 67 low-income and middle-income countries. *Lancet Global Health* 2017; 5: e875–87.

FIG. 7.4

Funding needs (per capita) to progressively expand services towards UHC and reach other SDG health targets by 2030, by country income group^a



^a Income groups are defined as of July 2016. Per person health costs are reported as population-weighted mean values per income group per year.

Source: Data from Stenberg K, Hanssen O, Tan-Torres Edejer T et al. Financing transformative health systems towards achievement of the health Sustainable Development Goals: a model for projected resource needs in 67 low-income and middle-income countries. *Lancet Global Health* 2017; 5: e875–87.

event of unforeseen shocks. Although this is protective, the aim of comprehensive social protection is for poverty reduction, and sustainable and inclusive economic growth.

Social protection is an indicator under SDG 1, which is “End poverty in all its forms everywhere”. SDG Target 1.3 is to “Implement nationally appropriate social protection systems and measures for all, including floors, and by 2030 achieve substantial coverage of the poor and the vulnerable”.¹

Poverty and TB are intricately linked. People with TB can face severe direct and indirect financial and economic costs that affect their ability to access care and complete treatment, while poverty is a well-recognized risk factor for TB. As such, social protection for people with TB and their households is part of Pillar 2 of the End TB Strategy (Chapter 2).

The 2016 and 2017 editions of the *Global tuberculosis report*,^{2,3} and the 2015 WHO guide to implementing the

¹ The inclusion of a target for social protection floors in the SDGs follows recommendations on the topic by the International Labour Organization (ILO) in 2012. For further details, see International Labour Organization. R202 – Social protection floors recommendation, 2012 (no. 202). Geneva: ILO (101st ILC session); 2012 (https://www.ilo.org/dyn/normlex/en/f?p=NORMLEXPUB:12100:0::NO::P12100_INSTRUMENT_ID:3065524, accessed 11 July 2018).

² World Health Organization. *Global tuberculosis report 2016* (WHO/HTM/TB/2016.13). Geneva: WHO; 2016 (<http://apps.who.int/iris/bitstream/10665/250441/1/9789241565394-eng.pdf>, accessed 11 July 2018).

³ World Health Organization. *Global tuberculosis report 2017* (WHO/HTM/TB/2017.23). Geneva: WHO; 2017 (<http://apps.who.int/iris/bitstream/handle/10665/259366/9789241565516-eng.pdf>, accessed 11 July 2018).

TABLE 7.2
Social protection policy and cash transfer schemes in 30 high TB burden countries

<input checked="" type="checkbox"/> YES <input type="checkbox"/> NOT IDENTIFIED	STRATEGY/POLICY ^a	REGISTRY ^b	TARGETING ^c	
			POOR AND VULNERABLE	DISABLED
Angola	<input checked="" type="checkbox"/>	<input type="checkbox"/>	<input type="checkbox"/>	<input type="checkbox"/>
Bangladesh	<input checked="" type="checkbox"/>	<input checked="" type="checkbox"/>	<input checked="" type="checkbox"/>	<input checked="" type="checkbox"/>
Brazil	<input checked="" type="checkbox"/>	<input checked="" type="checkbox"/>	<input checked="" type="checkbox"/>	<input checked="" type="checkbox"/>
Cambodia	<input checked="" type="checkbox"/>	<input checked="" type="checkbox"/>	<input checked="" type="checkbox"/>	<input type="checkbox"/>
Central African Republic	<input type="checkbox"/>	<input type="checkbox"/>	<input type="checkbox"/>	<input type="checkbox"/>
China	<input checked="" type="checkbox"/>	<input checked="" type="checkbox"/>	<input checked="" type="checkbox"/>	<input checked="" type="checkbox"/>
DR Congo	<input type="checkbox"/>	<input type="checkbox"/>	<input type="checkbox"/>	<input type="checkbox"/>
Congo	<input checked="" type="checkbox"/>	<input checked="" type="checkbox"/>	<input checked="" type="checkbox"/>	<input type="checkbox"/>
DPR Korea	<input type="checkbox"/>	<input type="checkbox"/>	<input type="checkbox"/>	<input type="checkbox"/>
Ethiopia	<input checked="" type="checkbox"/>	<input checked="" type="checkbox"/>	<input checked="" type="checkbox"/>	<input type="checkbox"/>
India	<input checked="" type="checkbox"/>	<input checked="" type="checkbox"/>	<input checked="" type="checkbox"/>	<input checked="" type="checkbox"/>
Indonesia	<input checked="" type="checkbox"/>	<input checked="" type="checkbox"/>	<input checked="" type="checkbox"/>	<input checked="" type="checkbox"/>
Kenya	<input checked="" type="checkbox"/>	<input checked="" type="checkbox"/>	<input checked="" type="checkbox"/>	<input checked="" type="checkbox"/>
Lesotho	<input checked="" type="checkbox"/>	<input checked="" type="checkbox"/>	<input checked="" type="checkbox"/>	<input type="checkbox"/>
Liberia	<input checked="" type="checkbox"/>	<input checked="" type="checkbox"/>	<input checked="" type="checkbox"/>	<input type="checkbox"/>
Mozambique	<input checked="" type="checkbox"/>	<input checked="" type="checkbox"/>	<input checked="" type="checkbox"/>	<input checked="" type="checkbox"/>
Myanmar	<input checked="" type="checkbox"/>	<input type="checkbox"/>	<input checked="" type="checkbox"/>	<input type="checkbox"/>
Namibia	<input checked="" type="checkbox"/>	<input type="checkbox"/>	<input checked="" type="checkbox"/>	<input checked="" type="checkbox"/>
Nigeria	<input checked="" type="checkbox"/>	<input checked="" type="checkbox"/>	<input checked="" type="checkbox"/>	<input type="checkbox"/>
Pakistan	<input checked="" type="checkbox"/>	<input checked="" type="checkbox"/>	<input checked="" type="checkbox"/>	<input type="checkbox"/>
Papua New Guinea	<input checked="" type="checkbox"/>	<input checked="" type="checkbox"/>	<input type="checkbox"/>	<input checked="" type="checkbox"/>
Philippines	<input checked="" type="checkbox"/>	<input checked="" type="checkbox"/>	<input checked="" type="checkbox"/>	<input type="checkbox"/>
Russian Federation	<input checked="" type="checkbox"/>	<input type="checkbox"/>	<input checked="" type="checkbox"/>	<input checked="" type="checkbox"/>
Sierra Leone	<input checked="" type="checkbox"/>	<input checked="" type="checkbox"/>	<input checked="" type="checkbox"/>	<input type="checkbox"/>
South Africa	<input checked="" type="checkbox"/>	<input checked="" type="checkbox"/>	<input checked="" type="checkbox"/>	<input checked="" type="checkbox"/>
Thailand	<input checked="" type="checkbox"/>	<input checked="" type="checkbox"/>	<input checked="" type="checkbox"/>	<input checked="" type="checkbox"/>
UR Tanzania	<input type="checkbox"/>	<input type="checkbox"/>	<input checked="" type="checkbox"/>	<input type="checkbox"/>
Viet Nam	<input checked="" type="checkbox"/>	<input checked="" type="checkbox"/>	<input checked="" type="checkbox"/>	<input checked="" type="checkbox"/>
Zambia	<input checked="" type="checkbox"/>	<input checked="" type="checkbox"/>	<input checked="" type="checkbox"/>	<input checked="" type="checkbox"/>
Zimbabwe	<input checked="" type="checkbox"/>	<input checked="" type="checkbox"/>	<input checked="" type="checkbox"/>	<input checked="" type="checkbox"/>

^a Strategy/policy indicates whether a national strategy, and/or policy/legislative framework for social protection was identified.

^b Registry indicates whether a social and/or beneficiary registry system for social assistance programmes is in place.

^c Targeting indicates whether there are any identified non-contributory cash transfer schemes specifically for poor and vulnerable groups and the disabled. Poor and vulnerable groups include transfers to individuals and households based on socio-economic criteria and may include children, the elderly, poor people with disabilities, and those facing food insecurity; disabled refers to persons having a disability regardless of socio-economic status.

Sources: WHO Global TB Programme Country Social Protection Briefs; additional information on registries: The state of social safety nets 2015 (<http://documents.worldbank.org/curated/en/415491467994645020/The-state-of-social-safety-nets-2015>); Social registries for social assistance and beyond: a guidance note & assessment tool (<http://documents.worldbank.org/curated/en/698441502095248081/pdf/117971-REVISED-PUBLIC-Discussion-paper-1704.pdf>).

End TB Strategy,¹ explained core principles related to social protection in the context of TB. They also provided country examples of steps being taken to establish and/or strengthen social protection systems in general, including social assistance schemes of particular relevance to TB-affected households and communities.

¹ World Health Organization. Implementing the End TB Strategy: the essentials (WHO/HTM/TB/2015.31). Geneva: WHO; 2015 (http://www.who.int/tb/publications/2015/end_tb_essential.pdf?ua=1, accessed 11 July 2018).

The need for expanded policy and legal frameworks, financing and programme strengthening to ensure that support is available on a systematic and sustainable basis was highlighted. In addition, examples were provided of social support in the context of TB treatment services, such as transport, vouchers, food packages, and psychosocial support, and cash-transfer schemes (including those in which TB patients are linked to general schemes). Documentation on social protection for TB patients is increasing.

A summary of the elements of social protection of particular relevance to TB patients and their households that are in place in the 30 high TB burden countries is provided in **Table 7.2**, using information compiled by WHO's Global TB Programme in 2017 and 2018. The focus is on three major elements of social protection schemes that may be of substantial relevance to efforts to improve social protection for TB patients and households. These are: the existence of overall policy, strategy or legislative frameworks; a register of target populations or beneficiaries; and some level of coverage of cash transfers for specific poor or vulnerable populations or disabled persons. Of the 30 countries, 26 have a policy document, strategy or relevant legislation for social protection, and most have some form of cash transfer system.

Data on coverage levels of cash transfers are not easy to obtain, but available data suggest that coverage in many of the 30 high TB burden countries remains low and fragmented. More positively, India is an example of a country that took major steps in 2017–2018 to expand TB-specific cash transfers and linkages to broader nutrition schemes.

Stakeholder workshops to discuss the results from national surveys of costs faced by TB patients and their households (**Section 7.3**) provide an opportunity to discuss how to improve social protection for people with TB, for example through improved models of care, direct social support and better linkage to social services and other forms of social protection.

7.3 National surveys of costs faced by TB patients and their households (TB patient cost surveys)

One of the three high-level targets of the End TB Strategy is that no TB patients or their households should face costs due to TB that are catastrophic (**Chapter 2**). Monitoring of progress towards this target can also inform monitoring of progress towards UHC. The distinction between the indicator of catastrophic total costs due to TB disease and the broader indicator of catastrophic spending on health (**Section 7.1.2**) is explained in **Box 7.1**.

In 2015, WHO established a standardized protocol¹ for conducting a national survey to assess the direct and indirect costs incurred by TB patients and their households (TB patient cost surveys). Based on experience in pathfinding countries that conducted the first surveys, the protocol was refined and expanded into a handbook in 2017.²

¹ World Health Organization. Protocol for survey to determine direct and indirect costs due to TB and to estimate proportion of TB-affected households experiencing catastrophic costs due to TB: field testing version. Geneva: WHO; 2015 (http://www.who.int/tb/advisory_bodies/impact_measurement_taskforce/meetings/tf6_background_5a_patient_cost_surveys_protocol.pdf, accessed 21 June 2018).

² World Health Organization. Tuberculosis patient cost surveys: a handbook. Geneva: WHO; 2017 (http://www.who.int/tb/publications/patient_cost_surveys/en/, accessed 11 July 2018).

TB patient cost surveys have two primary objectives:

- to document the magnitude and main drivers of different types of costs incurred by TB patients (and their households), in order to guide policies to reduce financial barriers to accessing care and minimize the adverse socioeconomic impact of TB; and
- to determine the baseline and periodically measure the percentage of TB patients (and their households) treated in the NTP network who incur catastrophic total costs due to TB.

In the context of these surveys, catastrophic costs for TB patients and their households have been defined as direct medical and non-medical costs plus income losses that sum to 20% or more of household income.

WHO recommends conducting a baseline survey by 2020 at the latest, especially in high TB burden countries.

7.3.1 Global progress in implementation of surveys

The status of progress in implementing survey planning and implementation is shown in **Fig. 7.5**. By July 2018, 11 countries had completed national surveys:³ China (2016), Fiji (2017), Ghana (2016), Kenya (2017), Mongolia (2017), Myanmar (2015), Nigeria (2017), the Philippines (2016), Republic of Moldova (2016), Timor-Leste (2017) and Viet Nam (2016).⁴

In July 2018, surveys were underway in four countries: India, Sudan, Uganda and Zimbabwe. There are 13 countries in which surveys are scheduled to start in 2018 or in 2019: Brazil, Burkina Faso, Colombia, Dominican Republic, the Democratic Republic of the Congo, Ethiopia, Indonesia, Lao People's Democratic Republic, Papua New Guinea, Romania, Solomon Islands, South Africa, and the United Republic of Tanzania.

The main survey results for eight countries are shown in **Fig. 7.6**. The number in the centre of the circle shows the best estimate of the percentage of TB patients and their households that faced catastrophic costs, and the outer ring shows the distribution of costs using three major cost categories.

The percentage of TB-affected households that experienced total costs that were catastrophic ranged from 27% (95% confidence interval [CI]: 21–32%) in Kenya to 71% (95% CI: 68–73) in Nigeria.

The distribution of costs varied among countries. Nonetheless, it is evident that – despite the widespread norm of “free TB care” policies – the direct medical costs faced by TB-affected households can be high, and such costs accounted for a large proportion of total costs in some countries (e.g. in Mongolia and Myanmar). Minimizing direct medical costs borne by TB patients should be a high priority for NTPs and ministries of health.

³ Defined as having completed survey field work, analysis of data, and documentation of results (e.g. in a report).

⁴ The year indicates the year in which data collection was done.

BOX 7.1

The difference between “catastrophic total costs” for TB patients and their households, and the SDG indicator of catastrophic expenditures on health

It is important to distinguish between the indicator of “catastrophic spending on health” used within the SDG monitoring framework (SDG 3.8.2) and the indicator of “catastrophic total costs due to TB”.

The former is a population-based indicator that measures the share of the population incurring “catastrophic spending” on health, with a threshold defined as exceeding 10% and 25% of a household’s total consumption expenditure or income. Health expenditures are defined as direct expenditures on medical care. The denominator of the SDG indicator includes many people who had no contact with the health system and thus had zero expenditures on health.

The TB-specific indicator incorporates not only direct medical payments for diagnosis and treatment, but also direct non-medical payments (e.g. for transportation and lodging) and indirect costs (e.g. lost income). In addition, the TB-specific indicator is restricted to a particular population: diagnosed TB patients who are users of health services that are part of NTP networks.

As a consequence of these differences, the percentage of TB patients facing costs that are large in relation to their household expenditure or income is expected to be much higher than the percentage of the general population facing catastrophic expenditures on health. The two indicators should not be compared directly.

FIG. 7.5

National surveys of costs faced by TB patients and their households since 2016: progress and plans (as of July 2018)

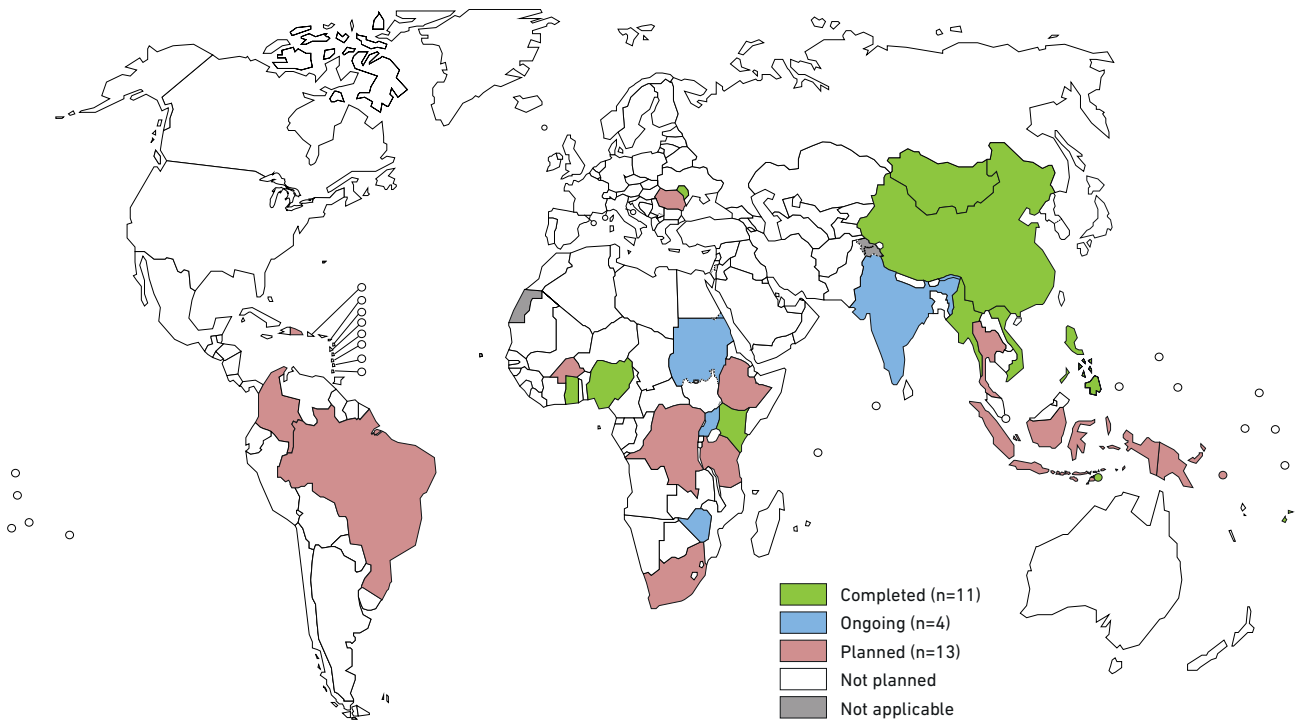
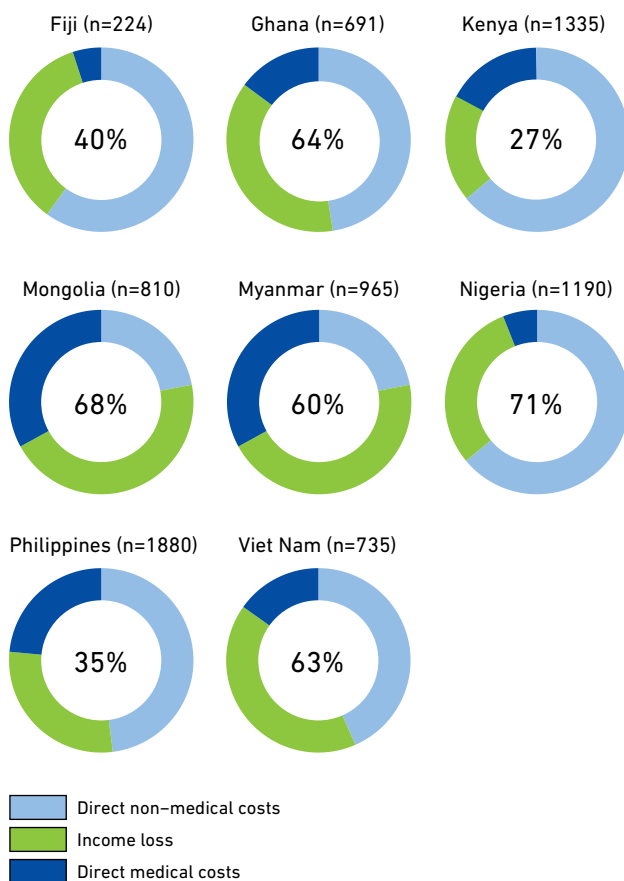


FIG. 7.6

Results from selected national surveys of costs faced by TB patients and their households, implemented 2015–2018. The number in the centre of the circle is the best estimate of the percentage of TB patients and their households that experienced total costs that were above 20% of their annual income (i.e. catastrophic total costs). The outer ring shows the distribution of costs in three major categories. The number of TB patients included in each survey is shown after the country name.



The surveys also show that actions to eliminate the indirect costs of care and to reduce income losses are needed. The combined cost of transportation, food, nutritional supplements and other non-medical expenditures (“direct non-medical costs”) accounted for the largest share of total costs in Fiji, Ghana, Nigeria, the Philippines and Viet Nam. Income losses associated with loss of employment or time lost while seeking or staying in care accounted for the largest single share of total costs in Mongolia, Myanmar and Timor-Leste.

All cost categories are influenced by the model of care, such as the extent to which hospitalization or outpatient care are relied upon, and the frequency with which attendance at health facilities is requested.

7.3.2 Policy and strategy implications of survey results

Results from TB patient cost surveys can inform policy and strategy in two major ways. First, costs can be mitigated by improving approaches to TB service delivery and financing, such as removal of user fees and the introduction of more patient-centred models of care. Second, any costs that remain after the optimization of health-care delivery can be mitigated by improved social protection measures, in collaboration with stakeholders across the social sector. Survey results should be used to stimulate the engagement of multisectoral partners and to facilitate policy discussion on both topics.

A multi-stakeholder consultation can be an effective way to initiate discussions about survey results and the actions needed in response. An early example was a multisectoral meeting in Viet Nam in March 2017, which was used to disseminate findings and formulate a joint action plan with the country’s Ministry of Labour and Social Affairs. Similar dissemination and stakeholder consultations have subsequently been held in Myanmar (2017), Kenya (July 2018) and Ghana (August 2018), resulting in multisectoral action to improve social support to TB patients and their households. The two most recent examples, in Ghana and Kenya, are described in more detail in [Box 7.2](#).

7.4 Addressing broader determinants of the TB epidemic

The influence of various social and economic determinants on the TB epidemic has long been recognized. In the late 1800s, cause-of-death data from national vital registration systems show that TB was one of the leading causes of death in some European countries. With social and economic development – such as improvements in incomes, housing and nutrition – numbers of TB cases and deaths started to decline in western Europe, North America and some other parts of the world around the turn of the 20th century, albeit slowly (1–2% per year).^{1,2} From the 1940s, the discovery, development and use of effective drug treatments substantially accelerated these trends, with national case rates (new cases per 100 000 population per year) falling by up to 10% per year and mortality rates falling even faster. The fastest historic declines in western Europe occurred during the 1950s and 1960s, in the context of UHC, rapid social and economic development and the availability of effective drug treatments.

¹ Styblo K, Meijer J, Sutherland I. The transmission of tubercle bacilli: its trend in a human population. *Bull World Health Organ.* 1969;41:137–78 (<https://www.ncbi.nlm.nih.gov/pubmed/5309081>, accessed 3 July 2018).

² Grange JM, Gandy M, Farmer P, Zumla A. Historical declines in tuberculosis: nature, nurture and the biosocial model. *Int J Tuberc Lung Dis.* 2001;5(3):208–12 (<https://www.ncbi.nlm.nih.gov/pubmed/11326817>, accessed 3 July 2018).

BOX 7.2

National surveys of costs faced by TB patients and their households in Ghana and Kenya: results and policy translation

Ghana

Ghana is a lower-middle-income country in which there has been economic growth for more than two decades and reductions in the proportion of the population living in poverty. However, social and health inequalities persist. Several social protection programmes have been implemented, and a National Health Insurance Scheme (NHIS) was introduced through an Act of Parliament in 2003. In 2017, active membership of the NHIS was estimated at 40% of the national population.

In 2016, the NTP worked with the London School of Hygiene and Tropical Medicine to conduct the first national survey of costs faced by TB patients and their households, according to the guidance in the WHO handbook on TB patient cost surveys.^a Ghana was the first country in Africa to implement such a survey.

The cross-sectional survey involved 691 patients with either drug-susceptible or multidrug-resistant (MDR) TB who were receiving treatment in health facilities in the public sector, across 25 clusters. The questionnaire included questions on costs, time losses, coping measures and asset ownership. Total costs were expressed as a percentage of annual household income; if total costs exceeded 20% of household income, the household was classified as “experiencing catastrophic total costs”.

The median cost that patients incurred per TB episode was US\$ 455, equivalent to about 25% of the average annual household income reported by survey participants. The share was significantly higher for MDR-TB patients. Non-medical costs (and particularly food) and lost income accounted for the largest share of the total cost, while medical costs accounted for less than 20%. Overall, 64% (95% CI: 61–68%) of TB-affected households, and 73% (95% CI: 61–82%) of households of patients with MDR-TB experienced total costs that were catastrophic. People with MDR-TB were pushed significantly further above the threshold for catastrophic costs than those with drug-susceptible TB.

The survey also found that TB-affected households were more likely than the general population to be poor at the time of diagnosis (46% versus 24%), and that the costs associated with TB care increased the proportion of TB-affected households living below the poverty line (from 46% to 60%). More than half (52%) of all patients were unable to pay for TB treatment from their existing income sources, and had to rely on savings, borrowing or sale of assets.

Among survey participants, 45% were enrolled in the NHIS at the time of diagnosis. A further 35% enrolled after diagnosis, and the remaining 20% were uninsured throughout. Only four patients (<1%) benefited from the Livelihood Empowerment Against Poverty (LEAP) programme, which provides conditional cash transfers to extremely poor households, with the aim of alleviating short-term poverty, and supporting long-term resiliency and human capacity development.

Shortly after the survey was completed, the NTP engaged key government agencies, to inform them about the findings and the need for multisectoral action to mitigate costs faced

by TB patients and their households, including through social support. Subsequently, a stakeholders meeting was used to disseminate survey findings and launch a national action plan to reduce and compensate for costs faced by TB patients and their households. The meeting was attended by representatives from the National Health Insurance Authority; the Ministry of Gender, Children and Social Protection; the Ghana Health Service; the Ghana AIDS Commission; the Ministry of Monitoring and Evaluation; the National Development Planning Commission; representatives of TB patients and civil society; and local and international partners.

Key intervention areas identified in the action plan to reduce direct medical costs are the inclusion of TB patients as one of the groups exempt from paying the NHIS premium under the category of indigent people, and further decentralization of TB services through the establishment of closer linkages with the Community-Based Health Planning and Services strategy. Interventions to reduce indirect costs include the expansion of existing social protection interventions to include TB patients, including their enrolment in LEAP, by making TB one of the criteria for eligibility; and the design of a social support package targeting the specific needs of TB patients, such as nutritional support and transport vouchers.

Further details are available in a recent publication.^b

Kenya

In 2017, the Kenya National Tuberculosis, Leprosy and Lung Disease Programme (NTLD) conducted the country's first national survey of costs faced by TB patients and their households. The NTLD survey included 1071 patients with drug-susceptible TB and 282 patients with drug-resistant TB.

The median cost was US\$ 256 for those with drug-susceptible TB and US\$ 1434 for those with drug-resistant TB. The median value for total household annual expenditure was US\$ 2550. The overall proportion of TB-affected households that faced catastrophic costs was 27% (95% CI: 21–32%), but this proportion was three times higher for those affected by drug-resistant TB (86%; 95% CI: 79–94%). Direct non-medical costs, particularly expenses on food and nutritional supplements beyond the patient's normal diet, were the largest cost driver, followed by productivity losses.

Significant predictors for experiencing catastrophic costs were being in a low income quintile, having no education, having a small household size, and drug-resistant TB. More than half (59%) of the patients in the survey were either severely or moderately malnourished, and only 14% were covered by the National Health Insurance Fund (NHIF) during their TB treatment.

Based on survey results, a stakeholder consultation was held in July 2018. This was used to disseminate survey findings and to discuss the actions needed to achieve the goal of eliminating catastrophic costs for TB patients and their households in Kenya. Government stakeholders represented at the consultation included the Ministry of Treasury and Planning, the Ministry of Devolution and ASAL (Arid and Semi Arid Lands), the Ministry of Labour and Social Protection, and the health committee of the national parliament. Also

participating were representatives from the NHIF and civil society, and former TB patients.

Six priority actions were identified, as follows:

1. Addition of TB as an eligibility criterion for existing social protection programmes, particularly cash-transfer programmes and systematic identification of TB patients eligible for social protection.
2. Ensuring systematic assessment of nutritional status and associated counselling for all TB patients, with provision of food support according to need, including for malnourished children of TB-affected households.
3. Inclusion of TB care in the NHIF benefit package, and increased NHIF coverage among TB patients through fast-track, premium-free enrolment.
4. Development and implementation of policies and laws to eliminate discrimination and ensure job security for TB patients, in collaboration with national authorities in the labour sector.

5. Engagement of all care providers in the provision of timely and quality-assured TB care, to reduce delays in accessing diagnosis and treatment.
6. Establishment of a high-level multi-stakeholder coordinating mechanism and forum for the implementation of the End TB Strategy.

During the consultation, the representative of the health committee of the national parliament pledged high-level political support to ensure implementation of the policy changes needed to eliminate catastrophic costs for TB-affected households in Kenya.

^a World Health Organization. Tuberculosis patient cost surveys: a handbook. Geneva: WHO; 2017 (http://www.who.int/tb/publications/patient_cost_surveys/en/, accessed 11 July 2018).

^b Pedrazzoli D, Siroka A, Boccia D, Bonsu F, Nartey K, Houben R et al. How affordable is TB care? Findings from a nationwide TB patient cost survey in Ghana. *Trop Med Int Health*. 2018; (<https://www.ncbi.nlm.nih.gov/pubmed/29851223>, accessed 11 July 2018).

The links between TB and poverty, social protection, the prevalence of undernutrition, diabetes, HIV, alcohol use, smoking, indoor air pollution and income per capita have been analysed, reviewed and summarized in recent publications (further details are provided in [Chapter 2](#)).^{1,2}

As explained in [Chapter 2](#), WHO has developed a TB-SDG monitoring framework that focuses attention on 14 indicators (from seven SDGs) that are associated with TB incidence. Monitoring of these indicators can be used to identify key influences on the TB epidemic at national level and inform the multisectoral actions required to end the TB epidemic.

For SDG 3 (health), the seven indicators selected for TB-SDG monitoring are:

- coverage of essential health services;
- percentage of total health expenditures that are out-of-pocket;
- health expenditure per capita;
- HIV prevalence;
- prevalence of smoking;
- prevalence of diabetes; and
- prevalence of alcohol use disorder.

For SDGs 1, 2, 7, 8, 10 and 11, the seven indicators selected for monitoring are:

- proportion of the population living below the international poverty line;
- proportion of the population covered by social protection floors or systems;
- prevalence of undernourishment;
- proportion of the population with primary reliance on clean fuels and technology;
- GDP per capita;
- Gini index for income inequality;³ and
- proportion of the urban population living in slums.

Collection and reporting of data for the 14 indicators does not require any additional data collection and reporting efforts by NTPs. Nor does it require data collection and reporting efforts that go beyond those to which countries have already committed in the context of the SDGs. Data are available from global databases, primarily those maintained by the United Nations (UN), WHO, the World Bank and the Joint United Nations Programme on HIV/AIDS (UNAIDS) (for further details, see [Annex 1](#)).

The most recent data for five of the seven SDG 3 indicators listed above are shown for the 30 high TB burden countries in [Table 7.3](#).⁴ For all of the indicators shown, a lower level is more desirable.

The most recent data for six of the seven selected SDG indicators beyond SDG 3 in the 30 high TB burden countries are shown in [Fig. 7.7](#).⁵ In this figure, the outer edge

¹ Lönnroth K, Jaramillo E, Williams B, Dye C, Raviglione M. Tuberculosis: the role of risk factors and social determinants. In: Blas E & Kurup A, editors. *Equity, social determinants and public health programmes*. WHO. 2010 (http://apps.who.int/iris/bitstream/10665/44289/1/9789241563970_eng.pdf, accessed 21 June 2018).

² Lönnroth K, Castro KG, Chakaya JM, Chauhan LS, Floyd K, Glaziou P et al. Tuberculosis control and elimination 2010–50: cure, care, and social development. *Lancet*. 2010;375(9728):1814–29 (<https://www.ncbi.nlm.nih.gov/pubmed/20488524>, accessed 21 June 2018).

³ The index can take values between 0 and 1, with 0 representing perfect equality and 1 representing perfect inequality.

⁴ Coverage of health services is not included because this indicator is covered in [Section 7.1](#); health expenditure per capita is also not shown, but data are included in [Annex 2](#).

⁵ GDP per capita is not included in [Fig. 7.7](#) because it is the only indicator that is not measured on a scale of 0–100. However, the latest value and recent trends in this indicator are shown in the country profiles in [Annex 2](#).

TABLE 7.3

Status of selected SDG 3 indicators, 30 high TB burden countries, latest available year

COUNTRY	OUT-OF-POCKET HEALTH EXPENDITURE (% OF CURRENT HEALTH EXPENDITURE)	HIV PREVALENCE (% OF POPULATION AGED 15–49 YEARS)	SMOKING PREVALENCE (% OF POPULATION AGED ≥15 YEARS)	DIABETES PREVALENCE (% OF POPULATION AGED ≥18 YEARS)	ALCOHOL USE DISORDERS, 12 MONTH PREVALENCE (% OF POPULATION AGED ≥15 YEARS)
Angola	33	1.9	—	8.5 7.8	8.6 1.4
Bangladesh	72	0.1	44.7 1	10 9.3	1.3 0.2
Brazil	28	0.6	17.9 10.1	7.8 8.7	8.2 3.2
Cambodia	59	0.6	33.7 2	7.4 6.9	7.6 1.4
Central African Republic	40	4.0	—	8.0 7.6	5.5 0.7
China	32	—	48.4 1.9	9.9 7.6	9.3 0.2
Congo	44	3.1	52.3 1.7	7.7 7.6	3.1 0.4
DPR Korea	—	—	—	5.8 5.9	5.1 0.9
DR Congo	37	0.7	—	6.2 6.1	7.4 0.9
Ethiopia	38	1.1	8.5 0.4	5.8 5.0	3.7 0.6
India	65	0.3	20.6 1.9	9.1 8.3	4.5 0.5
Indonesia	48	0.4	76.1 2.8	7.4 8.0	1.3 0.3
Kenya	33	5.4	20.4 1.2	5.8 6.2	5.8 0.8
Lesotho	17	25	53.9 0.4	7.3 9.9	7.5 1.1
Liberia	20	1.6	18.1 1.5	7.8 7.6	7.4 1.1
Mozambique	6.8	12	29.1 5.1	6.6 6.2	4.8 0.6
Myanmar	74	0.8	35.2 6.3	6.9 7.9	2.7 0.5
Namibia	8.3	14	34.2 9.7	7.3 7.5	8.8 1.7
Nigeria	72	2.9	10.8 0.6	6.3 6.0	3.8 0.4
Pakistan	66	0.1	36.7 2.8	13 12	0.5 0.1
Papua New Guinea	5.8	0.9	48.8 23.5	15 14	7.7 1.4
Philippines	54	0.1	41 7.8	7.1 7.3	7.7 1.4
Russian Federation	36	—	58.3 23.4	7.4 8.0	31 6.2
Sierra Leone	38	1.7	41.3 8.8	7.1 6.6	4.9 0.5
South Africa	7.7	19	33.2 8.1	9.7 13	10 1.5
Thailand	12	1.1	38.8 1.9	8.3 8.8	9.1 1.0
UR Tanzania	26	4.7	26.7 3.3	6.0 6.1	9.3 1.8
Viet Nam	43	0.4	45.9 1	5.5 5.1	8.7 0.9
Zambia	28	12	24.7 3.1	6.5 6.7	7.9 1.0
Zimbabwe	26	14	30.7 1.6	6.5 7.6	9.0 1.6

Both sexes Male Female

— indicates data not available.

Sources: World Bank Sustainable Development Goals Database (<http://datatopics.worldbank.org/sdgs/>) and WHO Global Health Observatory (<http://www.who.int/gho/en/>)

FIG. 7.7

Status of selected SDG indicators beyond SDG 3, 30 high TB burden countries, latest available year^a



Not in slums: Percentage of urban population not living in slums.
Income equality: An inverse GINI index is shown where 0 is perfect inequality and 100 is perfect equality.
Social protection: Percentage of population covered by social protection and labour programmes.
Not in poverty: Percentage of population living above the international poverty line.
Nutrition: Percentage of population not undernourished.
Clean fuels: Percentage of population with access to clean fuels and technologies for cooking.

^a Red dot indicates that there are no data for the selected indicator.

Source: World Bank Sustainable Development Goals Database (<http://datatopics.worldbank.org/sdgs/>).

of the hexagon (100) is the ideal value for each indicator. Therefore, better performance corresponds to a larger shaded-in region. To achieve this representation visually, the indicators “proportion of the urban population living in slums” and “proportion of the population living below the international poverty line” are inverted in Fig. 7.7. It should be highlighted that all indicator values in Fig. 7.7 are for the general population, as opposed to people with TB. Values for TB patients specifically (such as out-of-pocket expenditure and access to social protection) may differ from these general values.

Poverty is an underlying driver of national TB epidem-

ics, influencing several of the other indicators associated with TB incidence. This is illustrated by a recent modelling study, which showed that eliminating extreme poverty and providing social protection (both targets under SDG 1) could substantially reduce TB incidence. The main results are summarized in Box 7.3.

Table 7.3 and Fig. 7.7 show that many high TB burden countries, especially those in the low-income category, still face significant challenges to achieve a range of TB-related SDG targets. Furthermore, values for poor populations and vulnerable groups most at risk of developing TB are likely to be worse than national averages.

BOX 7.3

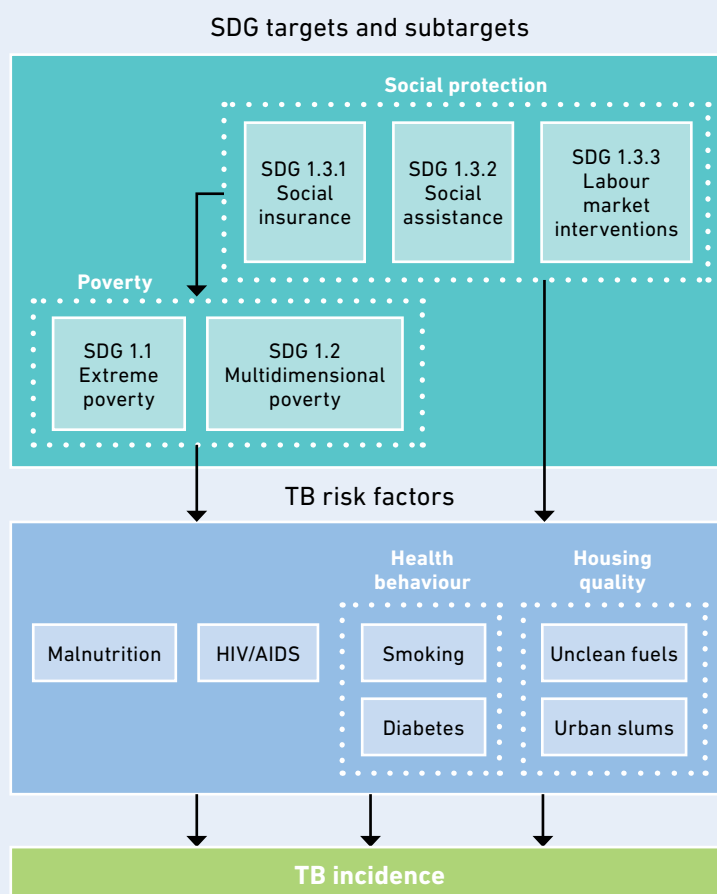
The impact of social protection and poverty elimination on global TB incidence: a modelling study

SDG 1 is to end poverty in all its forms everywhere, and associated targets include eliminating extreme poverty and ensuring nationally appropriate social protection systems and measures for all (including floors). Achievement of SDG 1 is likely to affect the TB epidemic through a range of pathways, as illustrated in Fig. B7.3.1.

A recent study used data from the SDG data repository and the WHO global TB database for 192 countries to explore how achievement of SDG 1 could affect TB incidence. Analyses were used to assess the strength of the association between the SDG 1 targets and TB incidence, as a basis for defining a simplified framework for modelling in which SDG 1 indicators were linked to TB incidence.

It was estimated that ending extreme poverty could reduce global TB incidence by 33% by 2035 (95% confidence interval: 16–45%), while expanding social protection coverage could reduce incidence by 76% (45–90%) by 2035. Together, both pathways were estimated to be able to reduce TB incidence by 84% (55–95%). Full achievement of SDG 1 could thus have a substantial impact on the global burden of TB.

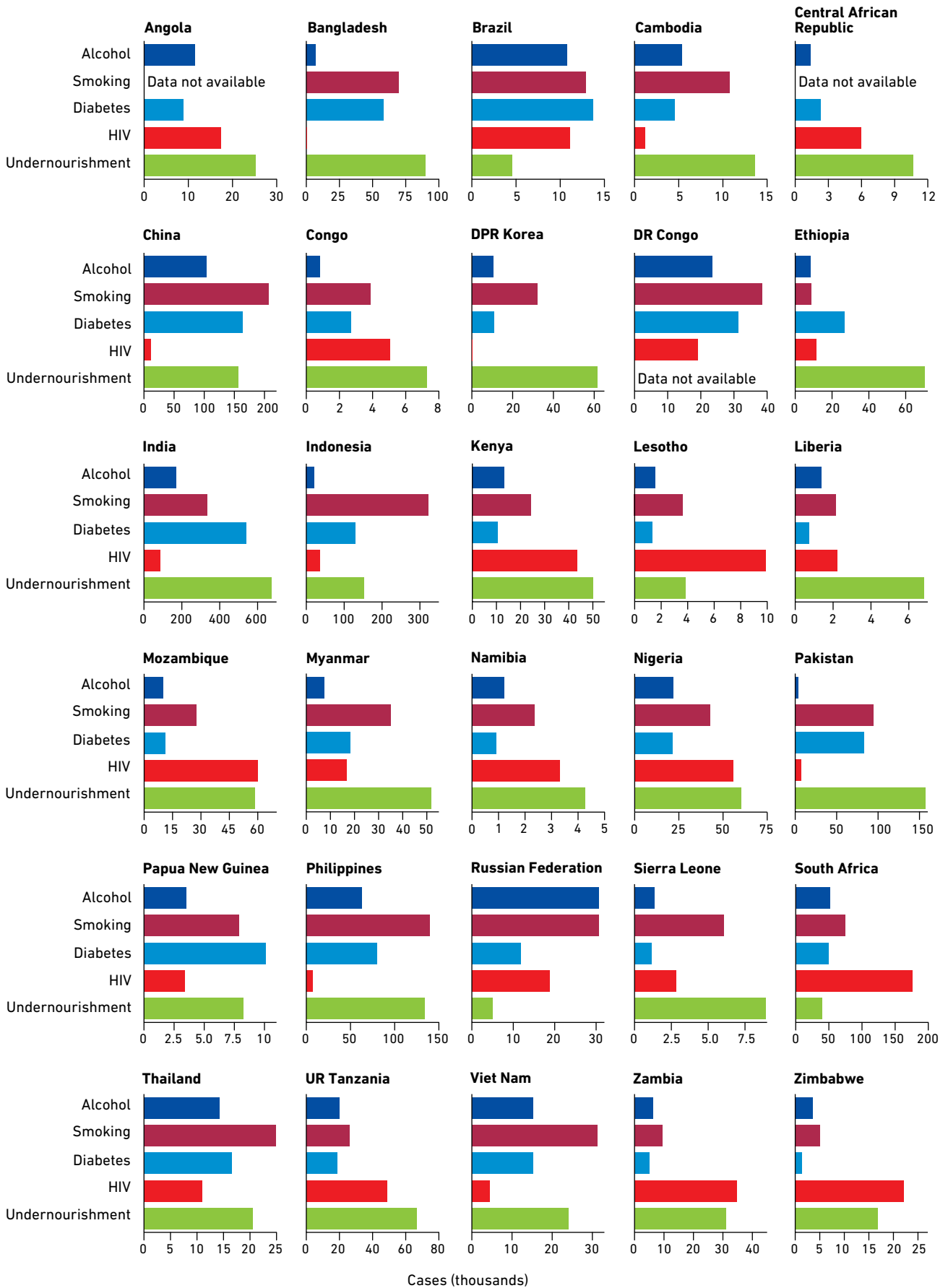
FIG. B7.3.1
Conceptual framework linking SDG 1 indicators to TB incidence



^a Carter DJ, Glaziou P, Lönnroth K, Siroka A, Floyd K, Weil D, Ravigliione M, Houben RMG, Boccia D. The impact of social protection and poverty elimination on global tuberculosis incidence: a statistical modelling analysis of Sustainable Development Goal 1. *Lancet Glob Health*. 2018 May;6(5):e514–22 (<https://www.ncbi.nlm.nih.gov/pubmed/29580761>, accessed 11 July 2018).

FIG. 7.8

Estimated number of TB cases attributable to five risk factors, 30 high TB burden countries, 2017



Sources: Lönnroth K, Castro KG, Chakaya JM et al. Tuberculosis control and elimination 2010–50: cure, care, and social development. Lancet. 2010 May 22;375(9728):1814–29; World Bank; and WHO Global TB Programme.

TABLE 7.4**TB cases attributable to selected risk factors, 2017**

RISK FACTOR	RELATIVE RISK ^a	EXPOSED (MILLIONS IN 2017)	GLOBAL POPULATION ATTRIBUTABLE FRACTION (%)	ATTRIBUTABLE TB CASES (MILLIONS IN 2017)
Undernourishment	3.1–3.3	734	18	1.9
HIV infection	20	36	8.7	0.88
Smoking	1.6–2.5	1 047	7.9	0.83
Diabetes	2.3–4.3	460	7.5	0.79
Harmful use of alcohol	1.9–4.6	407	4.7	0.49

^a Source: Lönnroth K, Castro KG, Chakaya JM et al. Tuberculosis control and elimination 2010–50: cure, care, and social development. *Lancet*. 2010 May 22;375(9728):1814–29. The relative risk for HIV infection is based on data from UNAIDS and estimates from this Global TB report.

Although it is typically not possible for NTPs to make progress on these issues alone, the UN high level meeting on TB in September 2018 (**Chapter 1**) and adaptation and use of the multisectoral accountability framework for TB developed in 2018 (**Chapter 2**) provide a basis for raising awareness and taking action on these issues.

Estimates of the number of incident TB cases attributable to five selected risk factors in 2017 are shown in **Table 7.4**. An estimated 1.9 million were attributable to undernourishment, 0.88 million to HIV infection, 0.83 million to smoking, 0.79 million to diabetes and 0.49 million to alcohol abuse. Applying the same method,

country-specific estimates of the number of incident TB cases attributable to the five risk factors in the 30 high TB burden countries are shown in **Fig. 7.8**. This figure shows that although various factors (undernutrition, the prevalence of alcohol use disorder, diabetes, HIV and smoking) contribute to the TB epidemic in the 30 high TB burden countries, there is considerable variation among countries in the relative contribution of these factors, and thus also variation in which factors need to be prioritized as part of national efforts to reduce the burden of TB disease.



A handheld DNA sequencer being used in Madagascar to identify TB resistance in sputum samples
Institut Pasteur de Madagascar

Chapter 8. TB research and development

KEY FACTS AND MESSAGES

Tuberculosis (TB) research and development is essential to achieve the global TB targets set in the Sustainable Development Goals and the End TB Strategy. A major technological breakthrough is required by 2025, so that the rate at which TB incidence falls can be dramatically accelerated compared with historic levels, to an average of 17% per year between 2025 and 2035.

Priorities for TB research and development include a vaccine to lower the risk of infection, a vaccine or new drug treatment to cut the risk of TB disease in the 1.7 billion people already latently infected, rapid diagnostics for use at the point of care and simpler, shorter drug regimens for treating TB disease.

“Intensified research and innovation” is the third pillar of the End TB Strategy.

A substantial increase in investment in TB research and development is needed. Funding has increased in recent years and reached a peak of US\$ 724 million in 2016, but this is only 36% of the estimated requirement of US\$ 2 billion per year.

The diagnostic pipeline is relatively stagnant. A small number of technologies emerged in 2017–2018 and several have not demonstrated adequate performance in field evaluation studies. There is still no single rapid, accurate and robust TB diagnostic test suitable for use at the point of care.

There are 20 drugs in Phase I, II or III trials for the treatment of drug-susceptible TB, multidrug-resistant TB or latent TB infection. There are 11 new compounds (an increase of 3 since August 2017): contezolid, delpazolid, GSK-3036656, macozinone, OPC-167832, pretomanid, Q203, SQ109, sutezolid, TBA-7371 and TBI-166. Two other drugs, bedaquiline and delamanid, have already received accelerated or conditional regulatory approval based on Phase IIb results. The seven repurposed drugs undergoing further testing are clofazimine, linezolid, levofloxacin, moxifloxacin, nitazoxanide, rifampicin (high dose) and rifapentine.

Various combination regimens with new or repurposed drugs are in Phase II or Phase III trials.

Twelve vaccine candidates are in clinical trials: four in Phase I, six in Phase II and two in Phase III. They include candidates to prevent the development of TB infection and disease, and candidates to help improve the outcomes of treatment for TB disease.

The global TB targets set in the Sustainable Development Goals (SDGs) and the End TB Strategy cannot be achieved without tuberculosis (TB) research and development. The SDG target is to “end the epidemic” by 2030; more specific targets for 2030 set in the End TB Strategy are a 90% reduction in TB deaths and an 80% reduction in TB incidence compared with levels in 2015, with targets for further reductions (95% and 90%, respectively) by 2035. Reaching these targets requires a major technological breakthrough by 2025, so that the rate at which TB incidence falls can be dramatically accelerated compared with historic levels, to an average of 17% per year from 2025–2035 (**Chapter 2**). Priorities for TB research and development include a vaccine to lower the risk of infec-

tion, a vaccine or new drug treatment to cut the risk of TB disease in the 1.7 billion people already latently infected,¹ rapid diagnostics for use at the point of care and simpler and shorter drug regimens for treating TB disease.^{2,3}

¹ Houben RM, Dodd PJ. The global burden of latent tuberculosis infection: a re-estimation using mathematical modelling. *PLoS Med*. 2016;13(10):e1002152 (<https://www.ncbi.nlm.nih.gov/pubmed/27780211>, accessed 30 July 2018).

² Shortening treatment regimens as well as decreasing treatment toxicity are particularly high priorities for drug-resistant TB.

³ WHO, in collaboration with partners has developed Target Product Profiles for TB treatment regimens (referred to as Target Regimen Profiles or TRPs). For further details, see: World Health Organization. Target Regimen Profiles for TB Treatment (WHO/HTM/TB/2016.16). Geneva: WHO; 2016. (<http://apps.who.int/iris/bitstream/handle/10665/250044/9789241511339-eng.pdf;jsessionid=5F6709C0669BB45006E43153C2B4DF06?sequence=1>, accessed 30 July 2018).

“Intensified research and innovation” is one of the three pillars of the WHO End TB Strategy. The two main components of this pillar are “discovery, development and rapid uptake of new tools, interventions and strategies” and “research to optimize implementation and impact, and promote innovations” (Chapter 2).

Technological breakthroughs and optimal use of available interventions require greater investment in TB research and development. The Stop TB Partnership’s “Global Plan to End TB 2016–2020” estimates that about US\$ 2 billion per year is needed.¹ To date, funding has not come close to this amount. In the 5 most recent years for which data have been published, annual funding was in the range of US\$ 600–700 million, reaching a peak of US\$ 724 million in 2016.² This was equivalent to only about 40% of the amount required. Although TB accounts for nearly 2% of disability-adjusted life-years (DALYs) lost, it receives only 0.25% of the estimated US\$ 265 billion spent on medical research annually,³ and less support than other global health threats such as HIV and malaria. Further details about recent funding for TB research and development are provided in Box 8.1.

Recognizing the need to accelerate efforts in TB research and development, “Pursuing science, research and innovation” was one of the four major topics of the Moscow Declaration to End TB. The commitments from Member States and the calls to partners included in the declaration are listed in Box 8.2. In April 2018, TB research and development was one of the topics prioritized for discussion at the first United Nations (UN) high-level meeting on TB in September 2018.⁴

This chapter is not intended as an exhaustive overview of current or recently completed TB research. As in previous global TB reports, it focuses on providing an overview of progress in the development of new TB diagnostics (Section 8.1), drugs (Section 8.2) and vaccines (Section 8.3), which account for most of the funding invested in TB research and development in recent years (Box 8.1). The status of the pipelines as of August 2018, based on recent publications as well as communications with the secretariats of the relevant working groups of the Stop TB Partnership and other stakeholders, is described and discussed. The last part of the chapter (Section 8.4) highlights how global and intercountry collaborations can help to expedite progress in TB research.

¹ The Global Plan to End TB, 2016–2020. Geneva: Stop TB Partnership; 2015 (<http://www.stoptb.org/global/plan/>, accessed 21 June 2018).

² Treatment Action Group. The ascent begins: tuberculosis research funding trends, 2005–2016. New York: Treatment Action Group; 2015 (<http://treatmentactiongroup.org/tbrd2017>, accessed 28 June 2018).

³ Moses H, Matheson DH, Cairns-Smith S, George BP, Palisch C, Dorsey ER. The anatomy of medical research: US and international comparisons. *JAMA*. 2015;313(2):174–89 (<https://www.ncbi.nlm.nih.gov/pubmed/25585329>, accessed 9 July 2018).

⁴ United Nations General Assembly. Resolution adopted by the General Assembly on 4 April 2018. A/RES/72/268. Scope, modalities, format and organization of the high level meeting on the fight against tuberculosis (http://www.un.org/en/ga/search/view_doc.asp?symbol=A/RES/72/268, accessed 9 July 2018).

BOX 8.1

Global investments in TB research and development: a recent analysis

As part of preparations for the first Global Ministerial Conference on TB held in November 2017, recent investments in TB research and development were assessed.^a

Key findings were as follows:

- A total of US\$ 4.6 billion was invested from 2009–2016, mostly for development of new diagnostics, drugs and vaccines (61%). The rest was for basic research (21%), operational research (10%) and infrastructure or “unspecified” projects (8%).
- Most funding from 2009 to 2016 was from a few countries, led by the United States of America (USA, with US\$ 1.8 billion), followed by the European Union and United Kingdom (US\$ 269 million each). Contributions from other countries did not exceed US\$ 62 million.
- The two largest funders were the US National Institutes of Health (NIH) and the Bill & Melinda Gates Foundation. Combined, they accounted for 37% of total investments from 2005 to 2015.
- Spending by the pharmaceutical industry decreased considerably, following the withdrawal of several companies from the field and the maturation of clinical development programmes for new TB drugs.
- Between 2011 and 2015, the top five recipients of funding were either product-development partnerships or research institutes and universities in the USA. Other recipients in the top 15 included the Medical Research Council (United Kingdom), the Foundation for Innovative New Diagnostics (FIND, Switzerland), the University of Cape Town (South Africa), the International Union Against Tuberculosis and Lung Disease (based in France) and the National Institute for Research in Tuberculosis (India).
- Insufficient investment in operational research has compromised the scale-up of new technologies and approaches, limiting their impact on the TB epidemic. Consistent with this finding, only 13% of all TB research publications in the past decade were related to operational or public health research.

^a World Health Organization. Global investments in tuberculosis research and development past, present, and future. Geneva: WHO; 2017 (<http://apps.who.int/iris/bitstream/handle/10665/259412/9789241513326-eng.pdf>, accessed 28 June 2018).

BOX 8.2

Commitments by Member States and calls to partners listed under the heading “Pursuing science, research and innovation” in the Moscow Declaration to End TB^a

Commitments by Member States

- Increasing national or regional capacity and funding, as needed, to urgently expand multidisciplinary TB research and innovation, and applied health research, by establishing and strengthening national TB research networks, including civil society and community-based mechanisms, considering TB research as a central element of national TB and research and development strategies, expanding existing networks to integrate TB research, and reducing research- and implementation-related regulatory impediments.
- Working, when relevant, across ministries, donors, the scientific community, the private sector, academia and other key stakeholders for the purpose of research:
 - for development and evaluation of rapid point-of-care diagnostics; new and more effective drugs; and shorter, high-quality and cost-effective treatment regimens for all forms of TB – including latent TB infection and drug-resistant TB – and; safe and effective TB vaccines by 2025; and
 - on environmental and social determinants of TB and effective interventions strategies.
- Improving, as appropriate, the coordination of research efforts nationally and globally, and ensuring that the emerging knowledge is promptly put into action, including by putting in place appropriate policy frameworks and implementing new medical technologies.
- Strengthening, as appropriate, surveillance systems, improving data collection and reporting at all levels, using innovative approaches and including surveillance in TB research agendas.

Calls to partner agencies

- WHO – in collaboration with global partners, research organizations, donors, the scientific community and countries – to consider developing a Global Strategy for TB Research, taking into consideration ongoing and new efforts, such as the TB Research Network stated in the BRICS Leaders Xiamen Declaration.^{b,c}

- WHO – in collaboration with global health and research partners and countries – to make further progress in enhancing cooperation and coordination of TB research and development, considering (where possible) drawing on existing research networks to integrate TB research, such as the new antimicrobial resistance (AMR) R&D collaboration hub proposed in the 2017 G20 Leaders' Declaration,^d in particular, to facilitate rapid scale-up of innovative approaches and tools for TB prevention, diagnosis, treatment and care.

A TB resolution adopted at the World Health Assembly in May 2018 reinforced the Moscow Declaration's call for the development of a global strategy for TB research and innovation, and requested WHO to develop such a strategy in collaboration with other stakeholders.^e The aim of the strategy is to accelerate progress in TB research and development, through strategic approaches such as fostering global collaborations, stimulating the development of innovative financing mechanisms for the most urgent priorities, promoting the sharing of data and information, and facilitating the rapid scale-up of new tools.

^a Moscow Declaration to End TB; First WHO Global Ministerial Conference on Ending TB in the Sustainable Development Era: A Multisectoral Response, November 2017. WHO and the Ministry of Health of the Russian Federation. http://www.who.int/tb/features_archive/Moscow_Declaration_to_End_TB_final_ENGLISH.pdf?ua=1, accessed 21 June 2018).

^b The BRICS countries are Brazil, Russian Federation, India, China and South Africa.

^c BRICS Leaders Xiamen Declaration. China: Ministry of Foreign Affairs of the People's Republic of China; 2017 (https://www.brics2017.org/English/Documents/Summit/201709/t20170908_2021.html, accessed 11 July 2018).

^d G20 Nations. G20 Leaders' Declaration: Shaping an interconnected world. Hamburg; G20 Nations; 2017 (<http://www.g20.utoronto.ca/2017/2017-G20-leaders-declaration.html>, accessed 11 June 2018).

^e World Health Organization. Preparation for a high-level meeting of the General Assembly on ending tuberculosis (WHA71.13), Seventy-first World Health Assembly. Geneva: WHO; 2018 (http://apps.who.int/gb/ebwha/pdf_files/WHA71/A71_R3-en.pdf, accessed 11 July 2018).

8.1 New diagnostics for TB

This section starts by providing an overview of the TB diagnostics pipeline, with particular attention to progress made in 2018. It then describes diagnostic tests, products and methods related to the detection of TB infection, disease and drug resistance that were reviewed by WHO in 2018, or that are scheduled for evaluation in 2019.

8.1.1 An overview of the diagnostics pipeline

Although the TB diagnostics pipeline appears robust, with several technologies in development, no new platform has emerged in 2018. Despite the promise from several manufacturers, there is still no single rapid, accurate and robust TB diagnostic test suitable for use at the point of care. This reflects serious underinvestment (especially in basic science), and persistent scientific and technical challenges that affect the development of new TB diagnostics.

An overview of the pipeline for TB diagnostics in August 2018 is shown in **Fig. 8.1**. The list of technologies is not necessarily complete, but does reflect technologies that have been documented in previous reports published by the Treatment Action Group¹ and Unitaid.² Technologies under development are primarily molecular-based; they include tests to detect TB for diagnosis and treatment monitoring, to detect drug resistance among confirmed TB cases, and to detect TB and drug resistance simultaneously in centralized high-throughput laboratories.

The pipeline remains largely unchanged from the one published in the 2017 edition of the global TB report,³ and highlights the urgent need for new technologies to minimize barriers to health-care access, ensure quality testing for difficult-to-diagnose groups, expand the spectrum of drug susceptibility testing (DST), and reduce the costs of diagnostic platforms and their maintenance.

The development of the Cepheid close-to-care platform – GeneXpert® Omni® (Omni) – continues to be delayed and is not expected to be available before 2019. Cepheid has developed an alternative to Omni – the GeneXpert® Edge® (Edge). The Edge is a single-module GeneXpert instrument connected to a tablet device for transfer of data; its specific features include an auxiliary battery that allows operation in more decentralized settings, at the same level as microscopy. The instrument is being developed to facilitate wider access to rapid molecular testing for TB and rifampicin resistance, and virology parameters for HIV and hepatitis C virus, for

example. The Edge is expected to be available from November 2018.

The next-generation Xpert MTB/RIF Ultra (Ultra) cartridge has shown significantly better performance (increased sensitivity) than the current Xpert MTB/RIF cartridge in detecting *Mycobacterium tuberculosis* in specimens with low numbers of bacilli. This was particularly the case for smear-negative, culture-positive specimens (e.g. those from people living with HIV), extrapulmonary specimens (notably cerebrospinal fluid) and specimens from children.⁴

Despite the improved performance of the Ultra cartridge, as confirmed by WHO, the transition to this next-generation cartridge has been limited, in part, due to the short shelf-life of the cartridge. South Africa is the only high TB burden country currently using Ultra as the initial TB diagnostic test. WHO has commissioned an updated systematic review of the performance of Ultra for use in the diagnosis of pulmonary and extrapulmonary TB in adults and children, and expects to refine and update policy guidelines for the use of Ultra in 2019.

Two cartridge-based technologies being developed in China and India, respectively, are in the pipeline. These technologies have the potential to be used as an alternative to the GeneXpert platform in primary health-care facilities. Both are similar to Xpert MTB/RIF in that they target the multicopy IS6110 and a single-copy gene target. Evidence about the performance of both technologies from well-designed validation studies is needed to enable WHO to review and assess their performance; ancillary solutions in areas such as maintenance and connectivity would also be needed to support roll-out.

The first of these two assays, the Truenat MTB assays® (Molbio Diagnostics, Bangalore, India), uses chip-based real-time polymerase chain reaction (PCR) for the detection of TB. A reflex assay has been developed to detect resistance to rifampicin in samples with positive results for *M. tuberculosis*. The Indian Council of Medical Research has recommended Truenat MTB to the Indian Central TB Division, based on local data. The Foundation for Innovative New Diagnostics (FIND) is planning a multicentre study of diagnostic accuracy at the level of peripheral microscopy centres, to assess the performance characteristics of the test in different epidemiological and geographical settings; the results will inform a WHO review.

Ustar Biotechnologies (China) has developed the EasyNAT TB® assay based on isothermal cross-priming amplification. The assay uses preloaded reagents in a single cartridge for integrated extraction, amplification and detection, using two separate chambers. Testing,

¹ Lessem E. The tuberculosis diagnostics pipeline. Treatment Action Group; 2017 (<http://www.pipelinerreport.org/sites/default/files/2017-Pipeline-Report-TB-Diagnostics.pdf>, accessed 22 June 2018).

² Unitaid. Tuberculosis – diagnostics technology landscape, 5th edition. Geneva: WHO; 2017 (<https://unitaid.eu/assets/2017-Unitaid-TB-Diagnostics-Technology-Landscape.pdf>, accessed 22 June 2018).

³ World Health Organization. Global tuberculosis report 2017 (WHO/HTM/TB/2017.23). Geneva: WHO; 2017 (<http://apps.who.int/iris/bitstream/handle/10665/259366/9789241565516-eng.pdf>, accessed 21 June 2018).

⁴ WHO meeting report of a technical expert consultation: non-inferiority analysis of Xpert MTB/RIF Ultra compared to Xpert MTB/RIF. Geneva: World Health Organization; 2017 (WHO/HTM/TB/2017.04; <http://apps.who.int/iris/bitstream/10665/254792/1/WHO-HTM-TB-2017.04-eng.pdf>, accessed 1 May 2018).

FIG. 8.1

An overview of progress in the development of TB diagnostics, August 2018^a

TECHNOLOGIES IN DEVELOPMENT	TECHNOLOGIES ENDORSED BY WHO	SCHEDULED FOR WHO EVALUATION IN 2018/19
<p>Molecular detection of TB and drug resistance</p> <ul style="list-style-type: none"> ■ Gendrive MTB/RIF ID, Epistem, UK ■ Xpert XDR-TB cartridge, Cepheid, USA ■ TruArray MDR-TB, Akkoni, USA ■ INFINITIMTB Assay, AutoGenomics, USA ■ FluoroType XDR-TB assay, Hain Lifescience, Germany ■ MeltPro TB assay, Zeesan Biotech, China ■ QuantuMDx, POC, UK <p>Tests for latent TB infection</p> <ul style="list-style-type: none"> ■ Diaskin test, Generium, Russian Federation ■ C-Tb test, Serum Institute of India, India 	<p>Molecular detection of TB and drug resistance</p> <ul style="list-style-type: none"> ■ Line probe assays for the detection of <i>Mycobacterium tuberculosis</i> (MTB), isoniazid and rifampicin resistance in acid-fast bacilli smear positive sputum or MTB cultures (FL-LPA), Hain Lifescience, Germany and Nipro, Japan ■ Line probe assays for the detection of resistance to fluoroquinolones and second-line injectable agents (SL-LPA), Hain Lifescience, Germany ■ TB LAMP for detection of TB, Eiken, Japan <p>Nonmolecular technologies</p> <ul style="list-style-type: none"> ■ Inteferon gamma release assay (IGRAs) for the diagnosis of latent TB infection (LTBI) Oxford Immunotec, UK, Qiagen, USA <p>Culture-based technologies</p> <ul style="list-style-type: none"> ■ Commercial liquid culture systems and rapid speciation ■ Culture-based phenotypic DST using 1% critical proportion in LJ,7H10,7H11 and MGIT media. <p>Microscopy</p> <ul style="list-style-type: none"> ■ Light and light-emitting diode microscopy (diagnosis and treatment monitoring) 	<p>Molecular detection of TB and drug resistance</p> <ul style="list-style-type: none"> ■ Molecular technologies for genotypic drug resistance testing (including sequencing technologies) ■ FluoroType MTBDR, Hain Lifescience, Germany ■ m2000 RealTime MTB System, Abbott, USA ■ BD Max MDR-TB, Becton Dickinson, USA ■ Roche cobas® MTB system, Roche Diagnostics, Basel, Switzerland <p>Radiology</p> <ul style="list-style-type: none"> ■ Computer aided detection (CAD)
ON THE MARKET (EVIDENCE FOR USE NOT SUBMITTED TO WHO FOR EVALUATION)		WHO POLICY UPDATES SCHEDULED FOR 2018/2019
<p>Molecular detection of TB and drug resistance</p> <ul style="list-style-type: none"> ■ iCubate System, iCubate, USA ■ Genechip, TB drug resistance array, Capital Bio, China ■ EasyNAT TB Diagnostic kit, Ustar Biotechnologies, China ■ Truelab/Truenat MTB, Molbio/bigtec Diagnostics, India <p>Culture-based drug susceptibility testing</p> <ul style="list-style-type: none"> ■ Sensititre™ MYCOTBI plate; ThermoFisher Scientific Inc., USA 		<p>Molecular detection of TB and drug resistance</p> <ul style="list-style-type: none"> ■ Alere Determine TB-LAM, Alere, USA (TB detection in people seriously ill with HIV) ■ Xpert MTB/RIF Ultra for detection of TB and rifampicin resistance in pulmonary, extrapulmonary and paediatric samples, Cepheid, USA

^a The list of technologies is not necessarily complete, but does reflect technologies that have been documented in previous reports published by the Treatment Action Group¹ and Unitaid²

¹ Lessem E. The tuberculosis diagnostics pipeline. Treatment Action Group; 2017 (<http://www.pipelinerreport.org/2017/tbdx>, accessed 27 June 2018).

² Unitaid. Tuberculosis – diagnostic technology landscape, 5th edition. Geneva: World Health Organization; 2017 (<https://unitaid.eu/assets/2017-Unitaid-TB-Diagnostics-Technology-Landscape.pdf>, accessed 27 June 2018).

which is currently limited to detection of TB (drug resistance cannot be detected), can be performed by loading cartridges into a battery-operated device for testing two samples simultaneously.

8.1.2 Critical concentrations for phenotypic drug-susceptibility testing of medicines used in the treatment of drug-resistant TB

In May 2018, WHO and FIND released a technical report with updated laboratory parameters for DST of medicines used to treat drug-resistant TB.¹ Culture-based phenotypic testing is the current reference method for DST of anti-TB medicines, and it relies on the testing of so-called critical concentrations of drugs, that is, the lowest concentration of an anti-TB medicine in vitro that will inhibit the growth of 99% of phenotypically wild-type strains of *M. tuberculosis*.

The report summarizes the current evidence used to

¹ Technical report on critical concentrations for drug susceptibility testing of medicines used in the treatment of drug-resistant tuberculosis. Geneva: WHO; 2018 (WHO/CDS/TB/2018.5) (<http://apps.who.int/iris/bitstream/10665/260470/1/WHO-CDS-TB-2018.5-eng.pdf> accessed 1 May 2018).

determine critical concentrations in different culture media for performing DST for specific medicines used in the treatment of drug-resistant TB. Critical concentrations for the new and repurposed anti-TB agents bedaquiline, delamanid, linezolid and clofazimine are included, in addition to revised consensus critical concentrations for fluoroquinolones and second-line injectable agents.

8.1.3 TB diagnostic tests, products and methods that may be evaluated by WHO in 2019

In 2019, there may be sufficient data for WHO to reassess the critical concentrations to use when testing for resistance to isoniazid and rifampicin, and to determine the role of DNA-sequencing technologies in the diagnosis of drug-resistant TB. There may also be sufficient data to review the performance of the lateral flow lipoarabinomannan assay (LF-LAM), centralized high-throughput testing platforms, molecular sequencing as a reference standard for DST, and a new test for extensively drug-resistant TB (XDR-TB).²

² XDR-TB is defined as MDR-TB plus resistance to at least one fluoroquinolone and a second-line injectable.

Critical concentrations for phenotypic drug-susceptibility testing of isoniazid and rifampicin

Anti-TB medicines such as isoniazid and the rifamycins (rifampicin, rifapentin and rifabutin) are essential for TB treatment. There is emerging evidence that some of the critical concentrations used to test for resistance to these medicines, established historically in solid culture media, may not necessarily apply to commercial liquid culture systems. Therefore, WHO has commissioned FIND to undertake a systematic review of available minimum inhibitory concentration data for phenotypically wild-type and non wild-type strains, including sequencing data for relevant resistance genes. The goal of the review is to evaluate the available evidence and, if indicated, to revise the critical concentrations used to test for resistance to these first-line anti-TB medicines on selected media.

Role of DNA-sequencing technologies

The successful diagnosis and treatment of drug-resistant TB depends upon universal access to accurate DST. Conventionally, the diagnosis of drug resistance in *M. tuberculosis* strains has relied heavily upon culture and DST in liquid or solid media in TB containment laboratories. Phenotypic results are only obtained after weeks to months of incubation, and establishing the stringent laboratory biosafety conditions required for these culture-based methods is a challenge.

Drug resistance in the *Mycobacterium tuberculosis* complex is mainly conferred through point mutations in specific gene targets in the bacterial genome. Given the challenges associated with performing phenotypic DST, molecular tests are increasingly being used to allow more rapid testing and thus earlier initiation of appropriate treatment for drug-resistant TB.

Rapid molecular tests such as Xpert MTB/RIF and line probe assays (LPAs) allow for the detection of a set of common resistance mutations in a few gene regions. However, for many drugs used in the treatment of drug-resistant TB, the mutations are spread over multiple gene regions. Moreover, not all mutations conferring resistance have been identified, and limited information is available for identifying the “high-confidence mutations” that would accurately predict clinical treatment outcomes.

An advantage of sequencing technologies, compared with the rapid molecular tests that are currently available, is that they are able to provide detailed information on resistance mutations for multiple gene regions. A first step that has been taken is to grade mutations according to the level of confidence in the association between resistance mutations and the conferred phenotypic drug resistance. FIND has recently completed a systematic review of the association between *M. tuberculosis* strains with specific mutations and minimal inhibitory concentrations data for drugs of interest. It is anticipated that WHO will be able to use this review to assess evidence

about the accuracy of genotypic DST compared with current phenotypic reference standards, and to develop a related list of priority mutations (so-called high-confidence mutations), which could also be used to aid test development.

WHO and FIND are preparing a technical guide that summarizes the characteristics of available DNA-sequencing technologies, along with current knowledge of the genetic basis of TB drug resistance. This guide will provide a foundation for building additional knowledge related to the molecular basis of resistance, and will inform future WHO policy guidelines on the use and interpretation of DNA-sequencing technologies.

The Relational Sequencing TB Knowledgebase (ReSeqTB) is a global, clinical knowledge base that curates, standardizes and unifies genotypic and phenotypic DST data, along with metadata on drug-resistant TB.¹ The database is regarded as a “living” knowledge base, in which sequencing, phenotypic and clinical outcome data for existing, repurposed and new TB drugs are continually gathered from countries and other stakeholders.

Lateral flow lipoarabinomannan assay

Tests based on the detection of the mycobacterial lipoarabinomannan (LAM) antigen in urine are available as potential point-of-care tests for TB. Urinary LAM assays are unsuitable for use as general screening tests for TB due to suboptimal sensitivity. However, compared with traditional diagnostic methods, they have demonstrated improved sensitivity for the diagnosis of TB among individuals coinfecting with HIV, especially among patients with low CD4 counts.

The urine LAM strip-test (Determine®-TB Alere, USA) is currently recommended by WHO in HIV-positive adults with CD4 counts less than or equal to 100 cells/UL and with signs and symptoms of TB.² Since 2015, new evidence has emerged that might justify the use of the test in a broader group of people living with HIV. WHO has initiated a review of the evidence and associated guidance, and a meeting to review the evidence is due to be held before the end of 2018.

Centralized high-throughput testing platforms

Several manufacturers have developed centralized testing PCR-based platforms suitable for high laboratory throughput. If sufficient evidence becomes available, the performance of these platforms could be assessed before the end of 2018. Examples of the platforms in development are the RealTime® platform, developed by Abbott; the FluoroType MTBDR®, launched by Hain

¹ <http://reseqtb.org> (accessed 19 April 2018).

² World Health Organization. The use of lateral flow urine lipoarabinomannan assay (LF-LAM) for the diagnosis and screening of active tuberculosis in people living with HIV: policy guidance (WHO/HTM/TB/2015.25). Geneva: WHO; 2015; http://apps.who.int/iris/bitstream/10665/193633/1/9789241509633_eng.pdf?ua=1&ua=1, accessed 1 May 2018).

Lifescience in early 2017; and platforms being developed by Becton Dickinson and Roche.

WHO, in consultation with FIND, has developed a two-step evaluation process for this purpose. The first step includes verification of the accuracy of the assays in detecting *M. tuberculosis* as well as resistance to rifampicin and isoniazid, based on independent evaluation of a standardized panel of strains, conducted at one or more WHO TB supranational reference laboratories (SRLs). The second step is evaluation of the clinical validity of the assays, based on testing of the platforms in two or three national reference laboratories in high TB burden settings, and comparing results with the reference standards of culture, phenotypic DST and molecular sequencing, as well as with Xpert MTB/RIF.

Tests for extensively drug-resistant TB

Molecular LPAs for second-line anti-TB medicines are currently the only rapid molecular tests recommended by WHO for detecting whether patients with confirmed multidrug-resistant TB (MDR-TB) or rifampicin-resistant TB have XDR-TB. However, LPAs require sophisticated laboratory infrastructure and equipment that is normally only available in centralized laboratories.

Cepheid is continuing to develop an XDR-TB cartridge to detect resistance to isoniazid, injectable agents and fluoroquinolones. If sufficient data become available, WHO will evaluate the XDR-TB cartridge using the same two-step approach described above for centralized high-throughput testing platforms.

Computer-aided detection systems

Chest radiography, or chest X-ray (CXR), is an important tool for TB triaging and screening, and is also a useful aid in the diagnosis of TB. A major limitation of CXR is that it requires experienced interpreters (usually radiologists or technicians) for interpreting the images. Even among trained readers, variation between readers is common, which further affects the accuracy of CXR. In many countries, few experienced CXR readers are available.

In recent years, computer-aided detection (CAD) systems have been developed that use digital technology to detect physiological and pathological conditions, including TB. These CAD systems incorporate computer algorithms that analyse a digital CXR and produce a standardized interpretation of the image. A score or report is generated that estimates the likelihood that the image is from an individual with active TB disease. CAD systems are trained on thousands of images, using machine-learning techniques. A research agenda to assess CAD systems is underway¹. FIND is establishing an archive of digital x-rays to assess the performance

of commercially available CAD solutions. If sufficient data become available, WHO will evaluate the use of CAD systems for detecting TB in early 2019.

Biomarkers for treatment monitoring

DNA-based molecular diagnostic tools provide rapid results for the diagnosis of active TB disease. Tests such as the Xpert MTB/RIF assay have a sensitivity approaching that of culture, although they remain unsuitable for treatment monitoring because they detect dead bacilli.²

St Andrew's University, Scotland has developed a novel approach for real-time monitoring of a patient's response to TB treatment. The Molecular Bacterial Load Assay (MBLA) is a real-time reverse transcriptase quantitative PCR (RT-qPCR) that uses 16S-ribosomal RNA (rRNA) targets to quantify the *M. tuberculosis* bacillary load over the course of treatment.

The 16S-rRNA targets are more abundant and stable than messenger RNA (mRNA), and yet are reduced during TB treatment, which makes this assay more sensitive for *M. tuberculosis* detection and monitoring of the response to treatment. Its ability to quantify a patient's bacillary load of *M. tuberculosis* in real time, and avoid the challenges related to culture e.g. time to results and contamination concerns, gives MBLA the potential to eventually replace microscopy and culture as a test for treatment monitoring. To date, the assay has been evaluated in four high TB burden, low-resource settings in Africa.³

The next crucial step is the development of a design-locked product that allows standardized evaluation. Laboratory-based studies of analytical sensitivity and stability need to be performed to further assess the limits of detection of the assay, and the correlation with currently recommended technologies for monitoring TB treatment response (especially culture). If the laboratory-based studies demonstrate good performance, clinical evaluation studies will be needed to assess the MBLA as a possible replacement for microscopy and culture monitoring during treatment.

8.2 New drugs and drug regimens to treat TB

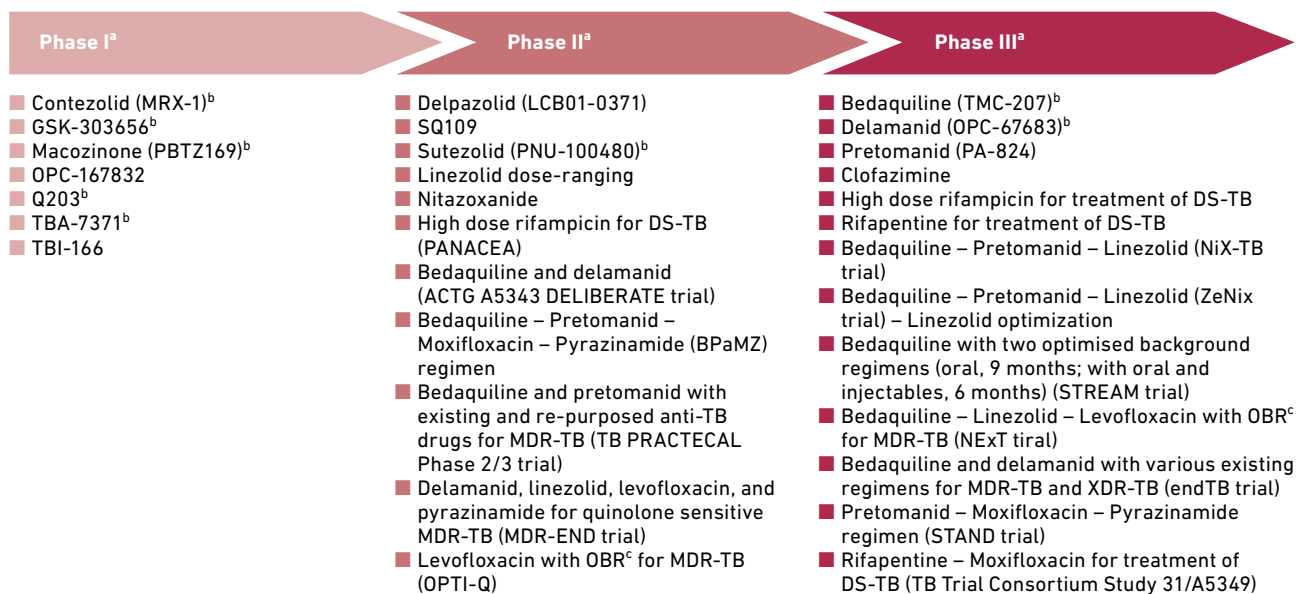
The pipeline for new anti-TB drugs in August 2018 is shown in Fig. 8.2. The pipeline has expanded in recent months, and there are now 20 drugs in Phase I, II or III trials, compared with 17 in August 2017.⁴ There are

¹ Ahmad Khan F, Pande T, Tessema B, Song R, Benedetti A, Pai M et al. Computer-aided reading of tuberculosis chest radiography: Moving the research agenda forward to inform policy. *European Respiratory Journal* 2017; 50(1) (<http://erj.ersjournals.com/content/50/1/1700953>). long, accessed 9 July 2018).

² World Health Organization. Automated real-time nucleic acid amplification technology for rapid and simultaneous detection of tuberculosis and rifampicin resistance: Xpert MTB/RIF assay for the diagnosis of pulmonary and extrapulmonary TB in adults and children – policy update (WHO/HTM/TB/2013.16). Geneva: WHO; 2013 (http://apps.who.int/iris/bitstream/10665/112472/1/9789241506335_eng.pdf, accessed 1 May 2018).

³ Gillespie SH, Sabiti W, Oravcova K. Mycobacterial load assay. *Methods Mol Biol.* 2017;1616:89–105 (<https://www.ncbi.nlm.nih.gov/pubmed/28600763>, accessed 9 July 2018).

⁴ P 139 of World Health Organization. Global tuberculosis report 2017 (WHO/HTM/TB/2017.23). Geneva: WHO; 2017 (<http://apps.who.int/iris/bitstream/handle/10665/259366/9789241565516-eng.pdf>, accessed 21 June 2018).

FIG. 8.2**The global clinical development pipeline for new anti-TB drugs and regimens, August 2018**

^a New drug compounds are listed first, followed by repurposed drugs and then by regimens.

^b New chemical class.

^c Optimized background regimen.

Source: Adapted from the Working Group on New TB Drugs pipeline. More information on these products and other ongoing projects can be found at <http://www.newtbdrugs.org/pipeline.php>

11 new compounds (up by three since August 2017): contezolid, delpazolid, GSK-3036656, macozinone, OPC-167832, pretomanid, Q203, SQ109, sutezolid, TBA-7371 and TBI-166.¹ Two other drugs (bedaquiline and delamanid) have already received accelerated or conditional regulatory approval based on Phase IIb trial results. Seven repurposed drugs are undergoing further testing: clofazimine, linezolid, levofloxacin, moxifloxacin, nitazoxanide, rifampicin (high dose) and rifapentine. The 20 compounds are described in more detail in [Section 8.2.1](#) and [Section 8.2.2](#).

New TB regimens are also being tested. These are described in [Section 8.2.3](#).

8.2.1 New compounds

Bedaquiline

WHO issued interim policy guidance on the use of bedaquiline for the treatment of adults with MDR-TB in 2013, based on Phase IIb trial results.² The recommendation to use bedaquiline as part of longer MDR-TB treatment regimens was conditional upon proper patient selection, a regimen design following WHO recommendations, close

monitoring of treatment, active TB drug safety monitoring and management, and informed consent according to local requirements. The recommendation was maintained following a review of data from observational studies in 2016.³ In 2018, additional data for patients treated with bedaquiline-containing regimens were analysed as part of a major update to WHO guidance on the treatment of drug-resistant TB.⁴

The safety and efficacy of bedaquiline when used as part of short MDR-TB regimens (i.e. 6 and 9 months duration), compared with the updated current standard of care recommended by WHO (i.e. a shortened regimen) is being investigated in the second stage of the Phase III Standardised Treatment Regimen of Anti-TB Drugs for Patients with MDR-TB (STREAM) trial.⁵ Recruitment started in March 2016, and the first results are expected in 2020. A study on the use of bedaquiline to treat children with MDR-TB is being implemented in the Philippines, the Russian Federation and South Africa. Bedaquiline is also being used in trials of all-oral treatment regimens ([Section 8.2.3](#)), and investigation of its use in the treatment of drug-susceptible TB has started.

¹ Most of the new compounds are being developed by not-for-profit organizations, academic institutions, small businesses or government agencies that lack the secure funding and resources available to major pharmaceutical companies. This makes the process of progression through trials and then registration more uncertain.

² World Health Organization. The use of bedaquiline in the treatment of multidrug-resistant tuberculosis: interim policy guidance (WHO/HTM/TB/2013.6). Geneva: WHO; 2013 (http://apps.who.int/iris/bitstream/10665/84879/1/9789241505482_eng.pdf, accessed 12 June 2018).

³ World Health Organization. Report of the Guideline Development Group Meeting on the use of bedaquiline in the treatment of multidrug-resistant tuberculosis, 2016 revision (WHO/HTM/TB/2017.01). Geneva: WHO; 2017 (<http://apps.who.int/iris/bitstream/10665/254712/1/WHO-HTM-TB-2017.01-eng.pdf>, accessed 9 July 2018).

⁴ Further details are provided in [Box 4.2](#) in [Chapter 4](#).

⁵ Moodley R, Godec TR, Team ST. Short-course treatment for multidrug-resistant tuberculosis: the STREAM trials. *Eur Respir Rev*. 2016;25(139):29–35 (<https://www.ncbi.nlm.nih.gov/pubmed/26929418>, accessed 12 June 2018).

Contezolid

MRX4, a prodrug¹ of Contezolid (MRX-I), is in a Phase 1 trial in the USA. MRX-1 is an oxazolidinone antibiotic that is potent against Gram-positive pathogens. Orally administered MRX-1 has shown the same or better efficacy than linezolid in systemic and local-infection mouse models.

Delamanid

WHO issued interim policy guidance on the use of delamanid for the treatment of adults with MDR-TB in 2014, based on Phase IIb trial results.² A conditional recommendation was made to use delamanid as part of longer MDR-TB treatment regimens for adults. The recommendation to use delamanid as part of longer MDR-TB treatment regimens was conditional upon proper patient selection, a regimen design following WHO recommendations, close monitoring of treatment, active TB drug safety monitoring and management, and informed consent according to local requirements. Following the release of results for children and adolescents treated for MDR-TB using delamanid in 2016, WHO's guidance on the use of delamanid in adults was expanded to include patients aged 6–17 years.³

In November 2017, final results from a Phase III trial assessing the safety and efficacy of delamanid as an addition to an optimized background regimen for adults with MDR-TB were reported to WHO by the manufacturer Otsuka Pharmaceutical, Japan. WHO conducted an expedited external expert review of the new data and issued a position statement in January 2018,⁴ which stated that the conditional guidance on delamanid remained valid, but that delamanid should only be added to a longer MDR-TB treatment regimen when the regimen cannot otherwise be composed according to WHO recommendations. In 2018, additional data from the Phase III trial were analysed by WHO alongside data from other studies of patients treated with delamanid-containing regimens as part of a major update to WHO guidance on the treatment of drug-resistant TB.⁵

As with bedaquiline, delamanid is being used in trials

¹ A prodrug is a derivative of drug molecules that undergo transformation into a pharmacologically active drug once inside the body. From: Rautio J, Kumpulainen H, Heimbach T, Oliyai R, Oh D, Jarvinen T et al. Prodrugs: design and clinical applications. *Nat Rev Drug Discov.* 2008;7(3):255–70 (<https://www.ncbi.nlm.nih.gov/pubmed/18219308>, accessed 9 July 2018).

² World Health Organization. The use of delamanid in the treatment of multidrug-resistant tuberculosis. Interim policy guidance (WHO/HTM/TB/2014.23). Geneva, WHO. 2014. (http://apps.who.int/iris/bitstream/10665/137334/1/WHO_HTM_TB_2014.23_eng.pdf, accessed 12 June 2018).

³ World Health Organization. The use of delamanid in the treatment of multidrug-resistant tuberculosis in children and adolescents: interim policy guidance (WHO/HTM/TB/2016.14). Geneva, WHO. 2016. (<http://apps.who.int/iris/bitstream/10665/250614/1/9789241549899-eng.pdf>; accessed 12 June 2018).

⁴ World Health Organization. WHO position statement on the use of delamanid for multidrug-resistant tuberculosis (WHO/CDS/TB/2018.1). Geneva: WHO; 2018 (<http://www.who.int/tb/publications/2018/WHOPositionStatementDelamanidUse.pdf>, accessed 21 June 2018).

⁵ Further details are provided in **Box 4.2** in **Chapter 4**.

of all-oral treatment regimens (**Section 8.2.3**). The use of delamanid in addition to an optimized background regimen to treat children less than 6 years old is also being investigated in other trials. Studies of its use in the prevention of drug-resistant TB among contacts of people with MDR-TB are planned.

Delpazolid (LCB01–0371)

Delpazolid is a new oxazolidinone developed by LegoChem BioSciences. It entered a Phase II trial in the Republic of Korea in 2017.

GSK-3036656

GSK-3036656 is a new oxaborole compound developed by GlaxoSmithKline. A Phase I trial started in March 2017.

Macozinone

Macozinone (formerly PBTZ169) is a benzothiazinone developed by Nearmedic Plus. A Phase I trial has been completed and a Phase II trial has been started in the Russian Federation. A Phase I study with a new formulation was started in 2018 in Switzerland.

OPC-167832

OPC-167832 is a carbostyryl developed by Otsuka that is bactericidal against both growing and intracellular bacilli. A single ascending dose study has been completed. A multiple ascending dose and early bactericidal activity (EBA) study of OPC-167832 alone and in combination with delamanid is planned in 2018.

Pretomanid

Pretomanid is a nitroimidazole being developed by the Global Alliance for TB Drug Development (the TB Alliance). It is currently being tested as part of combination regimens for the treatment of both drug-susceptible and drug-resistant TB, including XDR-TB (**Section 8.2.3**).

Q203

Q203 is an imidazopyridine that has been developed by Qurient (Republic of Korea). Single doses of various sizes have been tested in Phase I trials, and a Phase II trial is planned.

SQ109

SQ109 is a new, small-molecule drug discovered by scientists at Sequella Inc. (USA) and the US National Institutes of Health (NIH). A Phase IIb/III trial in which the drug was added to a standard regimen for MDR-TB has been completed in seven clinical centres in the Russian Federation, and positive results in terms of safety, efficacy and tolerability were reported in a press release in March 2017. A Phase II trial in the USA is in the planning stages.

Sutezolid

Sutezolid (PNU-100480) is an oxazolidinone and an analogue of linezolid. Results from an EBA study presented

in 2012 showed a significant reduction in colony-forming unit counts compared with the baseline level after 14 days of treatment. In January 2017, the Medicines Patent Pool announced that it had signed a licence with Johns Hopkins University to facilitate the clinical development of sutezolid in combination with other drugs. On World TB Day 2017, the TB Alliance and the Medicines Patent Pool announced a licensing agreement for the clinical development of sutezolid.

TBA-7371

TBA-7371 is an inhibitor of the enzyme DprE1, which is essential in the synthesis of components of mycobacterial cell walls. This inhibitor has shown to be active against strains of *M. tuberculosis* resistant to known TB drugs. The TB Alliance is implementing a Phase I study in the USA.

TBI-166

TBI-166 was identified through a lead optimization effort by the TB Alliance in partnership with the Institute of Materia Medica, the Chinese Academy of Medical Sciences and the Peking Union Medical College in Beijing. This rimonophenazine compound has improved physicochemical and pharmacokinetic properties (to avoid discoloration of skin), and has efficacy similar to clofazimine. A Phase I clinical trial started in January 2018 in China.

8.2.2 Approved drugs being tested for new purposes

Clofazimine

Clofazimine is a rimonophenazine that is used to treat leprosy. Its use in MDR-TB treatment is being explored in preclinical models of TB infection, to better understand its anti-TB effects. Novartis, the company that manufactures the drug, has withdrawn support for Phase II trials; however, clofazimine continues to be tested as part of treatment regimens for MDR-TB in Phase III trials.

Levofloxacin

Levofloxacin is being tested in a Phase II study called Opti-Q, which is investigating the best dose of levofloxacin to use for treatment of MDR-TB in adults with smear- and culture-positive pulmonary TB. Four doses are being tested (11, 14, 17 and 20 mg/kg/day) as part of an optimized background regimen. Trial enrolment and follow-up (in Peru and South Africa) has been completed. Data analysis is underway.

Linezolid

Linezolid is a marketed oxazolidinone with potent activity against TB. It has been widely used in the treatment of drug-resistant TB, and there is good evidence that it improves culture conversion and cure rates when added to treatment regimens. Since the medicine has a narrow therapeutic window, and the optimal dosing strategy

remains unknown, the TB Alliance has implemented a Phase II trial to evaluate the mycobactericidal activity, safety, tolerability and pharmacokinetics of five doses of linezolid in adults with pulmonary TB.

Moxifloxacin

Moxifloxacin is included in several trials of new regimens for treatment of both drug-susceptible and drug-resistant TB (**Section 8.2.3**).

Nitazoxanide

Nitazoxanide is an anti-parasitic drug. Its activity against *M. tuberculosis* is being tested in a Phase II trial in Haiti.

Rifampicin (high dose)

Findings from the multi-arm, multi-stage TB (MAMS-TB) trial of the Pan-African Consortium for the Evaluation of Antituberculosis Antibiotics (PanACEA) were published in 2017.¹ These showed that 35 mg/kg of rifampicin given over 12 weeks is safe, and shortens the time to stable culture conversion from 62 to 48 days. The other trial arms – which included various combinations of 10 mg/kg or 20 mg/kg of rifampicin, moxifloxacin and SQ109 – did not achieve significant improvements compared with the control arm. When all the data are taken into consideration, the trial suggests that a 35 mg/kg dose of rifampicin given for 12 weeks is likely to improve treatment outcomes. This trial is the first multi-arm adaptive trial design to be successfully implemented in multiple sites in countries with a high burden of TB. It may help to pave the way for accelerated testing of new TB treatment regimens at reduced cost.

Rifapentine

The effectiveness of rifapentine in the treatment of drug-susceptible TB is being studied in three trials. The TB Trial Consortium (TBTC) Study 31/ACTG A5349 is a Phase III trial that is investigating the use of rifapentine, with or without moxifloxacin, to shorten the treatment of drug-susceptible pulmonary TB to 4 months. TBTC Study 35, a Phase II study of the pharmacokinetics of new water-dispersible paediatric formulations of rifapentine, is scheduled to start at two sites in South Africa in 2018. TBTC Study 37 is a Phase III trial of the safety and efficacy of 6 weeks of daily rifapentine for the treatment of latent TB infection (LTBI) in HIV-negative people in settings with a low burden of TB. It is being compared with a locally available 3–4 month rifamycin-based regimen.

¹ Boeree MJ, Heinrich N, Aarnoutse R, Diacon AH, Dawson R, Rehal S et al. High-dose rifampicin, moxifloxacin, and SQ109 for treating tuberculosis: a multi-arm, multi-stage randomized controlled trial. *Lancet Infect Dis*. 2017;17(1):39–49 (<https://www.ncbi.nlm.nih.gov/pubmed/28100438>; accessed 12 June 2018).

8.2.3 New regimens for the treatment of drug-susceptible or drug-resistant TB

New combinations of drugs are being tested in Phase II or Phase III trials.

ACTG A5343 DELIBERATE

The ACTG A5343 DELIBERATE trial is testing the cardiotoxicity of regimens containing delamanid and bedaquiline alone, and in combination in pharmacokinetic and drug–drug interaction studies. The trial is sponsored by NIH and the National Institute of Allergy and Infectious Diseases (NIAID).

endTB

The endTB trial started in 2017. It is comparing several treatment regimens for MDR-TB or XDR-TB with the current longer treatment regimen for MDR-TB recommended by WHO. The regimens being tested include bedaquiline or delamanid (or both), moxifloxacin or levofloxacin, and pyrazinamide plus linezolid or clofazimine (or both), in various combinations.

MDR-END

The MDR-END trial is investigating a 9–12 month regimen of delamanid, linezolid, levofloxacin and pyrazinamide for the treatment of MDR-TB among TB patients without resistance to fluoroquinolones.

NeXT

The NeXT trial is testing a 6–9 month injection-free regimen of bedaquiline, ethionamide or high-dose isoniazid, linezolid, levofloxacin and pyrazinamide for patients with MDR-TB, compared with the 21–24 months treatment regimen. Recruitment started in South Africa in 2016.

NiX-TB, ZeNix and BPamZ

The NiX-TB trial is investigating the safety and efficacy of a 6-month all-oral regimen combining bedaquiline, pretomanid and linezolid in patients with XDR-TB, and in patients who could not tolerate MDR-TB treatment or for whom this treatment failed. It is being implemented by the TB Alliance in South Africa. The primary end-point is bacteriological failure, relapse or clinical failure during a 6 month follow-up period after completion of treatment. A cure rate of 87% has been reported for the first 15 patients. A follow-on trial called ZeNix is exploring lower doses and shorter durations of linezolid to minimize toxicity.

A Phase IIc/III trial (called SimpliciTB) of bedaquiline, pretomanid, moxifloxacin and pyrazinamide (BPamZ), targeting patients with both drug-susceptible or MDR-TB, is also being implemented. The primary end-point is culture conversion at 2 months. A previous Phase IIb study of this BPamZ regimen showed almost 100% culture conversion at 2 months in patients with MDR-TB.

STREAM

The STREAM Stage 1 trial is comparing a 9-month regimen for MDR-TB with longer regimens of 18–24 months in Ethiopia, Mongolia, South Africa and Viet Nam. It is a Phase III trial for which enrolment and follow-up were completed in 2018.¹

Following the release of preliminary results, WHO conducted an expedited review of data provided by the STREAM trial consortium.² The analysis showed that a “favourable outcome” was slightly higher in patients treated with longer regimens than in patients on the shorter regimen; non-inferiority as defined by the study protocol could not be shown. The external experts who reviewed the interim findings recommended that national TB programmes and other stakeholders should continue to use the shorter MDR-TB regimen under the same conditions as those recommended in WHO guidance issued in 2016.³ The final results from the STREAM Stage 1 trial were used to inform an update of WHO guidance on the treatment of MDR-TB in 2018.⁴

TB-PRACTECAL

The TB-PRACTECAL trial is a Phase II/III trial to evaluate the safety and efficacy of 6-month regimens that contain bedaquiline, pretomanid and linezolid, with or without moxifloxacin or clofazimine, for the treatment of adults with MDR-TB or XDR-TB. Primary outcomes include 8-week culture conversion, and the development of unfavourable outcomes (treatment failure or recurrence, death, discontinuation or loss to follow-up during a 72-week follow-up period). It is being implemented by Médecins sans Frontières and other collaborators in Belarus, South Africa and Uzbekistan.

8.3 New vaccines to prevent TB

The bacille Calmette–Guérin (BCG) vaccine, first used in the 1920s, remains the only licensed vaccine for preventing TB. Despite high coverage of BCG vaccination as part of childhood immunization programmes (Chapter 5), the slow decline in TB incidence globally highlights the need for a much more effective vaccine that provides protection against all forms of TB in all age groups.

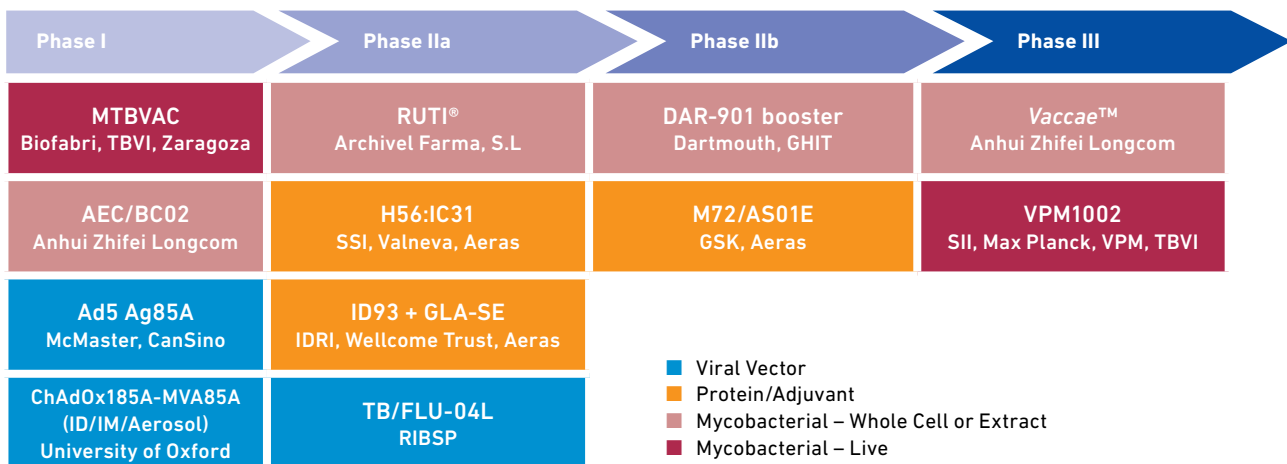
The status of the pipeline for new TB vaccines in August 2018, including the names of vaccine developers,

¹ Nunn AJ, Rusen ID, Van Deun A, Torrea G, Phillips PP, Chiang CY et al. Evaluation of a standardized treatment regimen of anti-tuberculosis drugs for patients with multi-drug-resistant tuberculosis (STREAM): study protocol for a randomized controlled trial. *Trials*. 2014;15:353 (<https://www.ncbi.nlm.nih.gov/pubmed/25199531>; accessed 12 June 2018).

² World Health Organization. WHO issues a position statement on the continued use of the shorter MDR-TB regimen following an expedited review of the STREAM Stage 1 preliminary results. Geneva: WHO; 2018 (http://www.who.int/tb/features_archive/WHO-position-statement-MDR-TB-regimen/en/; accessed 12 June 2018).

³ World Health Organization. WHO treatment guidelines for drug-resistant tuberculosis, 2016 update. Geneva: WHO; 2016 (<http://www.who.int/tb/areas-of-work/drug-resistant-tb/treatment/resources/en/>, accessed 3 July 2018).

⁴ Further details are provided in Box 4.2 in Chapter 4.

FIG. 8.3**The global development pipeline for new TB vaccines, August 2018^a**

^a Information was self-reported by vaccine sponsors, and the Stop TB Partnership Working Group on New TB Vaccines supported the review of their feedback.

is shown in **Fig. 8.3**. There are 12 vaccines in Phase I, II or III trials. Their main characteristics are summarized below.

8.3.1 Phase I trials

There are four vaccine candidates in Phase I trials.

Ad5 Ag85A

Ad5 Ag85A is an adenovirus serotype 5 vector expressing Ag85A. It has been evaluated for safety and immunogenicity in both BCG-naïve and previously BCG-immunized healthy volunteers in Canada. Overall, intramuscular administration was found to be safe, well tolerated and immunogenic in both trial groups, with more potent immunogenicity observed in volunteers who had been previously vaccinated with BCG. A safety and immunogenicity study of aerosol administration in BCG-vaccinated healthy volunteers has started.

AEC/BC02

AEC/BC02 is a freeze-dried recombinant vaccine expressing Ag85B and fusion protein ESAT6-CFP10. A Phase I study assessing safety and immunogenicity is underway in China, with sponsorship from Anhui Zhifei Longcom Biologic Pharmacy Co., Ltd.

ChAdOx185A – MVA85A (ID/IM/Aerosol)

ChAdOx185A is a simian adenovirus and MVA85A is a recombinant pox virus – both express antigen 85A. These candidates are being developed with the overall aim of generating a joint heterologous prime-boost regimen delivered through both systemic and mucosal routes.

A Phase I trial of intramuscular administration of ChAdOx185A in BCG-vaccinated adults in the United Kingdom, both alone and as part of a prime-boost strategy with MVA85A, has been completed. Phase I trials of

aerosol administration of ChAdOx185A in BCG-vaccinated adults are scheduled to start in the third quarter of 2018. Two studies of aerosol administration of MVA85A in BCG-vaccinated individuals have been completed, and a further study in people with LTBI is being implemented.

MTBVAC

MTBVAC is a live strain of *M. tuberculosis*, attenuated via deletions of the *phoP* and *fadD26* genes. The primary target population is neonates (as a BCG replacement vaccine); the secondary target population is adolescents and adults (as a booster vaccine). A Phase Ib trial in neonates was completed in 2018. Phase IIa trials in both target populations will start in 2018.

8.3.2 Phase II and Phase III clinical trials

There are eight vaccines in Phase II or Phase III trials. Another vaccine (H4:IC31) was in development until recently, and for this reason it is also described below.

DAR-901 booster

DAR-901 is a whole-cell, heat-inactivated, nontuberculous mycobacterial vaccine booster. It represents a new scalable manufacturing method for SRL172, a candidate vaccine that showed efficacy among adults living with HIV in a Phase II/III trial in the United Republic of Tanzania. It is now being tested in a Phase IIb prevention of (established) infection trial among BCG-primed adolescents, also in the United Republic of Tanzania. The trial is scheduled for completion in 2019.

H4:IC31

H4:IC31 is a subunit vaccine that contains a fusion protein of Ag85B and TB10.4, formulated with IC31 adjuvant. A pre-proof-of-concept Phase II trial to evaluate prevention of (established) infection among IGRA-negative,

HIV-negative adolescents who had undergone neonatal BCG vaccination was recently completed in South Africa.¹ The results showed that H4:IC31 induced increased immunogenicity specific to *M. tuberculosis* compared with placebo. However, statistical significance in preventing initial or sustained infection (measured using the QuantiFERON-TB Gold In-Tube test) was not shown. In the same study, a BCG revaccination arm showed a statistically significant reduction in the rate of sustained conversion compared with the placebo arm. Due to inadequate efficacy, further development and testing of the H4:IC31 candidate has been terminated. For this reason it is not shown in **Fig. 8.3**.

H56:IC31

H56:IC31 is an adjuvanted subunit vaccine that combines three *M. tuberculosis* antigens (Ag85B, ESAT-6 and Rv2660c) with the IC31[®] adjuvant, from Valneva Austria GmbH (Vienna Austria). Three Phase I or I/IIa trials of safety and immunogenicity have been completed. Two of these were in HIV-negative, BCG-vaccinated adults with and without LTBI, and without a history or any evidence of TB disease. The other was in HIV-negative pulmonary TB patients who had recently completed treatment. The trials showed that the vaccine had an acceptable safety profile and was immunogenic at all studied doses. Analysis of a Phase Ib trial evaluating the safety and immunogenicity of H4:IC31, H56:IC31 and BCG revaccination in adolescents is underway. In 2018, a Phase 2b trial assessing H56:IC31 for prevention of recurrence will be initiated in the United Republic of Tanzania and South Africa, co-sponsored by the Statens Serum Institut (SSI) and Aeras, with support from the European and Developing Countries Clinical Trial Partnership (EDCTP).

ID93 + GLA-SE

The ID93 + GLA-SE vaccine comprises four *M. tuberculosis* antigens associated with virulence (Rv2608, Rv3619 and Rv3620) or latency (Rv1813), and the adjuvant GLA-SE. A Phase IIa trial in HIV-negative TB patients that have recently completed treatment for pulmonary TB disease has been completed in South Africa, in preparation for a Phase IIb trial that will investigate the prevention of recurrence of disease in the same population. A Phase IIa trial in BCG-vaccinated healthy health-care workers, to assess prevention of infection, is underway.

M72/AS01E

M72/AS01E is a subunit vaccine that pairs two *M. tuberculosis* antigens (32A and 39A) with an adjuvant (AS01E). It is being tested in a Phase IIb efficacy trial in HIV-negative adults infected with *M. tuberculosis* in Kenya, South Africa and Zambia. The primary end-point is the

number of incident cases of active pulmonary TB disease not associated with HIV infection. Secondary end-points include safety and immunogenicity.

RUTI[®]

RUTI[®] is a non-live, polyantigenic vaccine based on cell-wall fragmented *M. tuberculosis* bacteria. It is intended as a therapeutic vaccine, to be used in conjunction with a short, intensive antibiotic treatment. A Phase I study in healthy volunteers and a Phase II study in people with LTBI have demonstrated a good safety profile, and found the vaccine to be immunogenic at all studied doses. The main target for RUTI is MDR-TB, and a Phase IIa study in patients with MDR-TB is being implemented.

TB/FLU-04L

TB/FLU-04L is a mucosal-vectored vaccine based on an attenuated replication-deficient influenza virus vector expressing antigens Ag85A and ESAT-6. It was designed as a prophylactic boost vaccine for infants, adolescents and adults. A Phase IIa trial in people with LTBI is being implemented.

Vaccae[™]

Vaccae[™] vaccine is a specified lysate that has been licensed by the China Food and Drug Administration as an immunotherapeutic agent to help shorten TB treatment for patients with drug-susceptible TB. A Phase III trial to assess its efficacy and safety in preventing TB disease in people with LTBI has been completed, and data analysis is underway. It is the largest TB vaccine trial undertaken in the past decade, involving 10 000 people aged 15–65 years.

VPM1002

VPM1002 is a live recombinant vaccine. A Phase II trial is being implemented in South Africa to assess the safety and immunogenicity of the vaccine in HIV-exposed and unexposed neonates, and the preparations for a subsequent Phase III trial are underway. A Phase II/III trial for prevention of TB recurrence in adults is being implemented in India.

8.4 Research collaborations and partnerships to end TB

Stakeholders with a clear role to play in advancing TB research include national and local governments, private companies, academia, civil society and TB-affected communities, research institutions, universities, and funding agencies. The Global Action Framework for TB Research, published by WHO in 2015, was developed to facilitate research collaborations and partnerships.² It has two main components: promoting TB research at a

¹ Nemes et al. 2018. Prevention of *M. tuberculosis* Infection with H4:IC31 Vaccine or BCG Revaccination. *N Engl J Med.* 379:138-149 (<https://www.ncbi.nlm.nih.gov/pubmed/29996082>, accessed 18 July 2018).

² World Health Organization. A global action framework for TB research in support of the third pillar of WHO's End TB Strategy. Geneva: WHO; 2015 (<http://www.who.int/tb/publications/global-framework-research/en/>, accessed 8 August 2017).

BOX 8.3

The BRICS TB Research Network

In November 2017, the governments of Brazil, the Russian Federation, India, China and South Africa announced the establishment of a collaborative TB research network, the aim of which is to accelerate TB research and innovation through cooperation mechanisms.

The establishment of the network follows previous BRICS commitments to coordinated action related to the TB response. In the 2013 Delhi Communiqué, BRICS agreed to collaborate to develop “capacity and infrastructure to reduce the prevalence and incidence of TB through innovation for new drugs, vaccines, diagnostics and promotion of consortia of TB researchers to collaborate on clinical trials of drugs and vaccines”.^a When BRICS ministers of health met in Brasilia in 2014, they agreed to cooperate on TB research and innovation, and they identified technology sharing, manufacturing capacity and TB financing as key priorities.^b In December 2016, they agreed to “the setting up of a BRICS network on TB research and creation of a research and development consortium on TB, HIV and malaria, including the possibility of international fund raising”.^c

The new network will include cooperation on TB research and innovation related to product development and validation, technology transfer, delivery of TB care and services, and capacity-building, based on the use of existing expertise and technological capacity within BRICS. Priority activities include leveraging domestic funds to support the development of common protocols and associated implementation of multicountry studies to accelerate the development of TB diagnostics, drugs and vaccines; and evaluating the epidemiological, clinical and economic impact of adopting innovative interventions at national level.

^a Government of India Ministry of Health and Family Welfare. Delhi communiqué. Delhi: Government of India; 2013 (<http://pib.nic.in/newsite/erelease.aspx?relid=91533>, accessed 11 June 2018).

^b BRICS Ministry of External Relations. Communiqué of the IV meeting of BRICS health ministers. Brasilia: BRICS Ministry of External Relations; 2014 (<http://brics.utoronto.ca/docs/141205-health.html>, accessed 11 June 2018).

^c Ministry of Health and Family Welfare, Government of India. 6th BRICS Health Ministers Meeting Delhi Communiqué. Delhi: Government of India Ministry of Health and Family Welfare; 2016 (<http://pib.nic.in/newsite/PrintRelease.aspx?relid=155596>, accessed 11 June 2018).

country level to find ways to address local barriers; and catalysing research at a global level through advocacy, knowledge sharing, research prioritization, collaboration and support to international networks for research and capacity-building.

One component of promoting research at the global level is the establishment of multicountry research networks. These can help to expedite innovation by fostering collaboration, leveraging resources and facilitating technology transfer, through public–private partnerships, north–south and south–south cooperation. Governments have an important role to play in strengthening such initiatives by creating policy frameworks that enable and promote research partnerships. At a G20 summit¹ in 2017, heads of state called for the establishment of a collaborative hub for research and development “to maximize the impact of new and existing anti-microbial basic and clinical research initiatives as well as product development” for drug-resistant infections, including TB.

A recent analysis of the authorship of research publications on TB between 2007 and 2016 suggested that north–south collaborations increased steadily during this period, while south–south collaborations were much less common.² With growing scientific and economic capacity in low- and middle-income countries, the capacity of countries with a high burden of TB to contribute to research is expected to increase. This is particularly so for the BRICS countries (Brazil, Russian Federation, India, China and South Africa), which account for more than 40% of the global TB disease burden in terms of both TB incidence and TB deaths, and about 50% of the global burden of drug-resistant TB (Chapter 3). During the Global Ministerial Conference on TB in November 2017, the establishment of a BRICS TB research network was announced; further details are provided in Box 8.3.

¹ G20 Nations. G20 Leaders’ Declaration: Shaping an interconnected world. Hamburg: G20 Nations; 2017 (<http://www.g20.utoronto.ca/2017/2017-G20-leaders-declaration.html>, accessed 11 June 2018).

² World Health Organization. Global investments in tuberculosis research and development past, present, and future. Geneva: WHO; 2017 (<http://apps.who.int/iris/bitstream/handle/10665/259412/9789241513326-eng.pdf>, accessed 28 June 2018).



An MDR-TB patient in Karachi, Pakistan

Hamad Darwish / WHO

ANNEX 1

The WHO global
TB database

A.1 Database contents

The 2018 global TB report is based on data collected annually from 216 countries and territories, including all 194 WHO Member States. These data are stored in a global TB database that is managed by the TB monitoring and evaluation unit of the Global TB Programme, at WHO headquarters.

In 2018, data were collected on the following topics: TB case notifications and treatment outcomes, including breakdowns by TB case type, age, sex, HIV status and drug resistance; laboratory diagnostic services; monitoring and evaluation, including surveillance and surveys specifically related to drug-resistant TB; TB preventive therapy; TB infection control; palliative care; engagement of all public and private care providers in TB prevention and care; community engagement; the budgets of national TB control programmes (NTPs); utilization of general health services (hospitalization and outpatient visits) during treatment; and NTP expenditures. A shortened version of the online questionnaire was used for high-income countries (that is, countries with a gross national income per capita of \geq US\$ 12 236 in 2016, as defined by the World Bank)¹ and/or low-incidence countries (defined as countries with an incidence rate of <20 cases per 100 000 population or <10 cases in total in 2016).

Countries reported data using a dedicated website (<https://extranet.who.int/tme>), which was opened for reporting in April 2018. Switzerland and countries in the European Union submitted data on notifications and treatment outcomes to the TESSy system managed by the European Centre for Disease Prevention and Control (ECDC). Data from TESSy were uploaded into the global TB database.

Additional data about the provision of treatment for latent TB infection to people newly enrolled in HIV care and antiretroviral therapy for HIV-positive TB patients were collected by the Joint United Nations Programme on HIV/AIDS (UNAIDS). These data were jointly validated by UNAIDS and the WHO's Global TB Programme and HIV department, and uploaded into the global TB database.

Following review and follow-up with countries, the data used for the main part of this report were those data available on **6 August 2018**. **Table A1.1** shows the number of countries and territories that had reported data by 6 August 2018.

TABLE A1.1

Reporting of data in the 2018 round of global TB data collection

	COUNTRIES AND TERRITORIES		WHO MEMBER STATES	
	NUMBER	NUMBER THAT REPORTED DATA	NUMBER	NUMBER THAT REPORTED DATA
African Region	47	46	47	46
Region of the Americas	46	40	35	34
Eastern Mediterranean Region	22	21	21	21
European Region	54	45	53	44
South-East Asia Region	11	11	11	11
Western Pacific Region	36	35	27	27
Global	216	198	194	183

Indicators in the Sustainable Development Goals associated with TB incidence were imported into the global TB database on **30 July 2018**. **Table A1.2** shows the data sources used.

A.2 Accessing TB data using the WHO Global TB Programme website

Most of the data held in the global TB database can be found by going to www.who.int/tb/data. This web page provides access to country profiles, comma-separated value (CSV) data files and data visualisations.

A2.1 Country profiles

Profiles can be viewed and downloaded for all 216 countries and territories that report TB data to WHO each year, and not just the 30 high burden countries shown in the printed version of the global TB report. The profiles can be generated on-demand directly from the global TB database and therefore may include updates received after publication of the global TB report.

TB financial profiles can be viewed and downloaded for over 100 countries and territories that report detailed TB financial data to WHO.

¹ <http://data.worldbank.org/about/country-classifications>

TABLE A1.2
Data sources for indicators in the Sustainable Development Goals associated with TB incidence

SDG INDICATOR	DISPLAY NAME IN PROFILE	DATA SOURCE	NAME AT SOURCE	SOURCE URL
1.1.1	Population living below the international poverty line (% of population)	UN SDG database	Proportion of population below the international poverty line	https://unstats.un.org/sdgs/indicators/database/?indicator=1.1.1
1.3.1	Population covered by social protection floors/systems (% of population)	World Bank	Coverage – All Social Protection and Labour (% of population)	http://data.worldbank.org/indicator/per_allsp.cov_pop_tot
2.1.1	Prevalence of undernourishment (% of population)	World Bank	Prevalence of undernourishment (% of population)	http://data.worldbank.org/indicator/SN.ITK.DEFC.ZS
3.3.1 (alternative)	HIV prevalence (% of population aged 15–49 years)	World Bank	Prevalence of HIV, total (% of population aged 15–49)	http://data.worldbank.org/indicator/SH.DYN.AIDS.ZS
3.4.1 (alternative)	Diabetes prevalence (% of population aged ≥18 years)	WHO-GHO	Raised fasting blood glucose (≥ 7.0 mmol/L or on medication) (age-standardized estimate)	http://apps.who.int/gho/data/node.main.A869 Direct links to CSV files: http://apps.who.int/gho/athena/data/data-coded.csv?target=GHO/NCD_GLUC_04&filter=AGEGROUP:*;COUNTRY:*;SEX:FMLE and http://apps.who.int/gho/athena/data/data-coded.csv?target=GHO/NCD_GLUC_04&filter=AGEGROUP:*;COUNTRY:*;SEX:MLE
3.5.2 (alternative)	Alcohol use disorders, 12 month prevalence (% in population aged ≥15 years)	WHO-GHO	Alcohol use disorders (15+), 12 month prevalence (%)	http://apps.who.int/gho/data/view.main.53040 Direct links to CSV files: http://apps.who.int/gho/athena/data/data-coded.csv?target=GHO/SA_0000001462&filter=COUNTRY:*;SEX:FMLE and http://apps.who.int/gho/athena/data/data-coded.csv?target=GHO/SA_0000001462&filter=COUNTRY:*;SEX:MLE
3.a.1 (alternative)	Smoking prevalence (% aged ≥15 years)	World Bank	Smoking prevalence, females (% of adults) and Smoking prevalence, males (% of adults)	http://data.worldbank.org/indicator/SH.PRV.SMOK.FE and http://data.worldbank.org/indicator/SH.PRV.SMOK.MA
3.8.2 (alternative)	Health expenditure per capita, PPP (current international \$)	World Bank	Health expenditure per capita, PPP (current international \$)	http://data.worldbank.org/indicator/SH.XPD.CHEX.PP.CD
3.8.2 (alternative)	Out-of-pocket health expenditure (% of current health expenditure)	World Bank	Out-of-pocket health expenditure (% of current health expenditure)	http://data.worldbank.org/indicator/SH.XPD.OOPC.CH.ZS
3.8.1	UHC Index of essential service coverage (% based on 16 tracer indicators including TB treatment)	WHO-GHO	UHC Index of essential service coverage (%)	http://apps.who.int/gho/data/node.main.INDEXOFESSENTIALSERVICECOVERAGE Direct links to CSV file: http://apps.who.int/gho/athena/data/data-coded.csv?target=GHO/UHC_INDEX_REPORTED&filter=COUNTRY:*
7.1.2	Access to clean fuels and technologies for cooking (% of population)	World Bank	Access to clean fuels and technologies for cooking (% of population)	http://data.worldbank.org/indicator/EG.CFT.ACCS.ZS
8.1.1 (alternative)	GDP per capita, PPP (constant 2011 international \$)	World Bank	GDP per capita, PPP (constant 2011 international \$)	http://data.worldbank.org/indicator/NY.GDP.PCAP.PP.KD
10.1.1 (alternative)	GINI index (0=perfect equality, 100=perfect inequality)	World Bank	GINI index (World Bank estimate)	http://data.worldbank.org/indicator/SI.POV.GINI
11.1.1	Population living in slums (% of urban population)	World Bank	Population living in slums (% of urban population)	http://data.worldbank.org/indicator/EN.POP.SLUM.UR.ZS

A2.2 CSV data files

These files are the primary resource for anyone interested in conducting their own analyses of the records in the global TB database. Data reported by countries, such as time series for case notifications and treatment outcomes and WHO's estimates of TB disease burden, can be downloaded as comma-separated value (CSV) files covering all years for which data are available. These CSV files can be imported into many spreadsheet, statistical analysis and database packages.

A data dictionary that defines each of the variables available in the CSV files is also available and can be downloaded.

The CSV files are generated on-demand directly from the global TB database, and therefore may include updates received after publication of the global TB report.

A2.3 Data visualisations

There are several interactive web pages that can be used to view maps, graphs and underlying data on TB case notifications, drug-resistant TB cases, treatment outcomes and WHO estimates of TB incidence and mortality.

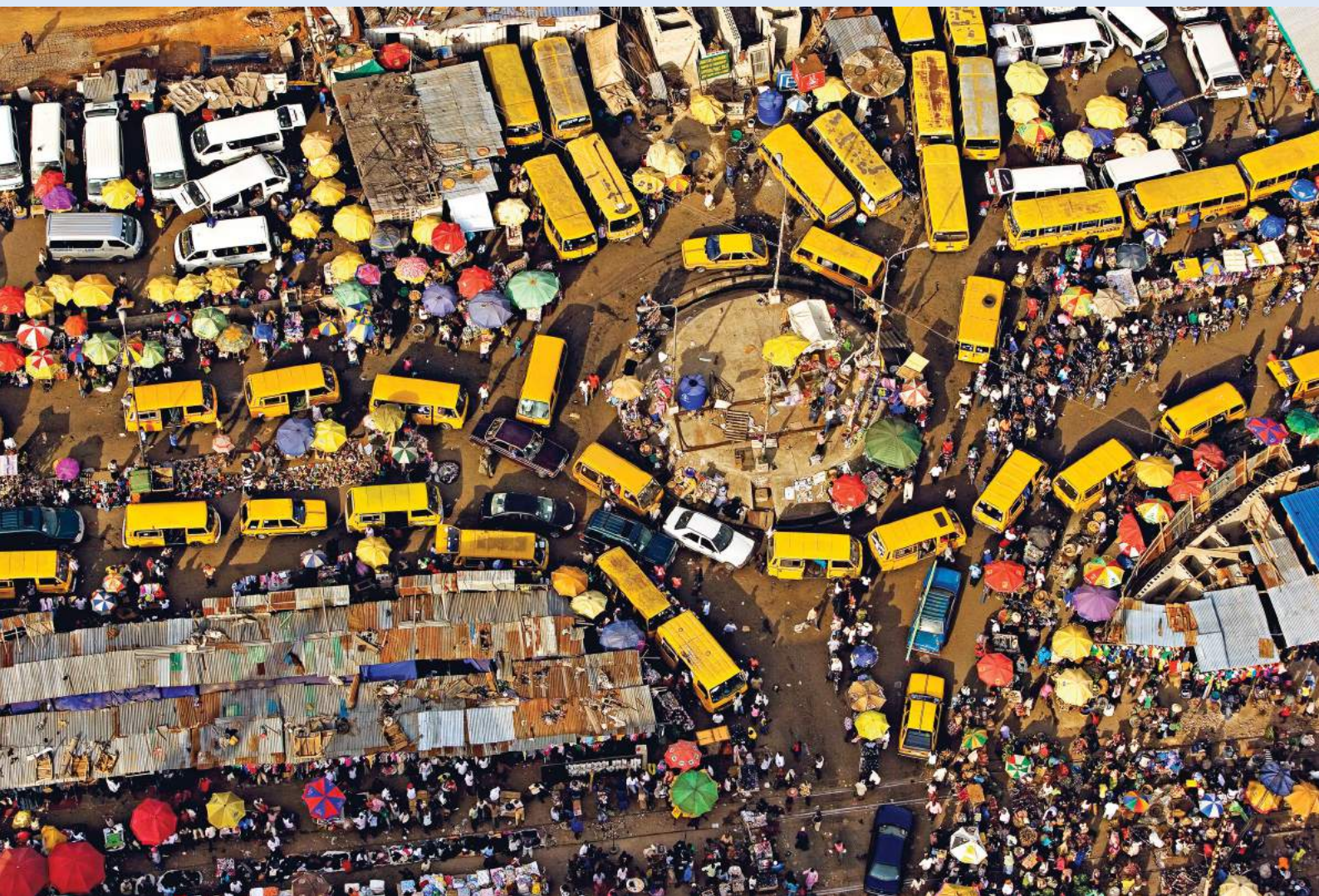
A.3 Accessing TB data using the WHO Global Health Observatory

The WHO Global Health Observatory (GHO) at www.who.int/gho/ is WHO's portal, providing access to data and analyses for monitoring the global health situation. It includes a data repository.

Data from WHO's global TB database can be viewed, filtered, aggregated and downloaded from within the GHO Data Repository at <http://apps.who.int/gho/data/node.main.1315>

The GHO data table headers include links to variable and indicator definitions. The data can be downloaded in many formats, including as CSV and Excel files.

There is also an Application Programme Interface (API) for analysts and programmers to use GHO data directly in their software applications. See <http://apps.who.int/gho/data/node.resources>



A busy market near Surulere in Lagos, Nigeria

Yann Arthus-Bertrand / Getty images

ANNEX 2

Country profiles

FOR 30 HIGH TB BURDEN COUNTRIES

**20 high TB burden countries
based on absolute number
of incident cases**

**10 high TB burden countries
based on severity of
disease burden
(incidence per capita)**

Angola

POPULATION 2017 **30 MILLION**

ESTIMATES OF TB BURDEN, ^a 2017

	NUMBER (THOUSANDS)	RATE (PER 100 000 POPULATION)
Mortality (excludes HIV+TB)	20 (12–31)	67 (39–103)
Mortality (HIV+TB only)	7.8 (3.9–13)	26 (13–44)
Incidence (includes HIV+TB)	107 (69–153)	359 (232–512)
Incidence (HIV+TB only)	18 (9.1–30)	61 (31–102)
Incidence (MDR/RR-TB) ^b	3.9 (1.6–7.2)	13 (5.5–24)

ESTIMATED TB INCIDENCE BY AGE AND SEX (THOUSANDS), ^a 2017

	0–14 YEARS	> 14 YEARS	TOTAL
Females	5.2 (4.7–5.6)	35 (27–43)	40 (31–50)
Males	5.7 (5.2–6.3)	61 (43–79)	67 (46–87)
Total	11 (9.5–12)	96 (60–131)	107 (69–153)

TB CASE NOTIFICATIONS, 2017

Total cases notified	57 877
Total new and relapse	54 401
— % tested with rapid diagnostics at time of diagnosis	
— % with known HIV status	59%
— % pulmonary	94%
— % bacteriologically confirmed among pulmonary	53%

UNIVERSAL HEALTH COVERAGE AND SOCIAL PROTECTION

TB treatment coverage (notified/estimated incidence), 2017	51% (36–79)
TB patients facing catastrophic total costs	
TB case fatality ratio (estimated mortality/estimated incidence), 2017	0.27 (0.14–0.42)

TB/HIV CARE IN NEW AND RELAPSE TB PATIENTS, 2017

	NUMBER	(%)
Patients with known HIV-status who are HIV-positive	3 613	11%
— on antiretroviral therapy	1 401	39%

DRUG-RESISTANT TB CARE, 2017

	NEW CASES	PREVIOUSLY TREATED CASES	TOTAL NUMBER ^c
Estimated MDR/RR-TB cases among notified pulmonary TB cases			2 000 (1 200–2 900)
Estimated % of TB cases with MDR/RR-TB	2.5% (1.1–4.3)	14% (10–18)	
% notified tested for rifampicin resistance	<1%	6%	534
MDR/RR-TB cases tested for resistance to second-line drugs			0
Laboratory-confirmed cases		MDR/RR-TB: 534, XDR-TB: 0	
Patients started on treatment ^d		MDR/RR-TB: 534, XDR-TB: 0	

TREATMENT SUCCESS RATE AND COHORT SIZE

	SUCCESS	COHORT
New cases registered in 2016	27%	53 980
Previously treated cases registered in 2016	28%	7 069
HIV-positive TB cases registered in 2016	0%	2 773
MDR/RR-TB cases started on second-line treatment in 2015	72%	227
XDR-TB cases started on second-line treatment in 2015		0

TB PREVENTIVE TREATMENT, 2017

% of HIV-positive people (newly enrolled in care) on preventive treatment	13%
% of children (aged < 5) household contacts of bacteriologically-confirmed TB cases on preventive treatment	

TB FINANCING, 2018

National TB budget (US\$ millions)	46
Funding source:	7% domestic, 12% international, 80% unfunded

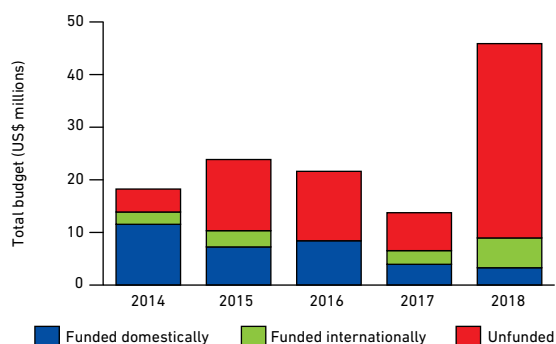
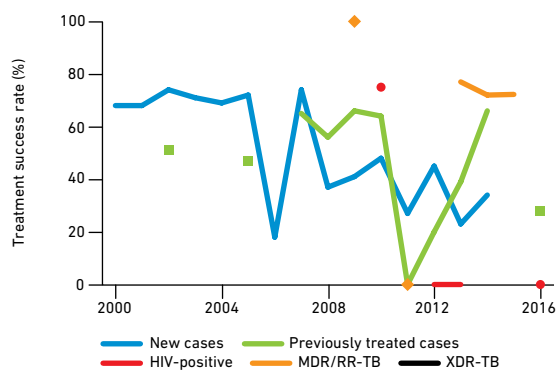
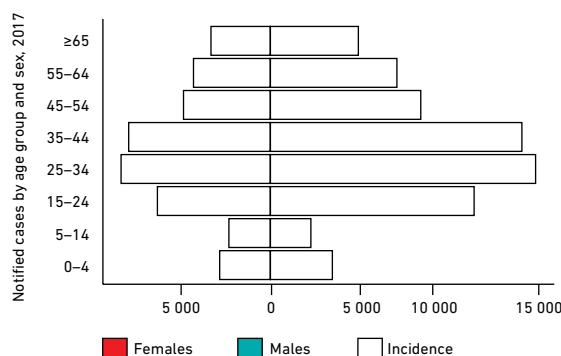
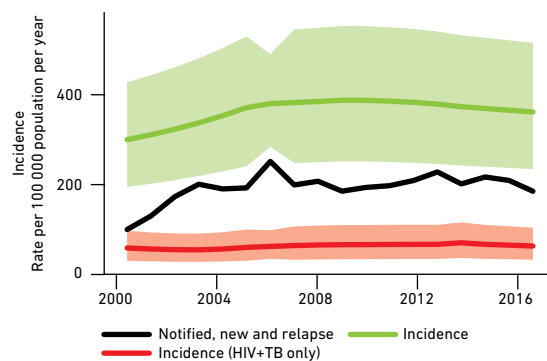
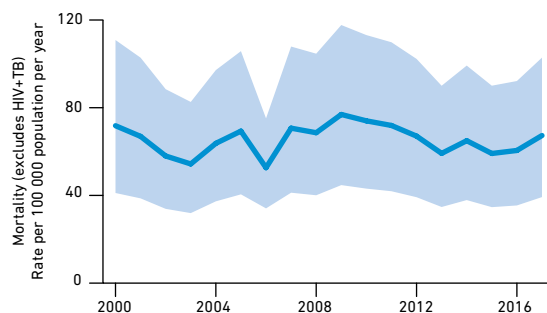
Data are as reported to WHO. Estimates of TB and MDR/RR-TB burden are produced by WHO in consultation with countries. Estimates are rounded and totals are computed prior to rounding.

^a Ranges represent uncertainty intervals.

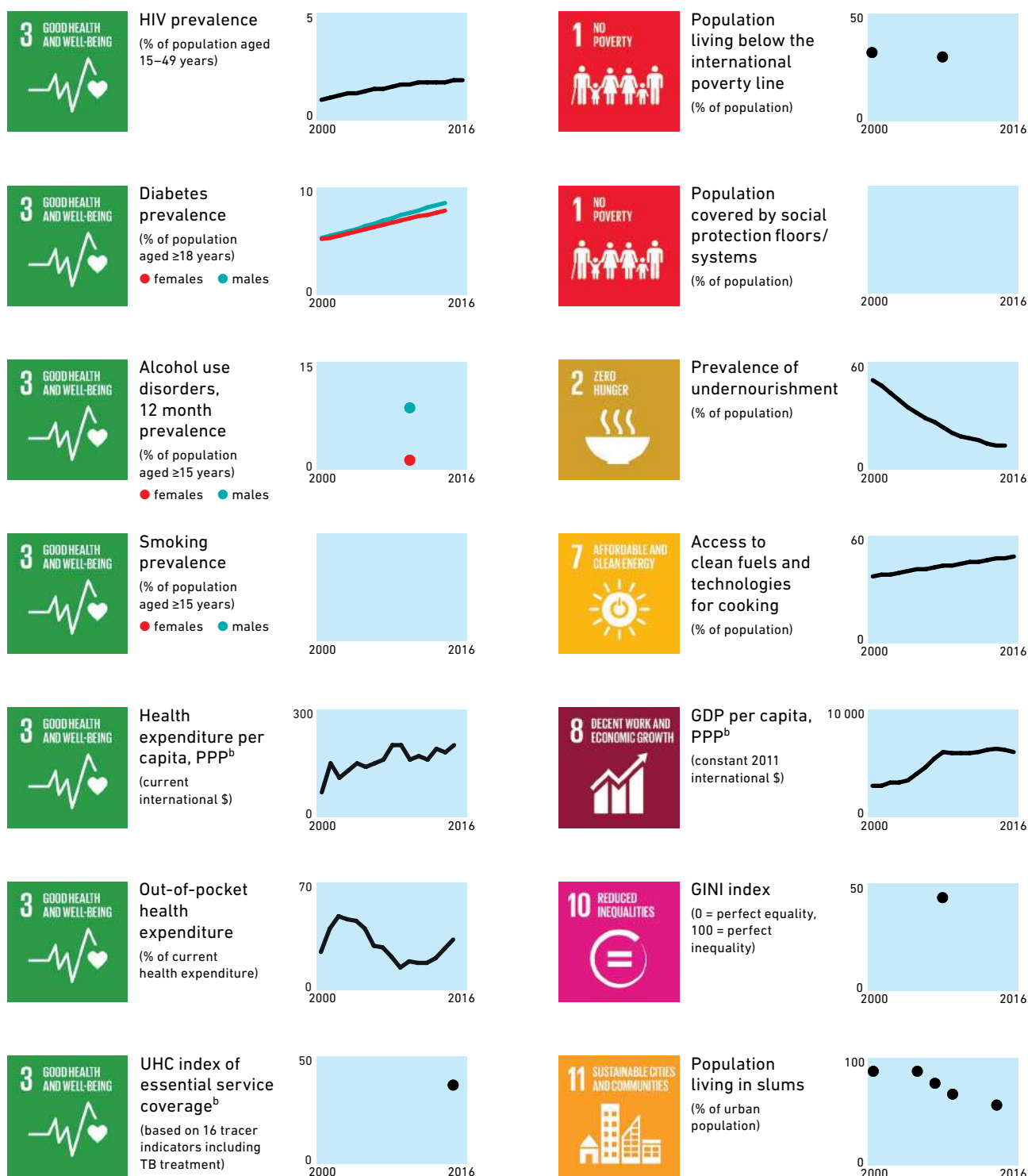
^b MDR is TB resistant to rifampicin and isoniazid; RR is TB resistant to rifampicin.

^c Includes cases with unknown previous TB treatment history.

^d Includes patients diagnosed before 2017 and patients who were not laboratory-confirmed.



INDICATORS IN THE SUSTAINABLE DEVELOPMENT GOALS ASSOCIATED WITH TB INCIDENCE^a



Targets for reductions in TB incidence and TB deaths set in WHO's End TB Strategy and the United Nations' Sustainable Development Goals (SDGs) are ambitious. Achieving them requires progress in reducing health-related risk factors for TB infection and disease, as well as broader social and economic determinants of TB infection and disease. WHO has developed a TB-SDG monitoring framework that comprises 14 indicators under seven SDGs for which there is evidence of an association with TB incidence. Further details are provided in [Chapter 2](#).

^a Data sources: SDG indicators database, The World Bank, World Health Organization. Missing values and empty boxes indicate data not available in these data sources.

^b GDP = gross domestic product; PPP = purchasing power parity; UHC = universal health coverage

Bangladesh

POPULATION 2017 **165 MILLION**

ESTIMATES OF TB BURDEN, 2017

	NUMBER (THOUSANDS)	RATE (PER 100 000 POPULATION)
Mortality (excludes HIV+TB)	59 (38–85)	36 (23–52)
Mortality (HIV+TB only)	0.17 (0.085–0.29)	0.11 (0.05–0.18)
Incidence (includes HIV+TB)	364 (265–479)	221 (161–291)
Incidence (HIV+TB only)	0.55 (0.27–0.92)	0.33 (0.17–0.56)
Incidence (MDR/RR-TB) ^b	8.4 (3.8–15)	5.1 (2.3–9)

ESTIMATED TB INCIDENCE BY AGE AND SEX (THOUSANDS)*, 2017

	0–14 YEARS	> 14 YEARS	TOTAL
Females	17 (16–18)	118 (98–137)	134 (110–158)
Males	18 (17–19)	212 (164–259)	230 (176–284)
Total	35 (32–38)	329 (237–421)	364 (265–479)

TB CASE NOTIFICATIONS, 2017

Total cases notified	244 201
Total new and relapse	242 639
— % tested with rapid diagnostics at time of diagnosis	<1%
— % with known HIV status	2%
— % pulmonary	81%
— % bacteriologically confirmed among pulmonary	74%

UNIVERSAL HEALTH COVERAGE AND SOCIAL PROTECTION

TB treatment coverage (notified/estimated incidence), 2017	67% (51–92)
TB patients facing catastrophic total costs	
TB case fatality ratio (estimated mortality/estimated incidence), 2017	0.17 (0.1–0.26)

TB/HIV CARE IN NEW AND RELAPSE TB PATIENTS, 2017

	NUMBER	(%)
Patients with known HIV-status who are HIV-positive	89	2%
— on antiretroviral therapy	84	94%

DRUG-RESISTANT TB CARE, 2017

	NEW CASES	PREVIOUSLY TREATED CASES	TOTAL NUMBER ^c
Estimated MDR/RR-TB cases among notified pulmonary TB cases			5 800 (3 800–7 800)
Estimated % of TB cases with MDR/RR-TB	1.6% (0.74–2.8)	29% (24–35)	
% notified tested for rifampicin resistance	18%	63%	49 943
MDR/RR-TB cases tested for resistance to second-line drugs			362
Laboratory-confirmed cases		MDR/RR-TB: 944, XDR-TB: 6	
Patients started on treatment ^d		MDR/RR-TB: 920, XDR-TB: 6	

TREATMENT SUCCESS RATE AND COHORT SIZE

	SUCCESS	COHORT
New and relapse cases registered in 2016	94%	222 252
Previously treated cases, excluding relapse, registered in 2016	86%	1 669
HIV-positive TB cases registered in 2016	49%	87
MDR/RR-TB cases started on second-line treatment in 2015	78%	880
XDR-TB cases started on second-line treatment in 2015		0

TB PREVENTIVE TREATMENT, 2017

% of HIV-positive people (newly enrolled in care) on preventive treatment	
% of children (aged < 5) household contacts of bacteriologically-confirmed TB cases on preventive treatment	21% (19–23)

TB FINANCING, 2018

National TB budget (US\$ millions)	66
Funding source:	14% domestic, 55% international, 31% unfunded

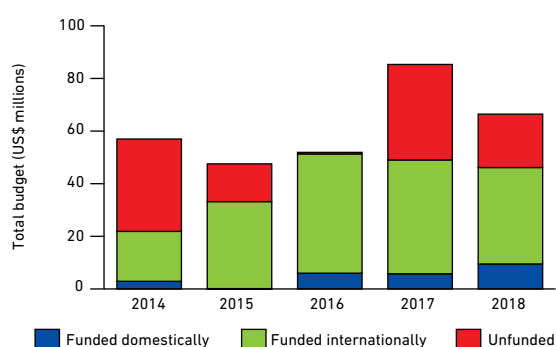
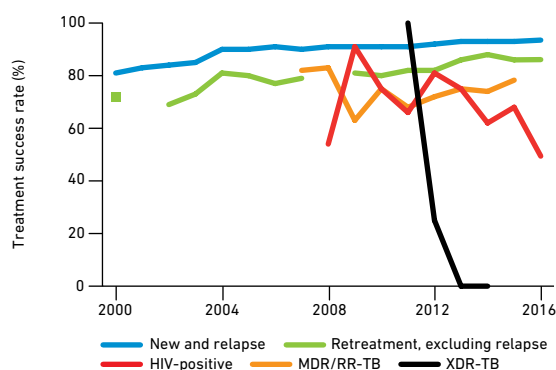
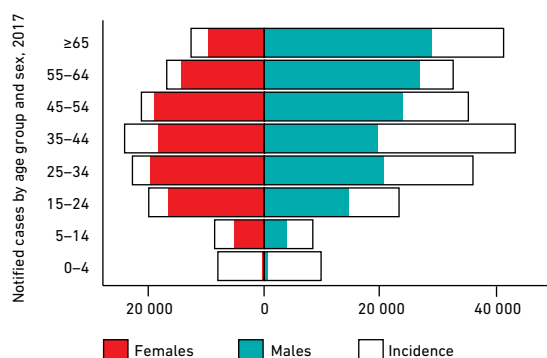
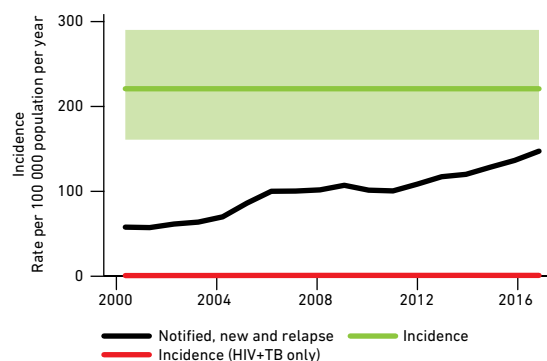
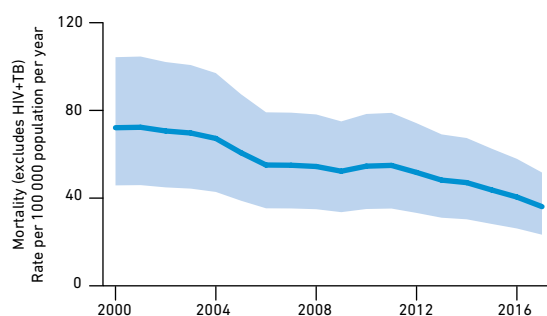
Data are as reported to WHO. Estimates of TB and MDR/RR-TB burden are produced by WHO in consultation with countries. Estimates are rounded and totals are computed prior to rounding.

^a Ranges represent uncertainty intervals.

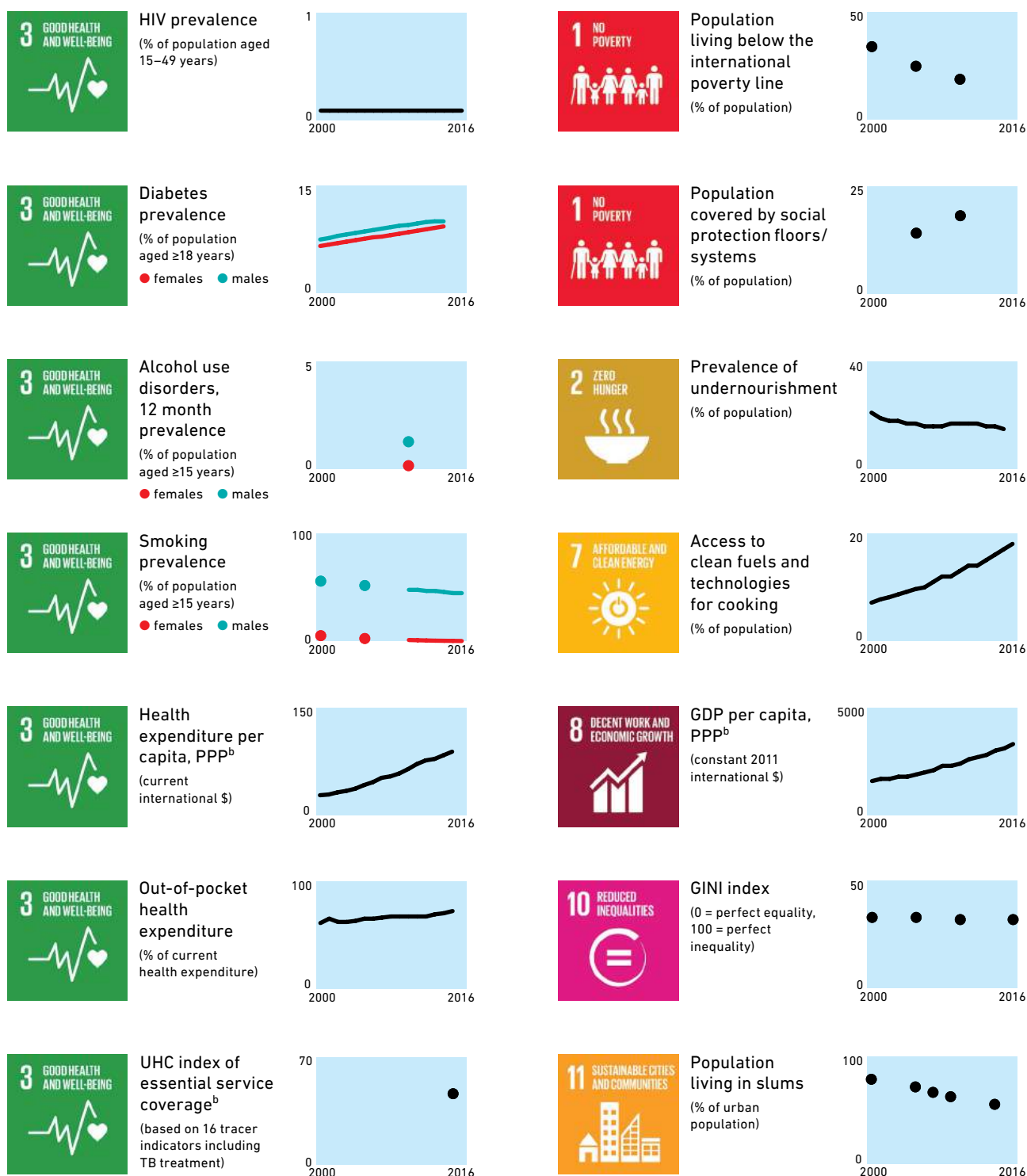
^b MDR is TB resistant to rifampicin and isoniazid; RR is TB resistant to rifampicin.

^c Includes cases with unknown previous TB treatment history.

^d Includes patients diagnosed before 2017 and patients who were not laboratory-confirmed.



INDICATORS IN THE SUSTAINABLE DEVELOPMENT GOALS ASSOCIATED WITH TB INCIDENCE^a



Targets for reductions in TB incidence and TB deaths set in WHO's End TB Strategy and the United Nations' Sustainable Development Goals (SDGs) are ambitious. Achieving them requires progress in reducing health-related risk factors for TB infection and disease, as well as broader social and economic determinants of TB infection and disease. WHO has developed a TB-SDG monitoring framework that comprises 14 indicators under seven SDGs for which there is evidence of an association with TB incidence. Further details are provided in [Chapter 2](#).

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^b GDP = gross domestic product; PPP = purchasing power parity; UHC = universal health coverage

ESTIMATES OF TB BURDEN, ^a 2017

	NUMBER (THOUSANDS)	RATE (PER 100 000 POPULATION)
Mortality (excludes HIV+TB)	5.1 (4.8–5.3)	2.4 (2.3–2.5)
Mortality (HIV+TB only)	1.9 (1.4–2.5)	0.91 (0.67–1.2)
Incidence (includes HIV+TB)	91 (78–105)	44 (37–50)
Incidence (HIV+TB only)	11 (9.3–13)	5.3 (4.5–6.3)
Incidence (MDR/RR-TB) ^b	2.4 (1.8–3.1)	1.2 (0.86–1.5)

ESTIMATED TB INCIDENCE BY AGE AND SEX (THOUSANDS), ^a 2017

	0–14 YEARS	> 14 YEARS	TOTAL
Females	5.2 (5–5.4)	24 (22–26)	29 (27–32)
Males	5.7 (5.5–5.9)	56 (50–63)	62 (54–70)
Total	11 (10–11)	80 (69–92)	91 (78–105)

TB CASE NOTIFICATIONS, 2017

Total cases notified	86 858
Total new and relapse	79 222
— % tested with rapid diagnostics at time of diagnosis	26%
— % with known HIV status	78%
— % pulmonary	87%
— % bacteriologically confirmed among pulmonary	72%

UNIVERSAL HEALTH COVERAGE AND SOCIAL PROTECTION

TB treatment coverage (notified/estimated incidence), 2017	87% (75–100)
TB patients facing catastrophic total costs	
TB case fatality ratio (estimated mortality/estimated incidence), 2017	0.08 (0.06–0.09)

TB/HIV CARE IN NEW AND RELAPSE TB PATIENTS, 2017

	NUMBER	(%)
Patients with known HIV-status who are HIV-positive	7 831	13%
— on antiretroviral therapy	3 742	48%

DRUG-RESISTANT TB CARE, 2017

	NEW CASES	PREVIOUSLY TREATED CASES	TOTAL NUMBER ^c
Estimated MDR/RR-TB cases among notified pulmonary TB cases			2 000 (1 600–2 400)
Estimated % of TB cases with MDR/RR-TB	1.5% (1.1–2)	8% (6–10)	
% notified tested for rifampicin resistance	30%	39%	27 716
MDR/RR-TB cases tested for resistance to second-line drugs			173
Laboratory-confirmed cases		MDR/RR-TB: 1 110, XDR-TB: 16	
Patients started on treatment ^d		MDR/RR-TB: 964, XDR-TB: 16	

TREATMENT SUCCESS RATE AND COHORT SIZE

	SUCCESS	COHORT
New and relapse cases registered in 2016	72%	75 223
Previously treated cases, excluding relapse, registered in 2016	39%	7 251
HIV-positive TB cases registered in 2016	53%	7 366
MDR/RR-TB cases started on second-line treatment in 2015	60%	954
XDR-TB cases started on second-line treatment in 2015	11%	27

TB PREVENTIVE TREATMENT, 2017

% of HIV-positive people (newly enrolled in care) on preventive treatment	
% of children (aged < 5) household contacts of bacteriologically-confirmed TB cases on preventive treatment	

TB FINANCING, 2018

National TB budget (US\$ millions)	57
Funding source:	85% domestic, 0% international, 15% unfunded

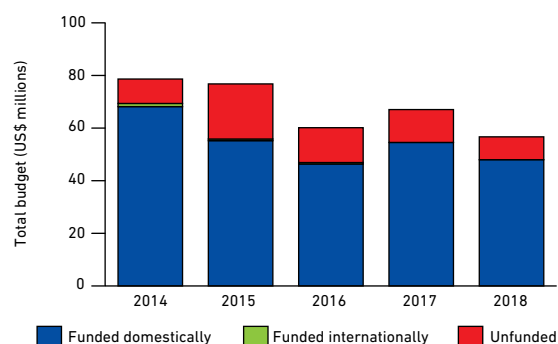
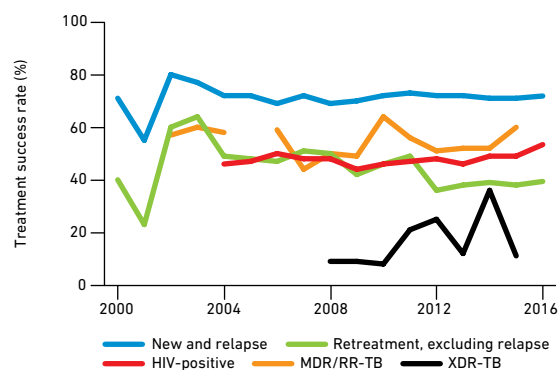
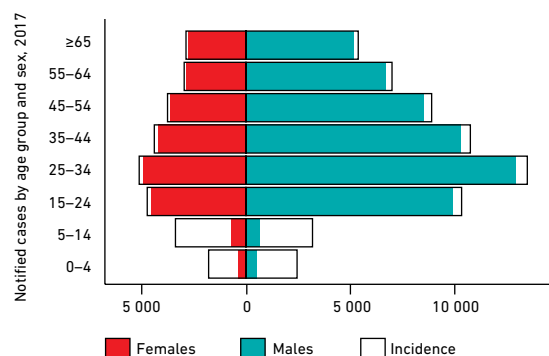
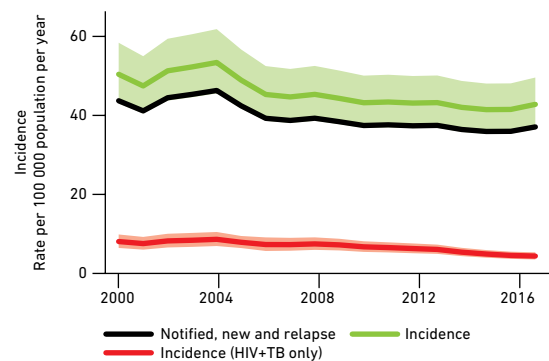
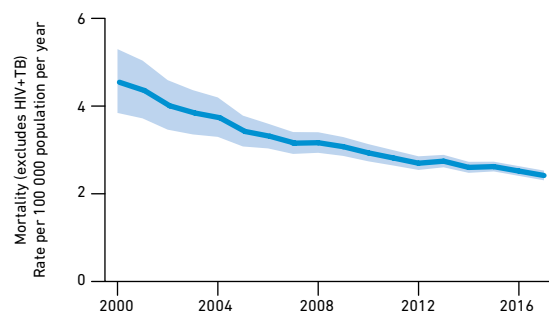
Data are as reported to WHO. Estimates of TB and MDR/RR-TB burden are produced by WHO in consultation with countries. Estimates are rounded and totals are computed prior to rounding.

^a Ranges represent uncertainty intervals.

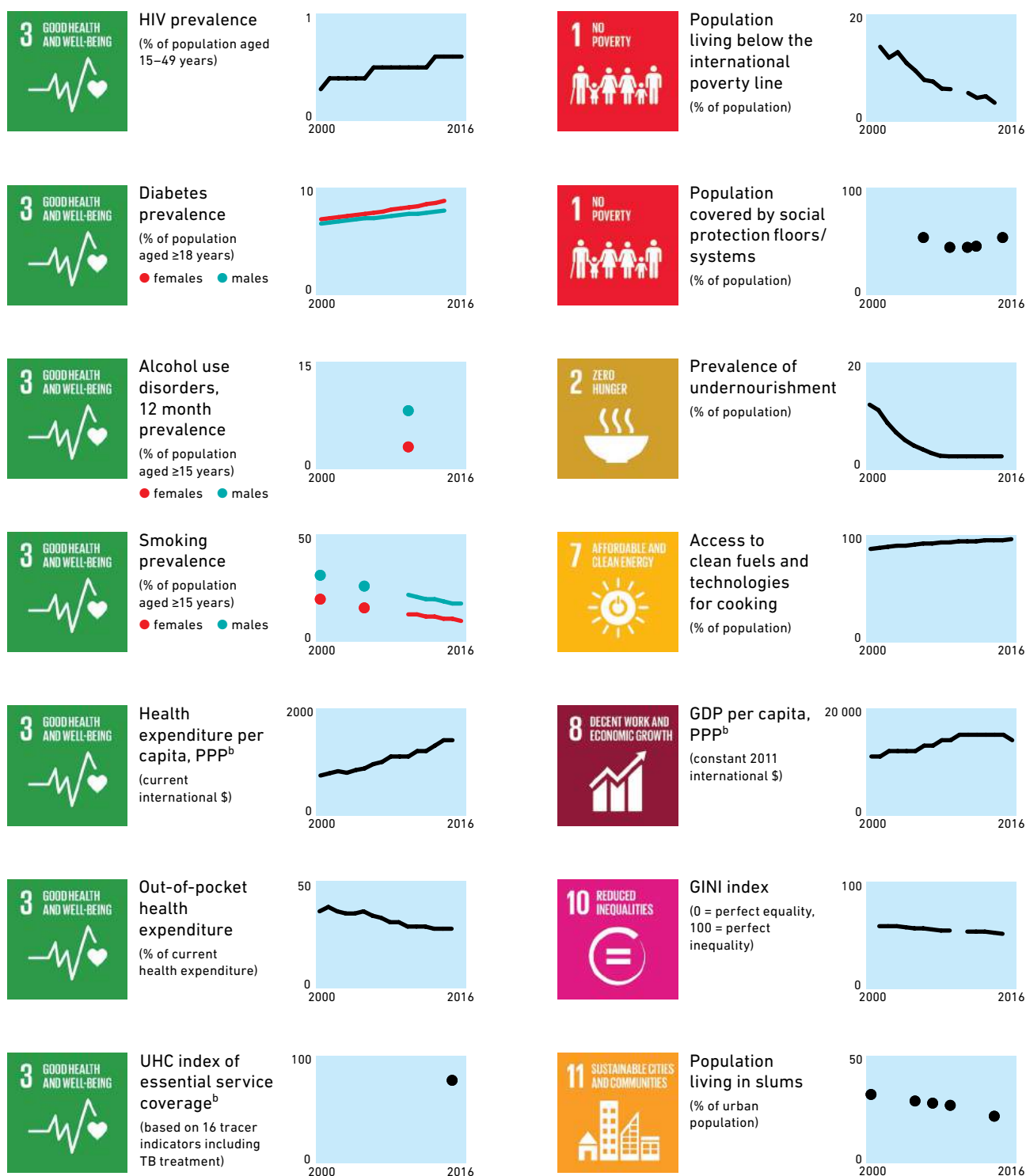
^b MDR is TB resistant to rifampicin and isoniazid; RR is TB resistant to rifampicin.

^c Includes cases with unknown previous TB treatment history.

^d Includes patients diagnosed before 2017 and patients who were not laboratory-confirmed.



INDICATORS IN THE SUSTAINABLE DEVELOPMENT GOALS ASSOCIATED WITH TB INCIDENCE^a



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ESTIMATES OF TB BURDEN, 2017

	NUMBER (THOUSANDS)	RATE (PER 100 000 POPULATION)
Mortality (excludes HIV+TB)	37 (33–41)	2.6 (2.4–2.9)
Mortality (HIV+TB only)	1.8 (0.84–3.1)	0.13 (0.06–0.22)
Incidence (includes HIV+TB)	889 (761–1 030)	63 (54–73)
Incidence (HIV+TB only)	12 (6.3–18)	0.82 (0.45–1.3)
Incidence (MDR/RR-TB) ^b	73 (55–94)	5.2 (3.9–6.6)

ESTIMATED TB INCIDENCE BY AGE AND SEX (THOUSANDS), 2017

	0–14 YEARS	> 14 YEARS	TOTAL
Females	47 (46–49)	241 (223–260)	289 (264–313)
Males	52 (50–54)	548 (484–613)	600 (526–674)
Total	99 (94–104)	790 (678–901)	889 (761–1 030)

TB CASE NOTIFICATIONS, 2017

Total cases notified	778 390
Total new and relapse	773 150
— % tested with rapid diagnostics at time of diagnosis	
— % with known HIV status	55%
— % pulmonary	95%
— % bacteriologically confirmed among pulmonary	32%

UNIVERSAL HEALTH COVERAGE AND SOCIAL PROTECTION

TB treatment coverage (notified/estimated incidence), 2017	87% (75–100)
TB patients facing catastrophic total costs	
TB case fatality ratio (estimated mortality/estimated incidence), 2017	0.04 (0.04–0.05)

TB/HIV CARE IN NEW AND RELAPSE TB PATIENTS, 2017

	NUMBER	(%)
Patients with known HIV-status who are HIV-positive	4 246	1%
— on antiretroviral therapy		

DRUG-RESISTANT TB CARE, 2017

	NEW CASES	PREVIOUSLY TREATED CASES	TOTAL NUMBER ^e
Estimated MDR/RR-TB cases among notified pulmonary TB cases ^d			58 000 (46 000–69 000)
Estimated % of TB cases with MDR/RR-TB	7.1% (5.6–8.7)	24% (20–28)	
% notified tested for rifampicin resistance	12%	69%	108 270
MDR/RR-TB cases tested for resistance to second-line drugs			
Laboratory-confirmed cases		MDR/RR-TB: 13 069, XDR-TB:	
Patients started on treatment ^e		MDR/RR-TB: 5 943, XDR-TB:	

TREATMENT SUCCESS RATE AND COHORT SIZE

	SUCCESS	COHORT
New and relapse cases registered in 2016	93%	778 493
Previously treated cases, excluding relapse, registered in 2016	81%	5 349
HIV-positive TB cases registered in 2016	51%	4 612
MDR/RR-TB cases started on second-line treatment in 2015	41%	5 691
XDR-TB cases started on second-line treatment in 2015		

TB PREVENTIVE TREATMENT, 2017

% of HIV-positive people (newly enrolled in care) on preventive treatment	
% of children (aged < 5) household contacts of bacteriologically-confirmed TB cases on preventive treatment	

TB FINANCING, 2018

National TB budget (US\$ millions)	609
Funding source:	87% domestic, 2% international, 11% unfunded

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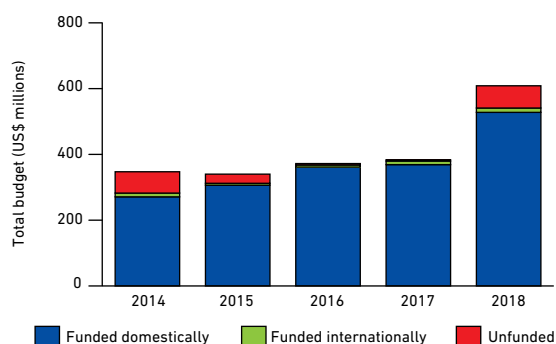
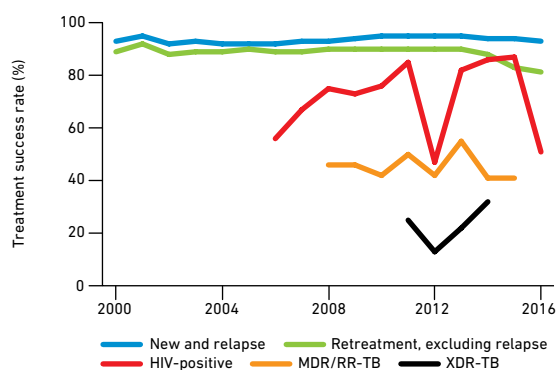
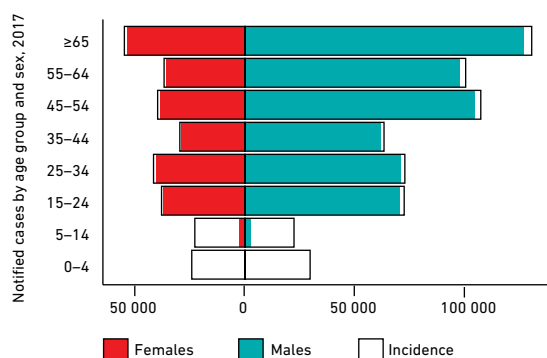
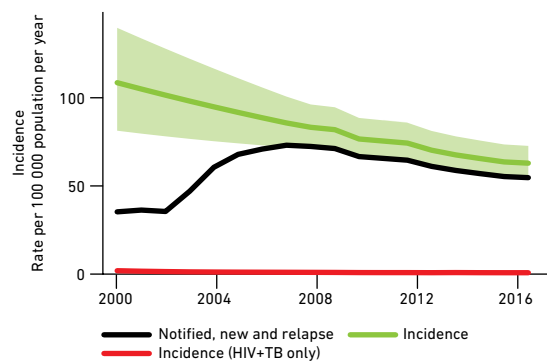
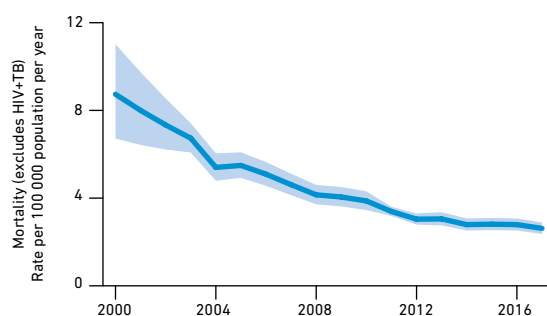
^a Ranges represent uncertainty intervals.

^b MDR is TB resistant to rifampicin and isoniazid; RR is TB resistant to rifampicin.

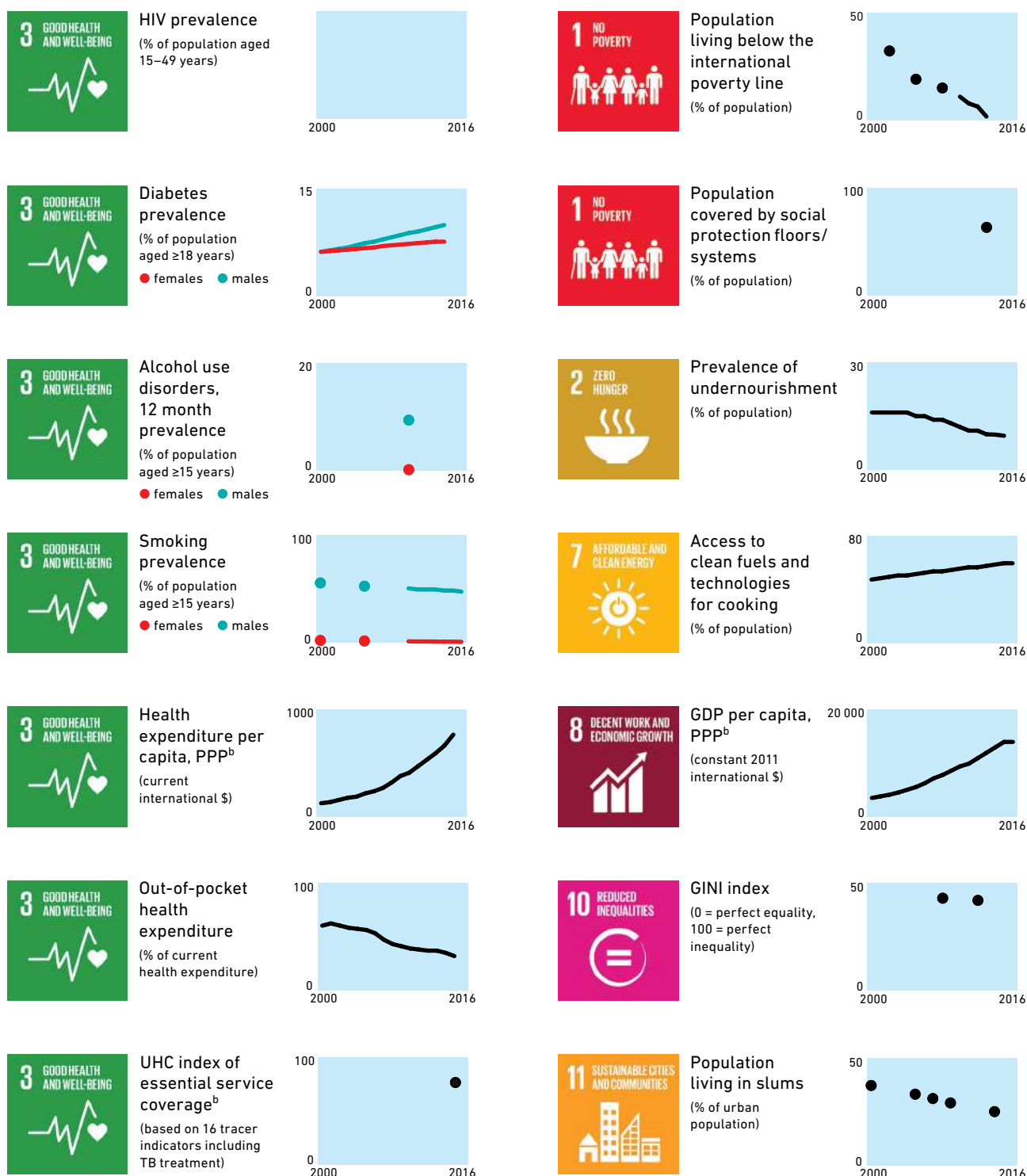
^c Includes cases with unknown previous TB treatment history.

^d The estimated number of MDR/RR-TB cases among bacteriologically confirmed pulmonary cases is 21 000 (17 000–25 000).

^e Includes patients diagnosed before 2017 and patients who were not laboratory-confirmed.



INDICATORS IN THE SUSTAINABLE DEVELOPMENT GOALS ASSOCIATED WITH TB INCIDENCE^a



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ESTIMATES OF TB BURDEN, ^a 2017

	NUMBER (THOUSANDS)	RATE (PER 100 000 POPULATION)
Mortality (excludes HIV+TB)	16 (11–22)	63 (43–86)
Mortality (HIV+TB only)	0.044 (0.022–0.072)	0.17 (0.09–0.28)
Incidence (includes HIV+TB)	131 (114–149)	513 (446–584)
Incidence (HIV+TB only)	0.17 (0.094–0.28)	0.69 (0.37–1.1)
Incidence (MDR/RR-TB) ^b	5.2 (2.6–8.8)	20 (10–34)

ESTIMATED TB INCIDENCE BY AGE AND SEX (THOUSANDS), ^a 2017

	0–14 YEARS	> 14 YEARS	TOTAL
Females	6.2 (6–6.4)	43 (40–47)	50 (45–54)
Males	8.4 (8.1–8.6)	73 (66–80)	81 (73–90)
Total	15 (14–15)	116 (101–131)	131 (114–149)

TB CASE NOTIFICATIONS, 2017

Total cases notified	107 103
Total new and relapse	100 553
— % tested with rapid diagnostics at time of diagnosis	
— % with known HIV status	
— % pulmonary	81%
— % bacteriologically confirmed among pulmonary	50%

UNIVERSAL HEALTH COVERAGE AND SOCIAL PROTECTION

TB treatment coverage (notified/estimated incidence), 2017	77% (68–88)
TB patients facing catastrophic total costs	
TB case fatality ratio (estimated mortality/estimated incidence), 2017	0.12 (0.08–0.17)

TB/HIV CARE IN NEW AND RELAPSE TB PATIENTS, 2017

	NUMBER	(%)
Patients with known HIV-status who are HIV-positive		
— on antiretroviral therapy		

DRUG-RESISTANT TB CARE, 2017

	NEW CASES	PREVIOUSLY TREATED CASES	TOTAL NUMBER ^e
Estimated MDR/RR-TB cases among notified pulmonary TB cases			4 100 (2 300–5 800)
Estimated % of TB cases with MDR/RR-TB	2.2% (0.82–4.2)	16% (9.1–25)	
% notified tested for rifampicin resistance		14%	2 116
MDR/RR-TB cases tested for resistance to second-line drugs			0
Laboratory-confirmed cases		MDR/RR-TB: 1 515, XDR-TB: 0	
Patients started on treatment ^d		MDR/RR-TB: 1 732, XDR-TB: 19	

TREATMENT SUCCESS RATE AND COHORT SIZE

	SUCCESS	COHORT
New cases registered in 2016		
Previously treated cases registered in 2016		
HIV-positive TB cases registered in 2016		
MDR/RR-TB cases started on second-line treatment in 2015	73%	325
XDR-TB cases started on second-line treatment in 2015		0

TB PREVENTIVE TREATMENT, 2017

% of HIV-positive people (newly enrolled in care) on preventive treatment	
% of children (aged < 5) household contacts of bacteriologically-confirmed TB cases on preventive treatment ^e	> 100%

TB FINANCING, 2018

National TB budget (US\$ millions)	84
Funding source:	7% domestic, 1% international, 92% unfunded

Data are as reported to WHO. Estimates of TB and MDR/RR-TB burden are produced by WHO in consultation with countries. Estimates are rounded and totals are computed prior to rounding.

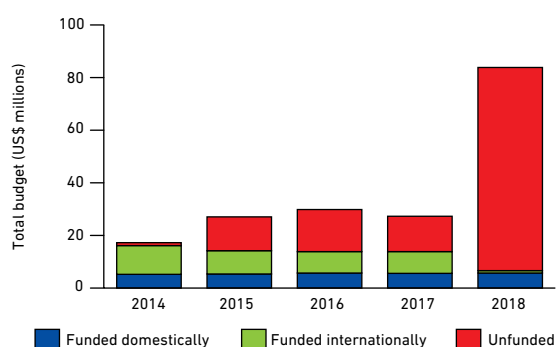
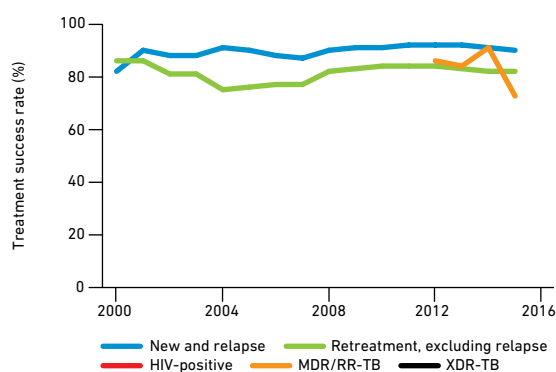
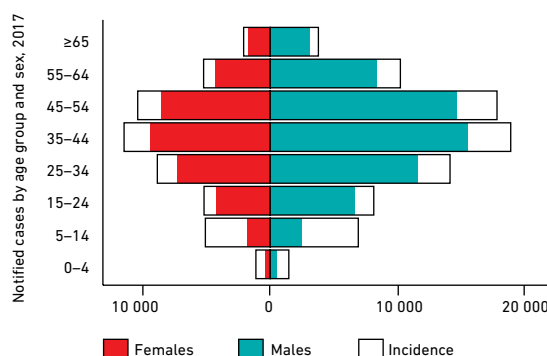
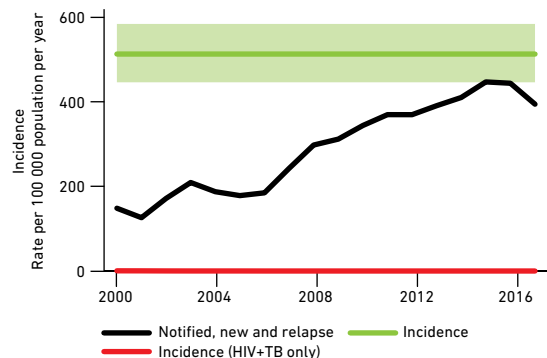
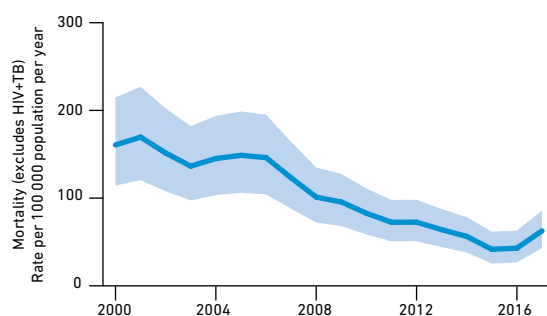
^a Ranges represent uncertainty intervals.

^b MDR is TB resistant to rifampicin and isoniazid; RR is TB resistant to rifampicin.

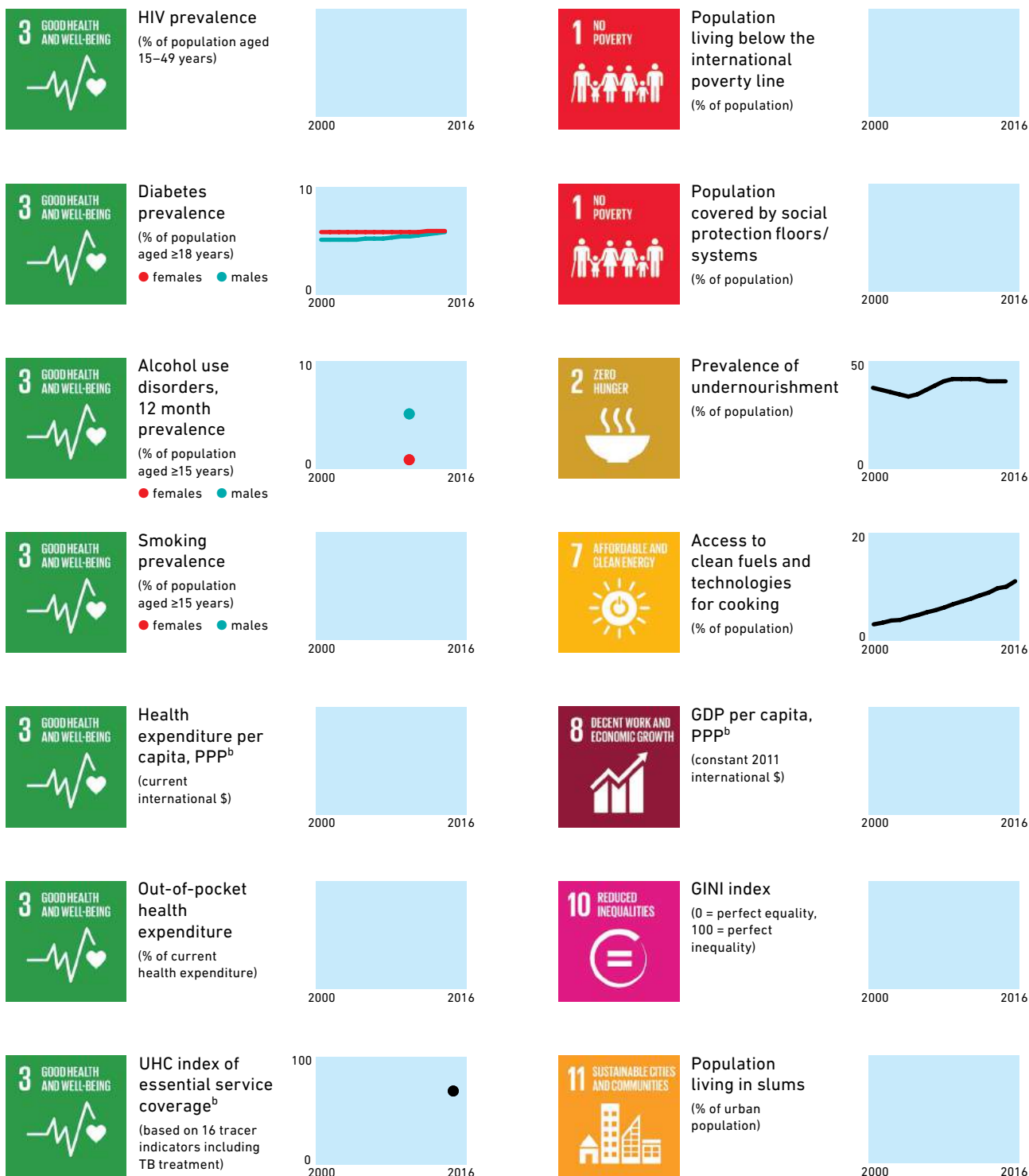
^c Includes cases with unknown previous TB treatment history.

^d Includes patients diagnosed before 2017 and patients who were not laboratory-confirmed.

^e Reasons for higher than expected coverage might be that the numerator did not exclude non-household contacts or children of five years and older.



INDICATORS IN THE SUSTAINABLE DEVELOPMENT GOALS ASSOCIATED WITH TB INCIDENCE^a



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^b GDP = gross domestic product; PPP = purchasing power parity; UHC = universal health coverage

Democratic Republic of the Congo

POPULATION 2017 **81 MILLION**

ESTIMATES OF TB BURDEN,^a 2017

	NUMBER (THOUSANDS)	RATE (PER 100 000 POPULATION)
Mortality (excludes HIV+TB)	49 (29–74)	60 (35–90)
Mortality (HIV+TB only)	7.5 (3.5–13)	9.2 (4.3–16)
Incidence (includes HIV+TB)	262 (169–374)	322 (208–460)
Incidence (HIV+TB only)	20 (13–29)	25 (16–35)
Incidence (MDR/RR-TB)**	7.5 (3.3–13)	9.2 (4–17)

ESTIMATED TB INCIDENCE BY AGE AND SEX (THOUSANDS),^a 2017

	0–14 YEARS	> 14 YEARS	TOTAL
Females	16 (14–17)	84 (65–102)	99 (76–123)
Males	17 (16–19)	145 (103–187)	162 (112–212)
Total	33 (28–37)	229 (145–313)	262 (169–374)

TB CASE NOTIFICATIONS, 2017

Total cases notified	151 832
Total new and relapse	150 085
— % tested with rapid diagnostics at time of diagnosis	
— % with known HIV status	64%
— % pulmonary	83%
— % bacteriologically confirmed among pulmonary	80%

UNIVERSAL HEALTH COVERAGE AND SOCIAL PROTECTION

TB treatment coverage (notified/estimated incidence), 2017	57% (40–89)
TB patients facing catastrophic total costs	
TB case fatality ratio (estimated mortality/estimated incidence), 2017	0.22 (0.11–0.35)

TB/HIV CARE IN NEW AND RELAPSE TB PATIENTS, 2017

	NUMBER	(%)
Patients with known HIV-status who are HIV-positive	9 688	10%
— on antiretroviral therapy	7 982	82%

DRUG-RESISTANT TB CARE, 2017

	NEW CASES	PREVIOUSLY TREATED CASES	TOTAL NUMBER ^e
Estimated MDR/RR-TB cases among notified pulmonary TB cases			3 400 (2 000–4 900)
Estimated % of TB cases with MDR/RR-TB	2.2% (1–3.5)	9.5% (8.8–10)	
% notified tested for rifampicin resistance	3%	57%	10 543
MDR/RR-TB cases tested for resistance to second-line drugs			147
Laboratory-confirmed cases		MDR/RR-TB: 893, XDR-TB: 19	
Patients started on treatment ^d		MDR/RR-TB: 839, XDR-TB: 15	

TREATMENT SUCCESS RATE AND COHORT SIZE

	SUCCESS	COHORT
New and relapse cases registered in 2016	89%	127 503
Previously treated cases, excluding relapse, registered in 2016	60%	789
HIV-positive TB cases registered in 2016		
MDR/RR-TB cases started on second-line treatment in 2015	90%	463
XDR-TB cases started on second-line treatment in 2015	10%	21

TB PREVENTIVE TREATMENT, 2017

% of HIV-positive people (newly enrolled in care) on preventive treatment	
% of children (aged < 5) household contacts of bacteriologically-confirmed TB cases on preventive treatment	15% (14–17)

TB FINANCING, 2018

National TB budget (US\$ millions)	74
Funding source:	2% domestic, 48% international, 49% unfunded

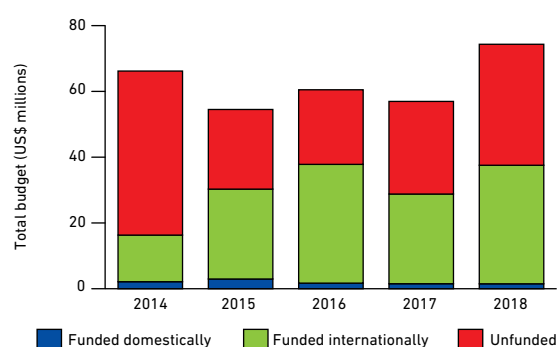
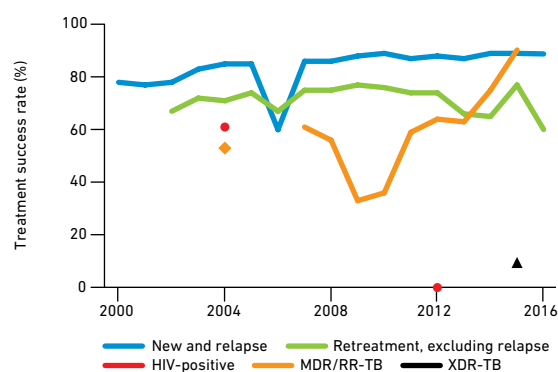
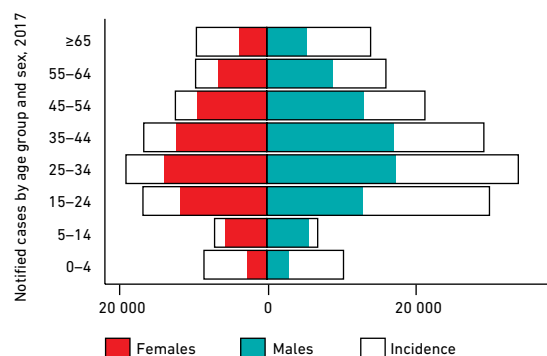
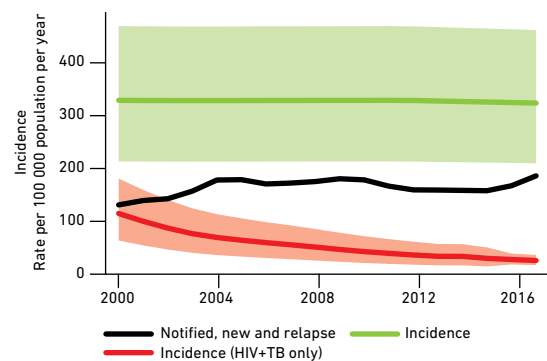
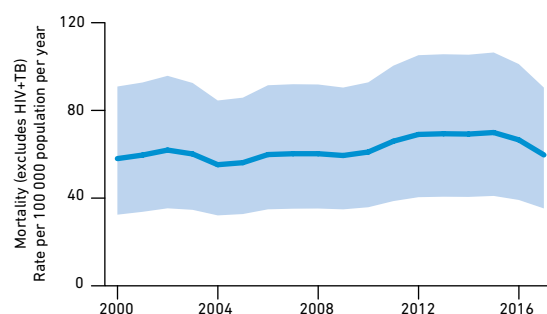
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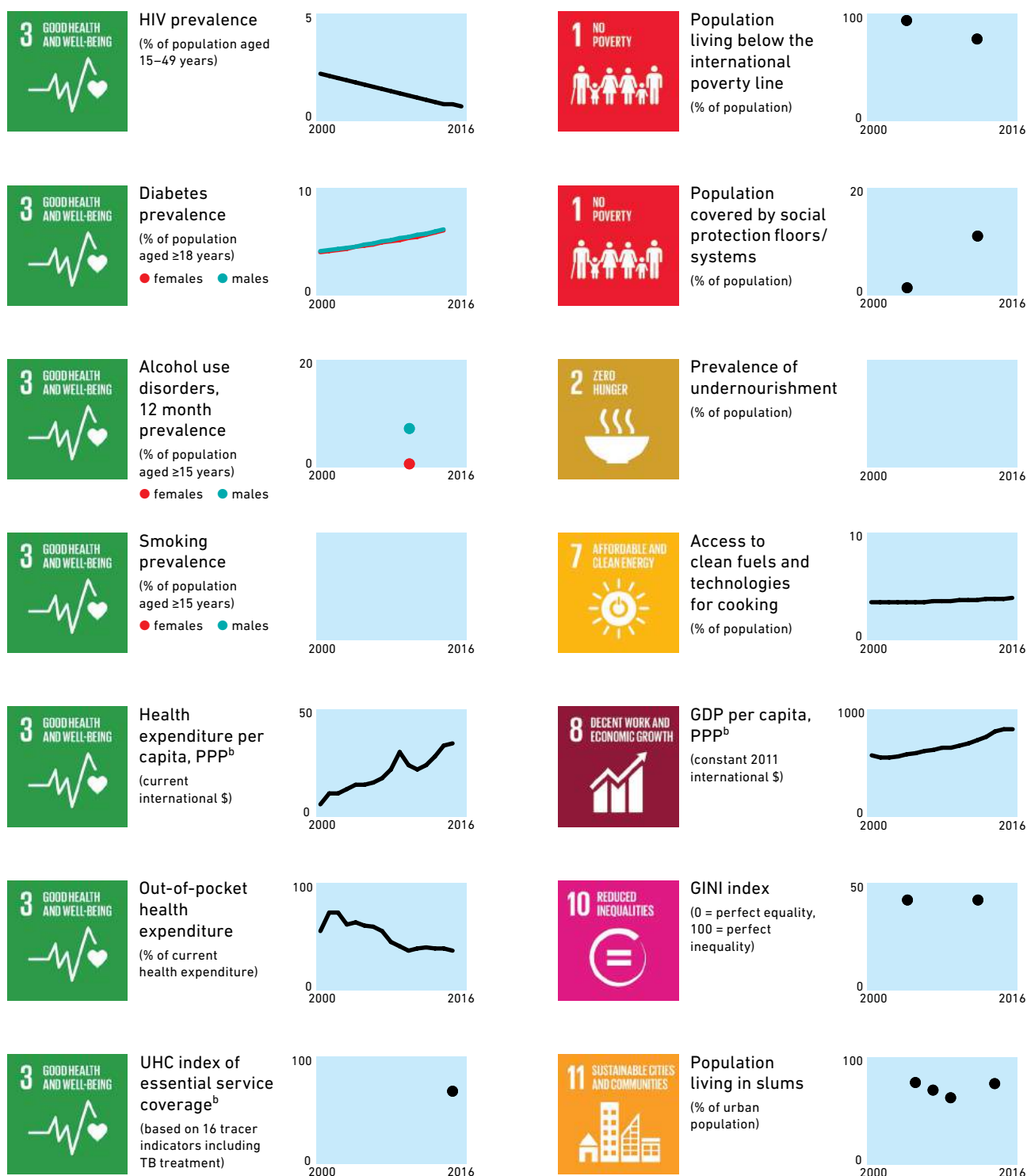
^b MDR is TB resistant to rifampicin and isoniazid; RR is TB resistant to rifampicin.

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Ethiopia

POPULATION 2017 **105 MILLION**

ESTIMATES OF TB BURDEN,^a 2017

	NUMBER (THOUSANDS)	RATE (PER 100 000 POPULATION)
Mortality (excludes HIV+TB)	25 (16–37)	24 (15–35)
Mortality (HIV+TB only)	3.6 (2.5–5)	3.5 (2.4–4.8)
Incidence (includes HIV+TB)	172 (121–232)	164 (115–221)
Incidence (HIV+TB only)	12 (8.6–17)	12 (8.2–16)
Incidence (MDR/RR-TB)**	5.5 (2.9–8.9)	5.2 (2.8–8.4)

ESTIMATED TB INCIDENCE BY AGE AND SEX (THOUSANDS),^a 2017

	0–14 YEARS	> 14 YEARS	TOTAL
Females	9.5 (8.7–10)	69 (55–83)	78 (61–95)
Males	10 (9.6–11)	83 (65–102)	94 (71–116)
Total	20 (18–22)	152 (106–198)	172 (121–232)

TB CASE NOTIFICATIONS, 2017

Total cases notified	117 705
Total new and relapse	116 725
— % tested with rapid diagnostics at time of diagnosis	
— % with known HIV status	86%
— % pulmonary	69%
— % bacteriologically confirmed among pulmonary	58%

UNIVERSAL HEALTH COVERAGE AND SOCIAL PROTECTION

TB treatment coverage (notified/estimated incidence), 2017	68% (50–96)
TB patients facing catastrophic total costs	
TB case fatality ratio (estimated mortality/estimated incidence), 2017	0.17 (0.1–0.26)

TB/HIV CARE IN NEW AND RELAPSE TB PATIENTS, 2017

	NUMBER	(%)
Patients with known HIV-status who are HIV-positive	7 272	7%
— on antiretroviral therapy	6 673	92%

DRUG-RESISTANT TB CARE, 2017

	NEW CASES	PREVIOUSLY TREATED CASES	TOTAL NUMBER ^e
Estimated MDR/RR-TB cases among notified pulmonary TB cases			2 700 (1 700–3 700)
Estimated % of TB cases with MDR/RR-TB	2.7% (1.6–4.1)	14% (6.7–25)	
% notified tested for rifampicin resistance			38 501
MDR/RR-TB cases tested for resistance to second-line drugs			205
Laboratory-confirmed cases		MDR/RR-TB: 680, XDR-TB: 4	
Patients started on treatment ^d		MDR/RR-TB: 680, XDR-TB: 4	

TREATMENT SUCCESS RATE AND COHORT SIZE

	SUCCESS	COHORT
New cases registered in 2016	90%	123 004
Previously treated cases registered in 2016		
HIV-positive TB cases registered in 2016		
MDR/RR-TB cases started on second-line treatment in 2015	75%	660
XDR-TB cases started on second-line treatment in 2015		

TB PREVENTIVE TREATMENT, 2017

% of HIV-positive people (newly enrolled in care) on preventive treatment	45%
% of children (aged < 5) household contacts of bacteriologically-confirmed TB cases on preventive treatment	11% (9.9–12)

TB FINANCING, 2018

National TB budget (US\$ millions)	93
Funding source:	11% domestic, 33% international, 56% unfunded

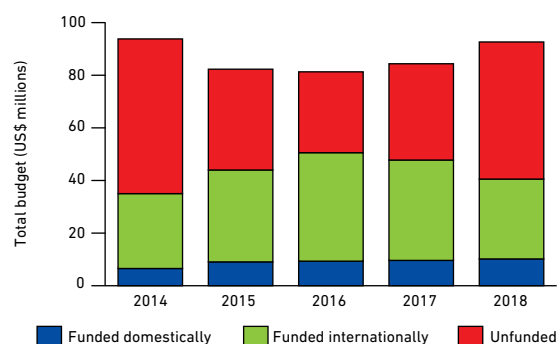
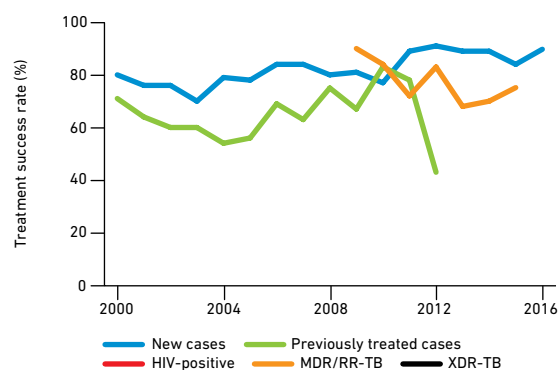
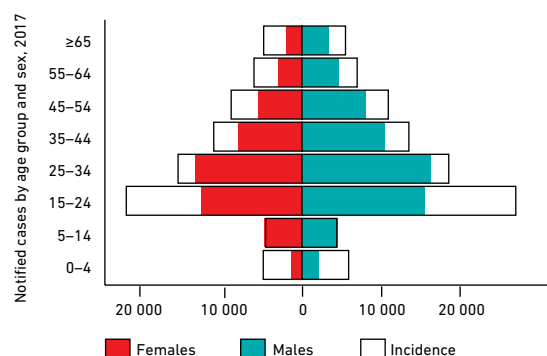
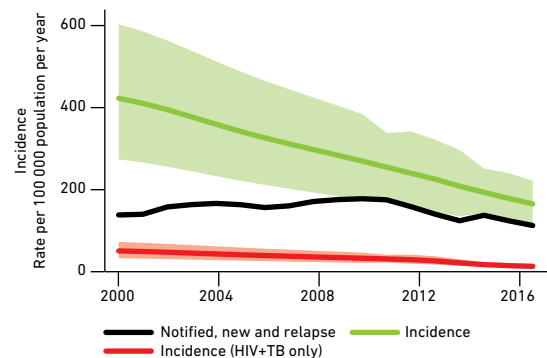
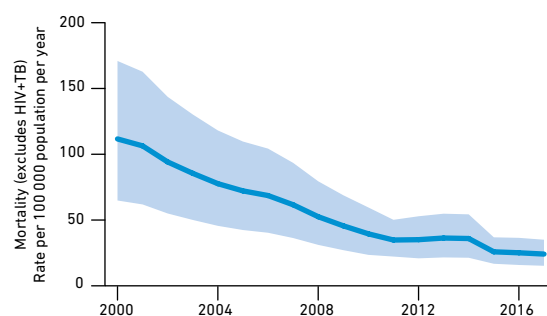
Data are as reported to WHO. Estimates of TB and MDR/RR-TB burden are produced by WHO in consultation with countries. Estimates are rounded and totals are computed prior to rounding.

^a Ranges represent uncertainty intervals.

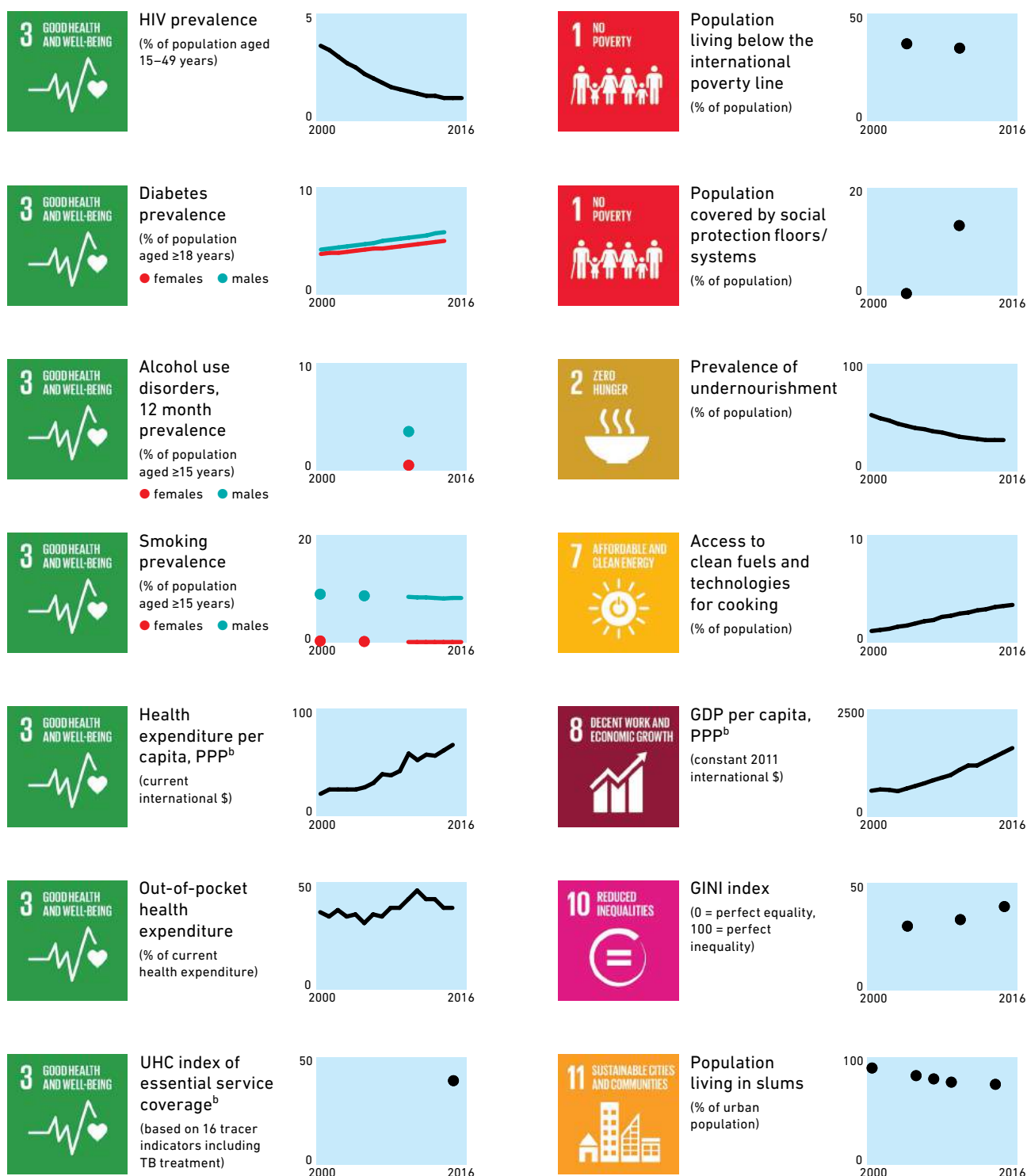
^b MDR is TB resistant to rifampicin and isoniazid; RR is TB resistant to rifampicin.

^c Includes cases with unknown previous TB treatment history.

^d Includes patients diagnosed before 2017 and patients who were not laboratory-confirmed.



INDICATORS IN THE SUSTAINABLE DEVELOPMENT GOALS ASSOCIATED WITH TB INCIDENCE^a



Targets for reductions in TB incidence and TB deaths set in WHO's End TB Strategy and the United Nations' Sustainable Development Goals (SDGs) are ambitious. Achieving them requires progress in reducing health-related risk factors for TB infection and disease, as well as broader social and economic determinants of TB infection and disease. WHO has developed a TB-SDG monitoring framework that comprises 14 indicators under seven SDGs for which there is evidence of an association with TB incidence. Further details are provided in [Chapter 2](#).

^a Data sources: SDG indicators database, The World Bank, World Health Organization. Missing values and empty boxes indicate data not available in these data sources.

^b GDP = gross domestic product; PPP = purchasing power parity; UHC = universal health coverage

ESTIMATES OF TB BURDEN,^a 2017

	NUMBER (THOUSANDS)	RATE (PER 100 000 POPULATION)
Mortality (excludes HIV+TB)	410 (381–441)	31 (28–33)
Mortality (HIV+TB only)	11 (6.5–16)	0.79 (0.48–1.2)
Incidence (includes HIV+TB)	2 740 (1 870–3 770)	204 (140–281)
Incidence (HIV+TB only)	86 (57–120)	6.4 (4.3–9)
Incidence (MDR/RR-TB)**	135 (78–208)	10 (5.8–16)

ESTIMATED TB INCIDENCE BY AGE AND SEX (THOUSANDS),^a 2017

	0–14 YEARS	> 14 YEARS	TOTAL
Females	107 (100–114)	847 (684–1 010)	954 (759–1 150)
Males	117 (109–126)	1 670 (1 220–2 120)	1 780 (1 290–2 280)
Total	224 (202–247)	2 510 (1 680–3 350)	2 740 (1 870–3 770)

TB CASE NOTIFICATIONS, 2017

Total cases notified	1 908 371
Total new and relapse	1 786 681
— % tested with rapid diagnostics at time of diagnosis	40%
— % with known HIV status	64%
— % pulmonary	85%
— % bacteriologically confirmed among pulmonary	60%

UNIVERSAL HEALTH COVERAGE AND SOCIAL PROTECTION

TB treatment coverage (notified/estimated incidence), 2017	65% (47–96)
TB patients facing catastrophic total costs	
TB case fatality ratio (estimated mortality/estimated incidence), 2017	0.16 (0.11–0.22)

TB/HIV CARE IN NEW AND RELAPSE TB PATIENTS, 2017

	NUMBER	(%)
Patients with known HIV-status who are HIV-positive	36 440	3%
— on antiretroviral therapy	28 651	79%

DRUG-RESISTANT TB CARE, 2017

	NEW CASES	PREVIOUSLY TREATED CASES	TOTAL NUMBER ^e
Estimated MDR/RR-TB cases among notified pulmonary TB cases			65 000 (54 000–76 000)
Estimated % of TB cases with MDR/RR-TB	2.8% (2–3.5)	12% (10–13)	
% notified tested for rifampicin resistance	32%	82%	720 051
MDR/RR-TB cases tested for resistance to second-line drugs			26 832
Laboratory-confirmed cases		MDR/RR-TB: 39 009, XDR-TB: 2 650	
Patients started on treatment ^d		MDR/RR-TB: 35 950, XDR-TB: 2 838	

TREATMENT SUCCESS RATE AND COHORT SIZE

	SUCCESS	COHORT
New and relapse cases registered in 2016	69%	1 763 876
Previously treated cases, excluding relapse, registered in 2016	70%	172 282
HIV-positive TB cases registered in 2016	75%	39 123
MDR/RR-TB cases started on second-line treatment in 2015	46%	26 966
XDR-TB cases started on second-line treatment in 2015	28%	2 130

TB PREVENTIVE TREATMENT, 2017

% of HIV-positive people (newly enrolled in care) on preventive treatment	10%
% of children (aged < 5) household contacts of bacteriologically-confirmed TB cases on preventive treatment	11% (10–12)

TB FINANCING, 2018

National TB budget (US\$ millions)	580
Funding source:	79% domestic, 21% international, 0% unfunded

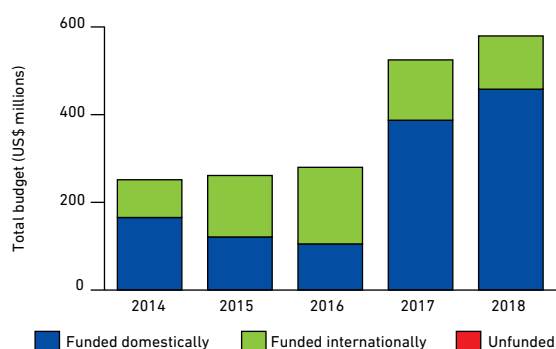
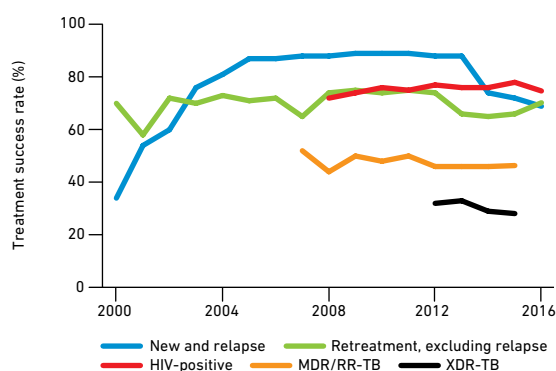
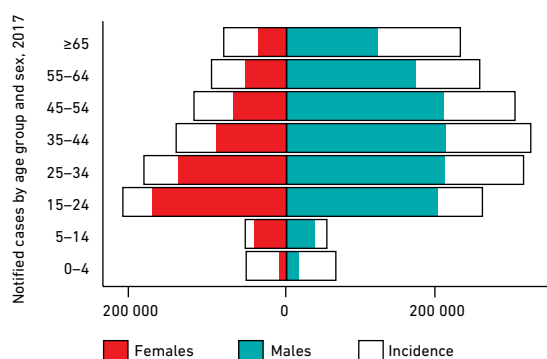
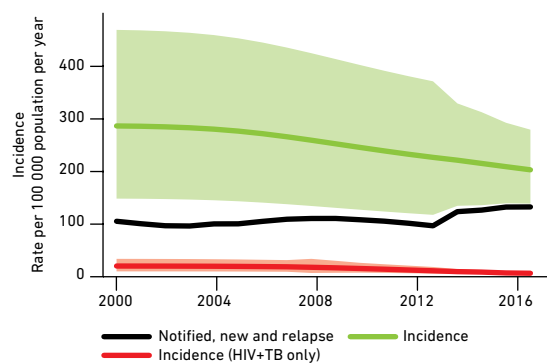
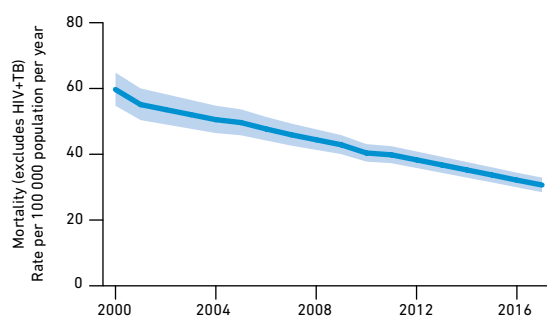
Data are as reported to WHO. Estimates of TB and MDR/RR-TB burden are produced by WHO in consultation with countries. Estimates are rounded and totals are computed prior to rounding.

^a Ranges represent uncertainty intervals. Estimates of TB incidence and mortality for India are interim in nature, pending results from the national TB prevalence survey planned for 2019/2020.

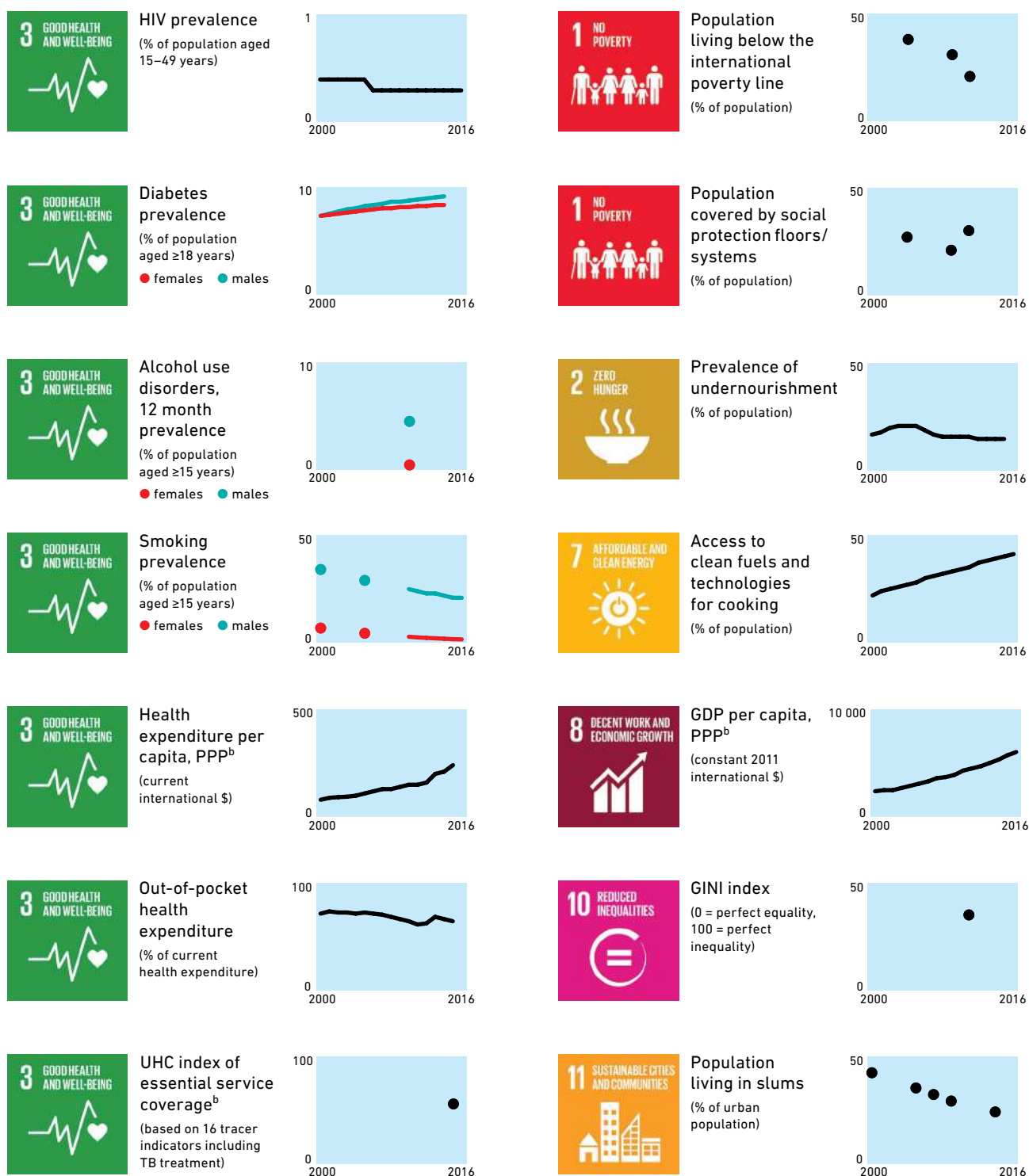
^b MDR is TB resistant to rifampicin and isoniazid; RR is TB resistant to rifampicin.

^c Includes cases with unknown previous TB treatment history.

^d Includes patients diagnosed before 2017 and patients who were not laboratory-confirmed.



INDICATORS IN THE SUSTAINABLE DEVELOPMENT GOALS ASSOCIATED WITH TB INCIDENCE^a



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^b GDP = gross domestic product; PPP = purchasing power parity; UHC = universal health coverage

Indonesia

POPULATION 2017 **264 MILLION**

ESTIMATES OF TB BURDEN,^a 2017

	NUMBER (THOUSANDS)	RATE (PER 100 000 POPULATION)
Mortality (excludes HIV+TB)	107 (100–114)	40 (38–43)
Mortality (HIV+TB only)	9.4 (5–15)	3.6 (1.9–5.8)
Incidence (includes HIV+TB)	842 (767–919)	319 (291–348)
Incidence (HIV+TB only)	36 (20–57)	14 (7.7–21)
Incidence (MDR/RR-TB) ^b	23 (16–31)	8.8 (6.2–12)

ESTIMATED TB INCIDENCE BY AGE AND SEX (THOUSANDS),^a 2017

	0–14 YEARS	> 14 YEARS	TOTAL
Females	23 (23–23)	326 (308–345)	349 (329–370)
Males	26 (26–27)	466 (435–497)	492 (458–526)
Total	49 (48–50)	792 (723–862)	842 (767–919)

TB CASE NOTIFICATIONS, 2017

Total cases notified	446 732
Total new and relapse	442 172
— % tested with rapid diagnostics at time of diagnosis	2%
— % with known HIV status	29%
— % pulmonary	90%
— % bacteriologically confirmed among pulmonary	54%

UNIVERSAL HEALTH COVERAGE AND SOCIAL PROTECTION

TB treatment coverage (notified/estimated incidence), 2017	53% (48–58)
TB patients facing catastrophic total costs	
TB case fatality ratio (estimated mortality/estimated incidence), 2017	0.14 (0.12–0.15)

TB/HIV CARE IN NEW AND RELAPSE TB PATIENTS, 2017

	NUMBER	(%)
Patients with known HIV-status who are HIV-positive	7 729	6%
— on antiretroviral therapy	2 244	29%

DRUG-RESISTANT TB CARE, 2017

	NEW CASES	PREVIOUSLY TREATED CASES	TOTAL NUMBER ^c
Estimated MDR/RR-TB cases among notified pulmonary TB cases			12 000 (8 600–15 000)
Estimated % of TB cases with MDR/RR-TB	2.4% (1.8–3.3)	13% (9–18)	
% notified tested for rifampicin resistance	16%	223%	112 743
MDR/RR-TB cases tested for resistance to second-line drugs			1 813
Laboratory-confirmed cases		MDR/RR-TB: 5 070, XDR-TB: 51	
Patients started on treatment ^d		MDR/RR-TB: 3 042, XDR-TB: 50	

TREATMENT SUCCESS RATE AND COHORT SIZE

	SUCCESS	COHORT
New and relapse cases registered in 2016	86%	364 671
Previously treated cases, excluding relapse, registered in 2016	71%	2 002
HIV-positive TB cases registered in 2016	64%	4 470
MDR/RR-TB cases started on second-line treatment in 2015	47%	1 565
XDR-TB cases started on second-line treatment in 2015	28%	32

TB PREVENTIVE TREATMENT, 2017

% of HIV-positive people (newly enrolled in care) on preventive treatment	16%
% of children (aged < 5) household contacts of bacteriologically-confirmed TB cases on preventive treatment	8.5% (7.8–9.3)

TB FINANCING, 2018

National TB budget (US\$ millions)	294
Funding source:	34% domestic, 16% international, 49% unfunded

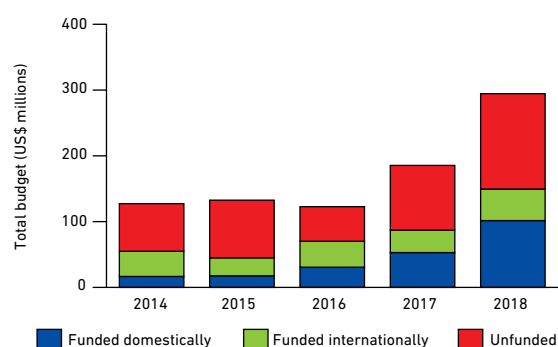
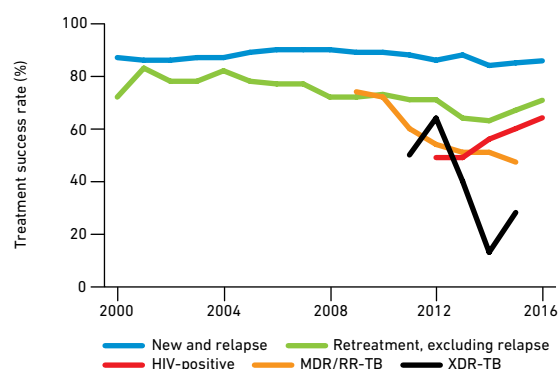
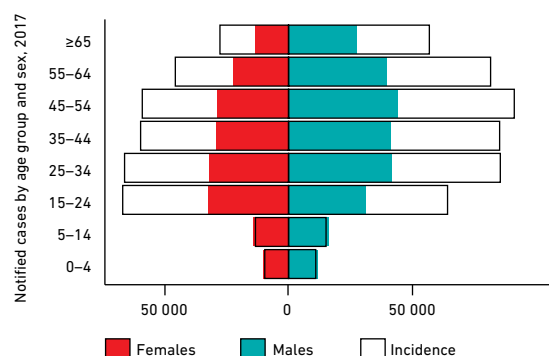
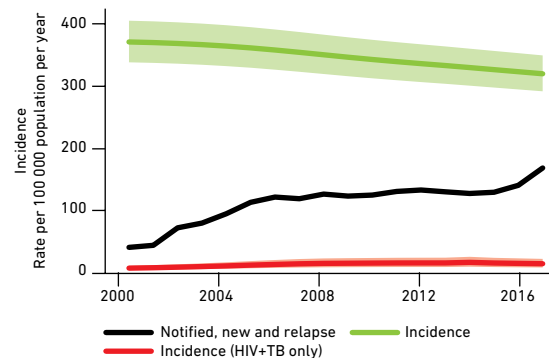
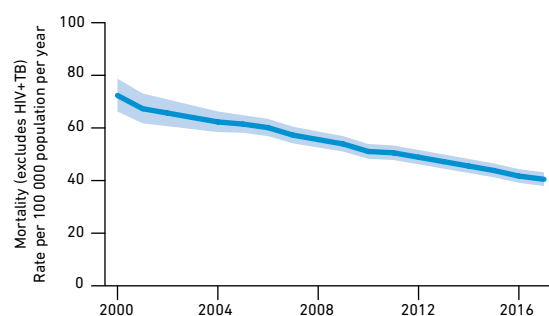
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^a Ranges represent uncertainty intervals.

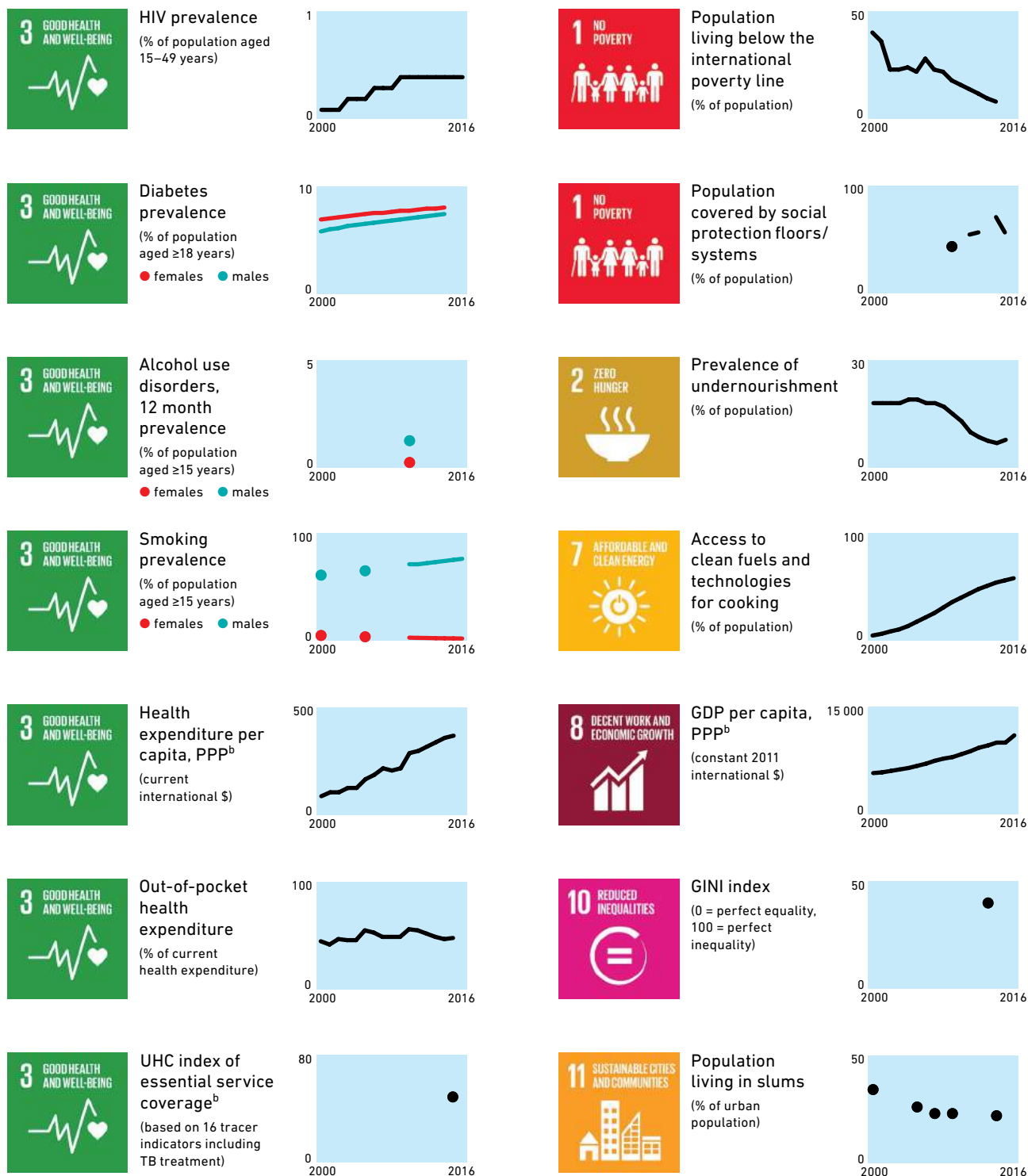
^b MDR is TB resistant to rifampicin and isoniazid; RR is TB resistant to rifampicin.

^c Includes cases with unknown previous TB treatment history.

^d Includes patients diagnosed before 2017 and patients who were not laboratory-confirmed.



INDICATORS IN THE SUSTAINABLE DEVELOPMENT GOALS ASSOCIATED WITH TB INCIDENCE^a



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^b GDP = gross domestic product; PPP = purchasing power parity; UHC = universal health coverage

ESTIMATES OF TB BURDEN, ^a 2017

	NUMBER (THOUSANDS)	RATE (PER 100 000 POPULATION)
Mortality (excludes HIV+TB)	25 (14–39)	50 (28–78)
Mortality (HIV+TB only)	18 (11–27)	37 (22–55)
Incidence (includes HIV+TB)	158 (97–235)	319 (195–472)
Incidence (HIV+TB only)	45 (27–68)	91 (55–137)
Incidence (MDR/RR-TB) ^b	2.8 (1.2–4.9)	5.6 (2.5–9.9)

ESTIMATED TB INCIDENCE BY AGE AND SEX (THOUSANDS), ^a 2017

	0–14 YEARS	> 14 YEARS	TOTAL
Females	9.9 (8.8–11)	43 (33–53)	53 (39–66)
Males	11 (9.7–12)	95 (63–127)	106 (68–143)
Total	21 (18–24)	138 (82–193)	158 (97–235)

TB CASE NOTIFICATIONS, 2017

Total cases notified	85 188
Total new and relapse	83 599
— % tested with rapid diagnostics at time of diagnosis	47%
— % with known HIV status	96%
— % pulmonary	84%
— % bacteriologically confirmed among pulmonary	67%

UNIVERSAL HEALTH COVERAGE AND SOCIAL PROTECTION

TB treatment coverage (notified/estimated incidence), 2017	53% (36–86)
TB patients facing catastrophic total costs, 2017	27% (21–32)
TB case fatality ratio (estimated mortality/estimated incidence), 2017	0.28 (0.15–0.45)

TB/HIV CARE IN NEW AND RELAPSE TB PATIENTS, 2017

	NUMBER	(%)
Patients with known HIV-status who are HIV-positive	22 992	29%
— on antiretroviral therapy	21 763	95%

DRUG-RESISTANT TB CARE, 2017

	NEW CASES	PREVIOUSLY TREATED CASES	TOTAL NUMBER ^c
Estimated MDR/RR-TB cases among notified pulmonary TB cases			1 100 (670–1 500)
Estimated % of TB cases with MDR/RR-TB	1.3% (0.74–2)	4.4% (3.7–5.2)	
% notified tested for rifampicin resistance	46%	52%	39 598
MDR/RR-TB cases tested for resistance to second-line drugs			198
Laboratory-confirmed cases			MDR/RR-TB: 390, XDR-TB: 1
Patients started on treatment ^d			MDR/RR-TB: 394, XDR-TB: 1

TREATMENT SUCCESS RATE AND COHORT SIZE

	SUCCESS	COHORT
New and relapse cases registered in 2016	81%	75 705
Previously treated cases, excluding relapse, registered in 2016	70%	1 048
HIV-positive TB cases registered in 2016	77%	22 502
MDR/RR-TB cases started on second-line treatment in 2015	73%	330
XDR-TB cases started on second-line treatment in 2015	100%	1

TB PREVENTIVE TREATMENT, 2017

% of HIV-positive people (newly enrolled in care) on preventive treatment	
% of children (aged < 5) household contacts of bacteriologically-confirmed TB cases on preventive treatment	26% (24–29)

TB FINANCING, 2018

National TB budget (US\$ millions)	42
Funding source:	33% domestic, 31% international, 36% unfunded

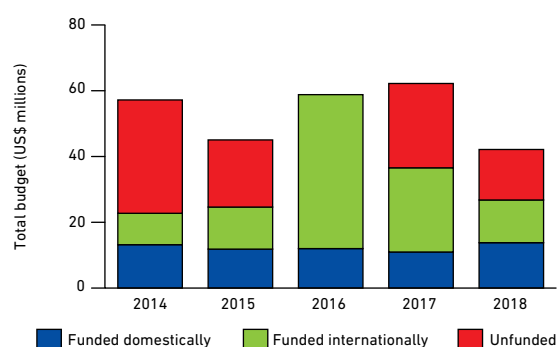
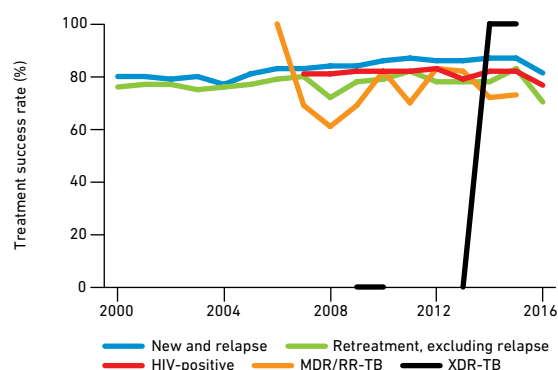
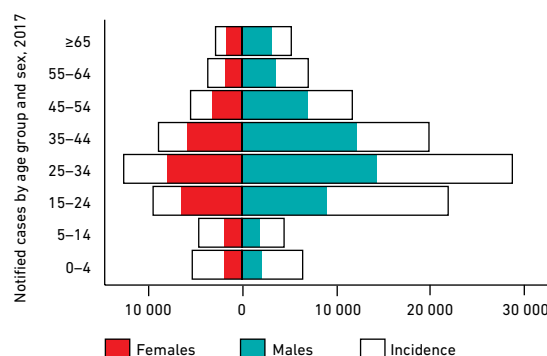
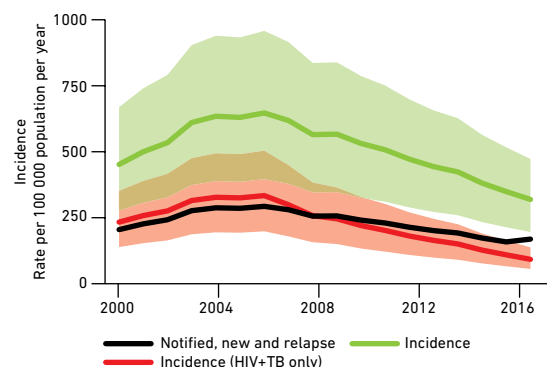
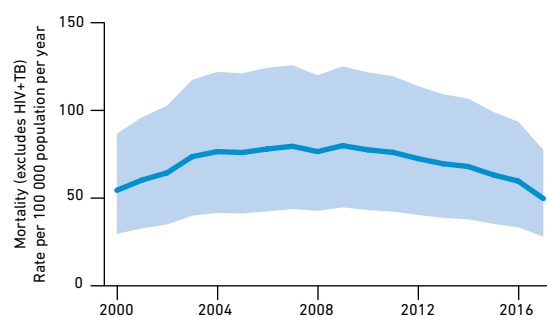
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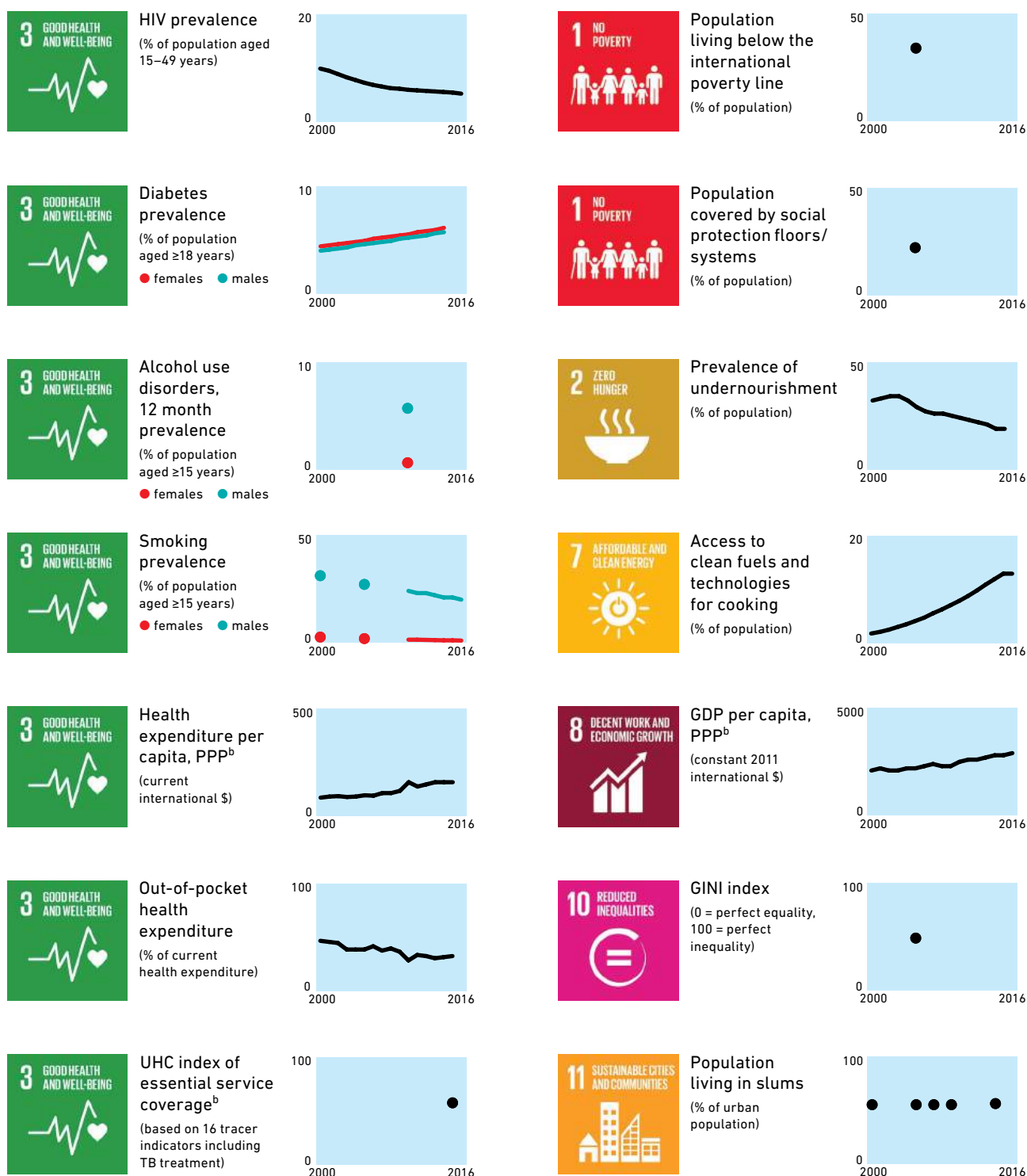
^b MDR is TB resistant to rifampicin and isoniazid; RR is TB resistant to rifampicin.

^c Includes cases with unknown previous TB treatment history.

^d Includes patients diagnosed before 2017 and patients who were not laboratory-confirmed.



INDICATORS IN THE SUSTAINABLE DEVELOPMENT GOALS ASSOCIATED WITH TB INCIDENCE^a



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^b GDP = gross domestic product; PPP = purchasing power parity; UHC = universal health coverage

Mozambique

POPULATION 2017 **30 MILLION**

ESTIMATES OF TB BURDEN,^a 2017

	NUMBER (THOUSANDS)	RATE (PER 100 000 POPULATION)
Mortality (excludes HIV+TB)	22 (13–33)	73 (43–111)
Mortality (HIV+TB only)	27 (17–39)	90 (56–131)
Incidence (includes HIV+TB)	163 (106–233)	551 (356–787)
Incidence (HIV+TB only)	66 (42–95)	221 (141–319)
Incidence (MDR/RR-TB)**	8.8 (4.6–14)	30 (15–48)

ESTIMATED TB INCIDENCE BY AGE AND SEX (THOUSANDS),^a 2017

	0–14 YEARS	> 14 YEARS	TOTAL
Females	11 (9.6–12)	51 (40–63)	62 (47–77)
Males	12 (11–13)	90 (64–115)	101 (70–133)
Total	23 (19–26)	141 (90–192)	163 (106–233)

TB CASE NOTIFICATIONS, 2017

Total cases notified	86 515
Total new and relapse	85 376
— % tested with rapid diagnostics at time of diagnosis	
— % with known HIV status	96%
— % pulmonary	93%
— % bacteriologically confirmed among pulmonary	40%

UNIVERSAL HEALTH COVERAGE AND SOCIAL PROTECTION

TB treatment coverage (notified/estimated incidence), 2017	52% (37–81)
TB patients facing catastrophic total costs	
TB case fatality ratio (estimated mortality/estimated incidence), 2017	0.31 (0.17–0.46)

TB/HIV CARE IN NEW AND RELAPSE TB PATIENTS, 2017

	NUMBER	(%)
Patients with known HIV-status who are HIV-positive	33 514	40%
— on antiretroviral therapy	31 790	95%

DRUG-RESISTANT TB CARE, 2017

	NEW CASES	PREVIOUSLY TREATED CASES	TOTAL NUMBER ^e
Estimated MDR/RR-TB cases among notified pulmonary TB cases			4 100 (2 500–5 700)
Estimated % of TB cases with MDR/RR-TB	3.7% (2.5–5.2)	20% (5.2–40)	
% notified tested for rifampicin resistance			
MDR/RR-TB cases tested for resistance to second-line drugs			209
Laboratory-confirmed cases			MDR/RR-TB: 861, XDR-TB: 31
Patients started on treatment ****			MDR/RR-TB: 907, XDR-TB: 31

TREATMENT SUCCESS RATE AND COHORT SIZE

	SUCCESS	COHORT
New and relapse cases registered in 2016	90%	70 510
Previously treated cases, excluding relapse, registered in 2016	86%	1 593
HIV-positive TB cases registered in 2016	87%	30 572
MDR/RR-TB cases started on second-line treatment in 2015	48%	646
XDR-TB cases started on second-line treatment in 2015	38%	16

TB PREVENTIVE TREATMENT, 2017

% of HIV-positive people (newly enrolled in care) on preventive treatment	
% of children (aged < 5) household contacts of bacteriologically-confirmed TB cases on preventive treatment ^e	> 100%

TB FINANCING, 2018

National TB budget (US\$ millions)	30
Funding source:	5% domestic, 95% international, 0% unfunded

Data are as reported to WHO. Estimates of TB and MDR/RR-TB burden are produced by WHO in consultation with countries. Estimates are rounded and totals are computed prior to rounding.

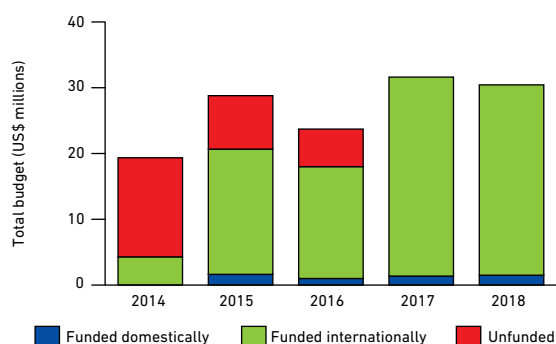
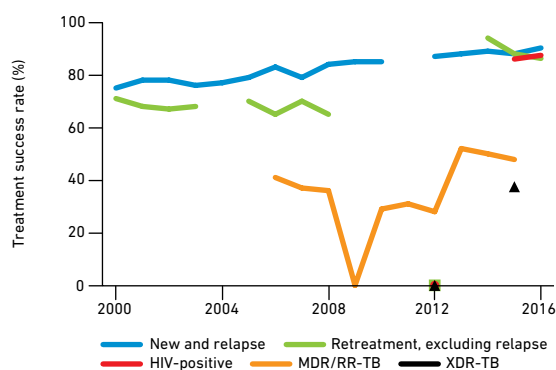
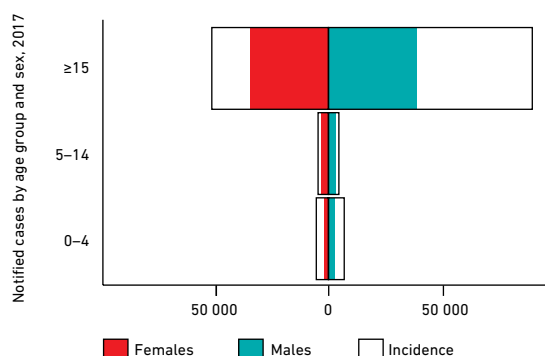
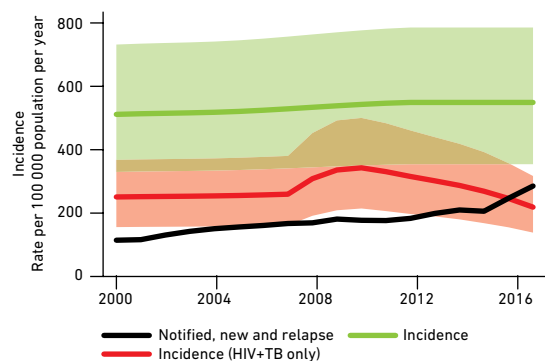
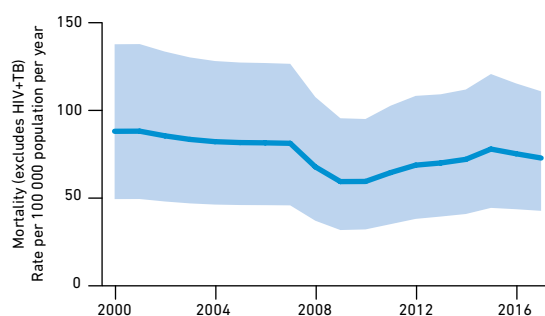
^a Ranges represent uncertainty intervals. Estimates of TB incidence and mortality for Mozambique will be reviewed after final results from their national TB prevalence survey are available in 2019.

^b MDR is TB resistant to rifampicin and isoniazid; RR is TB resistant to rifampicin.

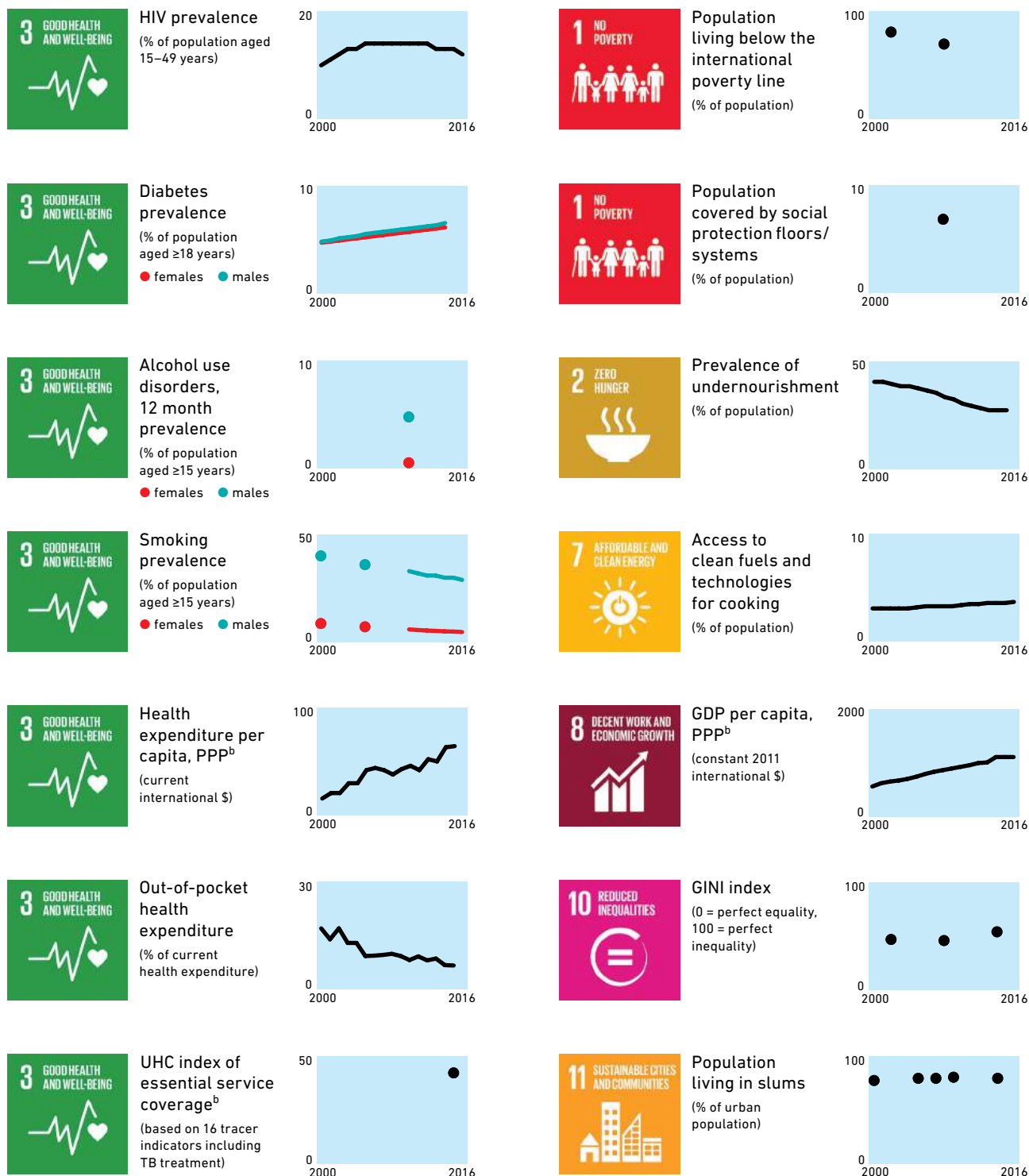
^c Includes cases with unknown previous TB treatment history.

^d Includes patients diagnosed before 2017 and patients who were not laboratory-confirmed.

^e Reasons for higher than expected coverage might be that the numerator did not exclude non-household contacts or children of five years and older.



INDICATORS IN THE SUSTAINABLE DEVELOPMENT GOALS ASSOCIATED WITH TB INCIDENCE^a



Targets for reductions in TB incidence and TB deaths set in WHO's End TB Strategy and the United Nations' Sustainable Development Goals (SDGs) are ambitious. Achieving them requires progress in reducing health-related risk factors for TB infection and disease, as well as broader social and economic determinants of TB infection and disease. WHO has developed a TB-SDG monitoring framework that comprises 14 indicators under seven SDGs for which there is evidence of an association with TB incidence. Further details are provided in [Chapter 2](#).

^a Data sources: SDG indicators database, The World Bank, World Health Organization. Missing values and empty boxes indicate data not available in these data sources.

^b GDP = gross domestic product; PPP = purchasing power parity; UHC = universal health coverage

Myanmar

POPULATION 2017 **53 MILLION**

ESTIMATES OF TB BURDEN,^a 2017

	NUMBER (THOUSANDS)	RATE (PER 100 000 POPULATION)
Mortality (excludes HIV+TB)	27 (18–39)	51 (33–73)
Mortality (HIV+TB only)	4.9 (3.5–6.6)	9.2 (6.6–12)
Incidence (includes HIV+TB)	191 (141–249)	358 (263–466)
Incidence (HIV+TB only)	17 (12–22)	31 (23–41)
Incidence (MDR/RR-TB) ^b	14 (8–21)	26 (15–39)

ESTIMATED TB INCIDENCE BY AGE AND SEX (THOUSANDS),^a 2017

	0–14 YEARS	> 14 YEARS	TOTAL
Females	11 (10–12)	57 (48–66)	68 (57–80)
Males	12 (11–13)	110 (87–134)	123 (95–150)
Total	23 (21–26)	168 (123–212)	191 (141–249)

TB CASE NOTIFICATIONS, 2017

Total cases notified	132 025
Total new and relapse	130 418
— % tested with rapid diagnostics at time of diagnosis	
— % with known HIV status	90%
— % pulmonary	90%
— % bacteriologically confirmed among pulmonary	41%

UNIVERSAL HEALTH COVERAGE AND SOCIAL PROTECTION

TB treatment coverage (notified/estimated incidence), 2017	68% (52–93)
TB patients facing catastrophic total costs, 2015	60% (57–63)
TB case fatality ratio (estimated mortality/estimated incidence), 2017	0.17 (0.11–0.25)

TB/HIV CARE IN NEW AND RELAPSE TB PATIENTS, 2017

	NUMBER	(%)
Patients with known HIV-status who are HIV-positive	10 164	9%
— on antiretroviral therapy	6 371	63%

DRUG-RESISTANT TB CARE, 2017

	NEW CASES	PREVIOUSLY TREATED CASES	TOTAL NUMBER ^c
Estimated MDR/RR-TB cases among notified pulmonary TB cases			8 700 (6 200–11 000)
Estimated % of TB cases with MDR/RR-TB	5.1% (3.4–7.2)	27% (16–40)	
% notified tested for rifampicin resistance	29%	63%	43 548
MDR/RR-TB cases tested for resistance to second-line drugs			165
Laboratory-confirmed cases		MDR/RR-TB: 3 281, XDR-TB: 28	
Patients started on treatment ^d		MDR/RR-TB: 2 666, XDR-TB: 9	

TREATMENT SUCCESS RATE AND COHORT SIZE

	SUCCESS	COHORT
New and relapse cases registered in 2016	88%	136 221
Previously treated cases, excluding relapse, registered in 2016	77%	2 051
HIV-positive TB cases registered in 2016	73%	10 489
MDR/RR-TB cases started on second-line treatment in 2015	80%	2 180
XDR-TB cases started on second-line treatment in 2015	43%	7

TB PREVENTIVE TREATMENT, 2017

% of HIV-positive people (newly enrolled in care) on preventive treatment	17%
% of children (aged < 5) household contacts of bacteriologically-confirmed TB cases on preventive treatment	2.1% (1.9–2.3)

TB FINANCING, 2018

National TB budget (US\$ millions)	58
Funding source:	4% domestic, 52% international, 45% unfunded

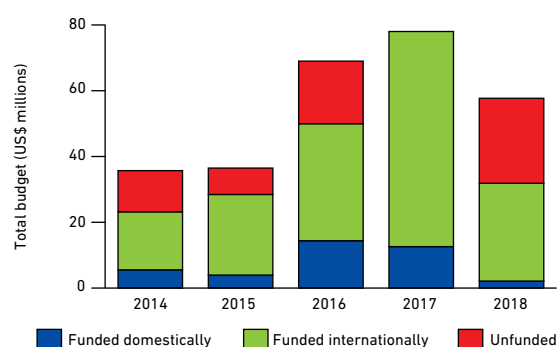
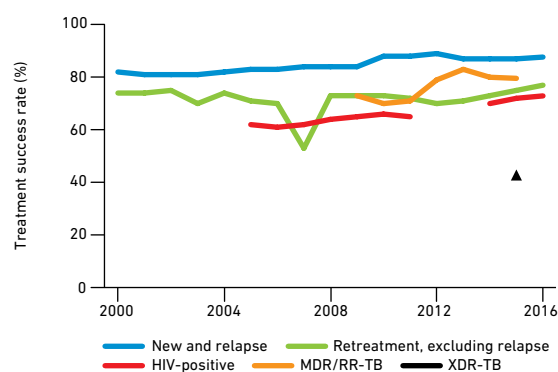
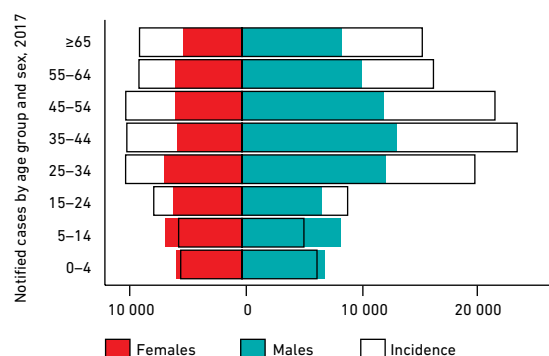
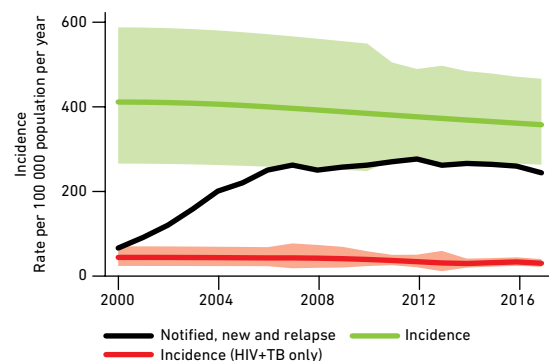
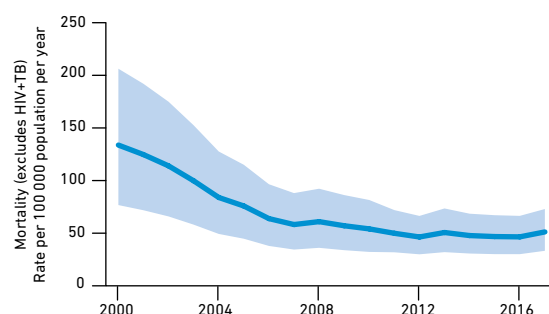
Data are as reported to WHO. Estimates of TB and MDR/RR-TB burden are produced by WHO in consultation with countries. Estimates are rounded and totals are computed prior to rounding.

^a Ranges represent uncertainty intervals. Estimates of TB incidence and mortality for Myanmar will be reviewed after final results from their national TB prevalence survey are available in 2019.

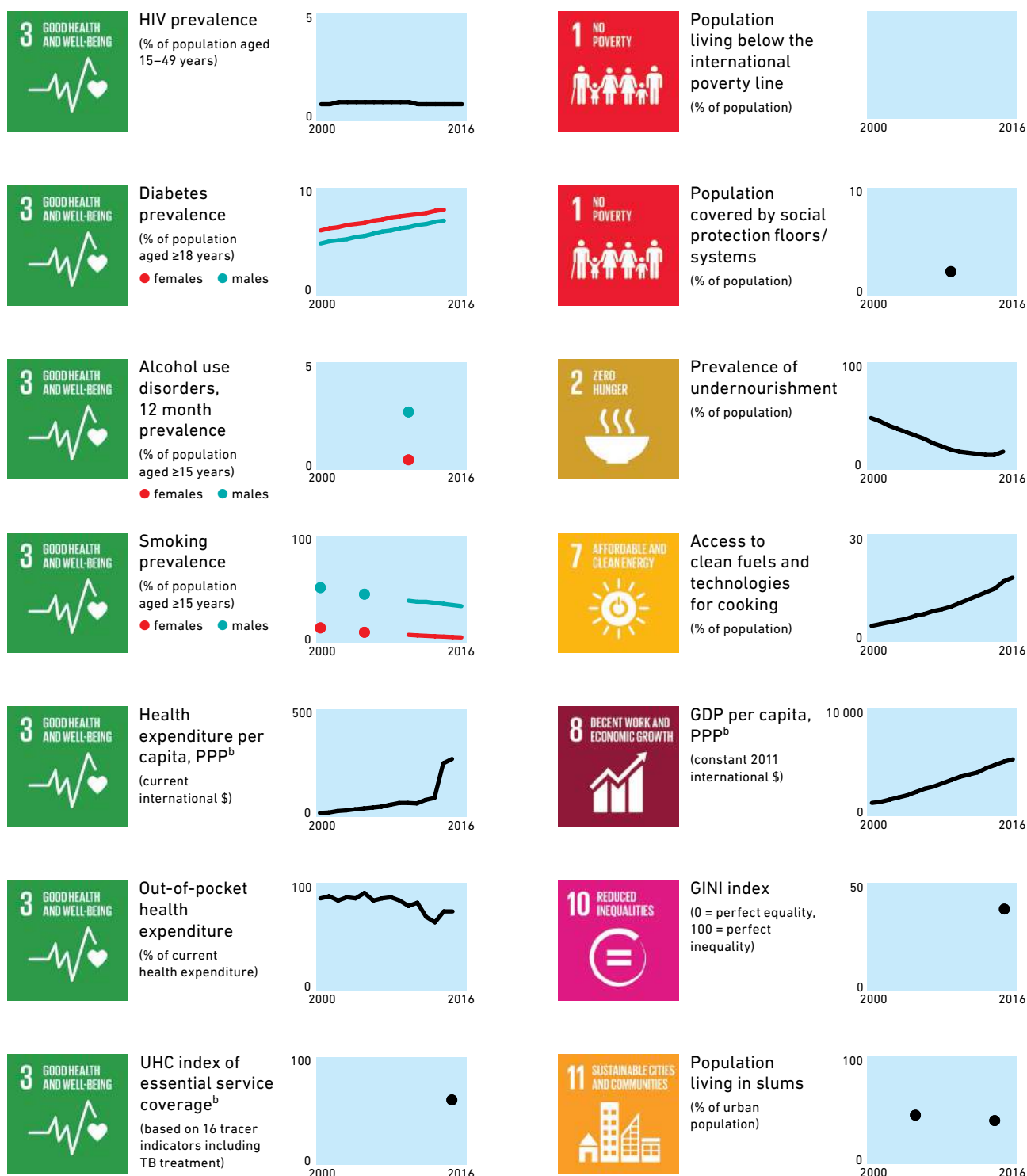
^b MDR is TB resistant to rifampicin and isoniazid; RR is TB resistant to rifampicin.

^c Includes cases with unknown previous TB treatment history.

^d Includes patients diagnosed before 2017 and patients who were not laboratory-confirmed.



INDICATORS IN THE SUSTAINABLE DEVELOPMENT GOALS ASSOCIATED WITH TB INCIDENCE^a



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^b GDP = gross domestic product; PPP = purchasing power parity; UHC = universal health coverage

Nigeria

ESTIMATES OF TB BURDEN, 2017

	NUMBER (THOUSANDS)	RATE (PER 100 000 POPULATION)
Mortality (excludes HIV+TB)	120 (70–183)	63 (36–96)
Mortality (HIV+TB only)	35 (21–52)	18 (11–27)
Incidence (includes HIV+TB)	418 (273–594)	219 (143–311)
Incidence (HIV+TB only)	58 (37–85)	30 (19–44)
Incidence (MDR/RR-TB) ^b	24 (14–36)	12 (7.3–19)

ESTIMATED TB INCIDENCE BY AGE AND SEX (THOUSANDS), 2017

	0–14 YEARS	> 14 YEARS	TOTAL
Females	27 (24–30)	123 (97–148)	150 (115–184)
Males	30 (27–33)	238 (169–307)	268 (186–351)
Total	57 (49–65)	361 (232–490)	418 (273–594)

TB CASE NOTIFICATIONS, 2017

Total cases notified	104 904
Total new and relapse	102 387
— % tested with rapid diagnostics at time of diagnosis	41%
— % with known HIV status	95%
— % pulmonary	96%
— % bacteriologically confirmed among pulmonary	78%

UNIVERSAL HEALTH COVERAGE AND SOCIAL PROTECTION

TB treatment coverage (notified/estimated incidence), 2017	24% (17–38)
TB patients facing catastrophic total costs, 2017	71% (68–73)
TB case fatality ratio (estimated mortality/estimated incidence), 2017	0.38 (0.2–0.59)

TB/HIV CARE IN NEW AND RELAPSE TB PATIENTS, 2017

	NUMBER	(%)
Patients with known HIV-status who are HIV-positive	13 516	14%
— on antiretroviral therapy	11 438	85%

DRUG-RESISTANT TB CARE, 2017

	NEW CASES	PREVIOUSLY TREATED CASES	TOTAL NUMBER ^c
Estimated MDR/RR-TB cases among notified pulmonary TB cases			5 400 (4 200–6 500)
Estimated % of TB cases with MDR/RR-TB	4.3% (3.2–5.5)	25% (19–31)	
% notified tested for rifampicin resistance	41%	61%	43 829
MDR/RR-TB cases tested for resistance to second-line drugs			691
Laboratory-confirmed cases		MDR/RR-TB: 2 286, XDR-TB: 14	
Patients started on treatment ^d		MDR/RR-TB: 1 786, XDR-TB: 10	

TREATMENT SUCCESS RATE AND COHORT SIZE

	SUCCESS	COHORT
New and relapse cases registered in 2016	86%	97 240
Previously treated cases, excluding relapse, registered in 2016	83%	3 193
HIV-positive TB cases registered in 2016	77%	14 794
MDR/RR-TB cases started on second-line treatment in 2015	78%	656
XDR-TB cases started on second-line treatment in 2015		0

TB PREVENTIVE TREATMENT, 2017

% of HIV-positive people (newly enrolled in care) on preventive treatment	39%
% of children (aged < 5) household contacts of bacteriologically-confirmed TB cases on preventive treatment	20% (18–21)

TB FINANCING, 2018

National TB budget (US\$ millions)	409
Funding source:	8% domestic, 16% international, 76% unfunded

Data are as reported to WHO. Estimates of TB and MDR/RR-TB burden are produced by WHO in consultation with countries. Estimates are rounded and totals are computed prior to rounding.

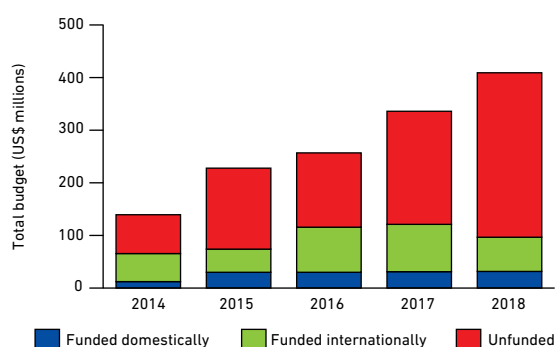
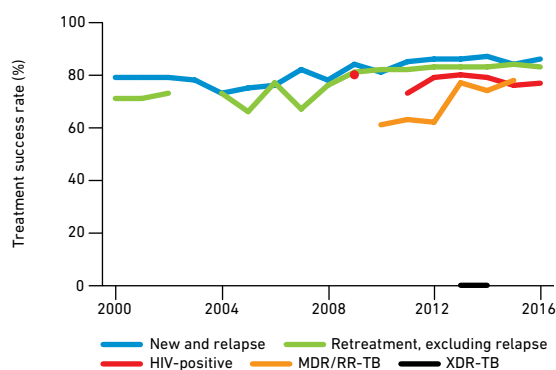
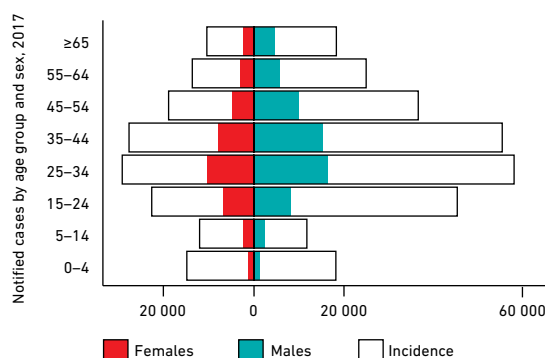
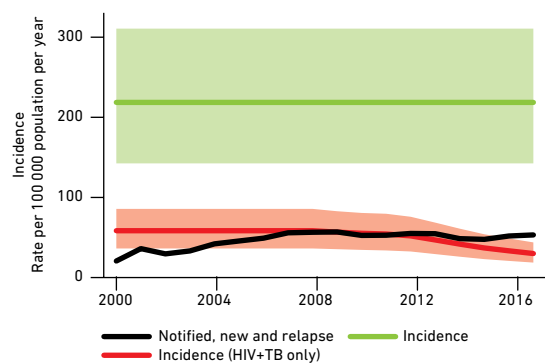
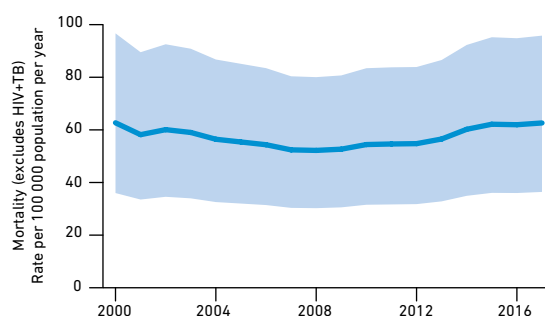
^a Ranges represent uncertainty intervals.

^b MDR is TB resistant to rifampicin and isoniazid; RR is TB resistant to rifampicin.

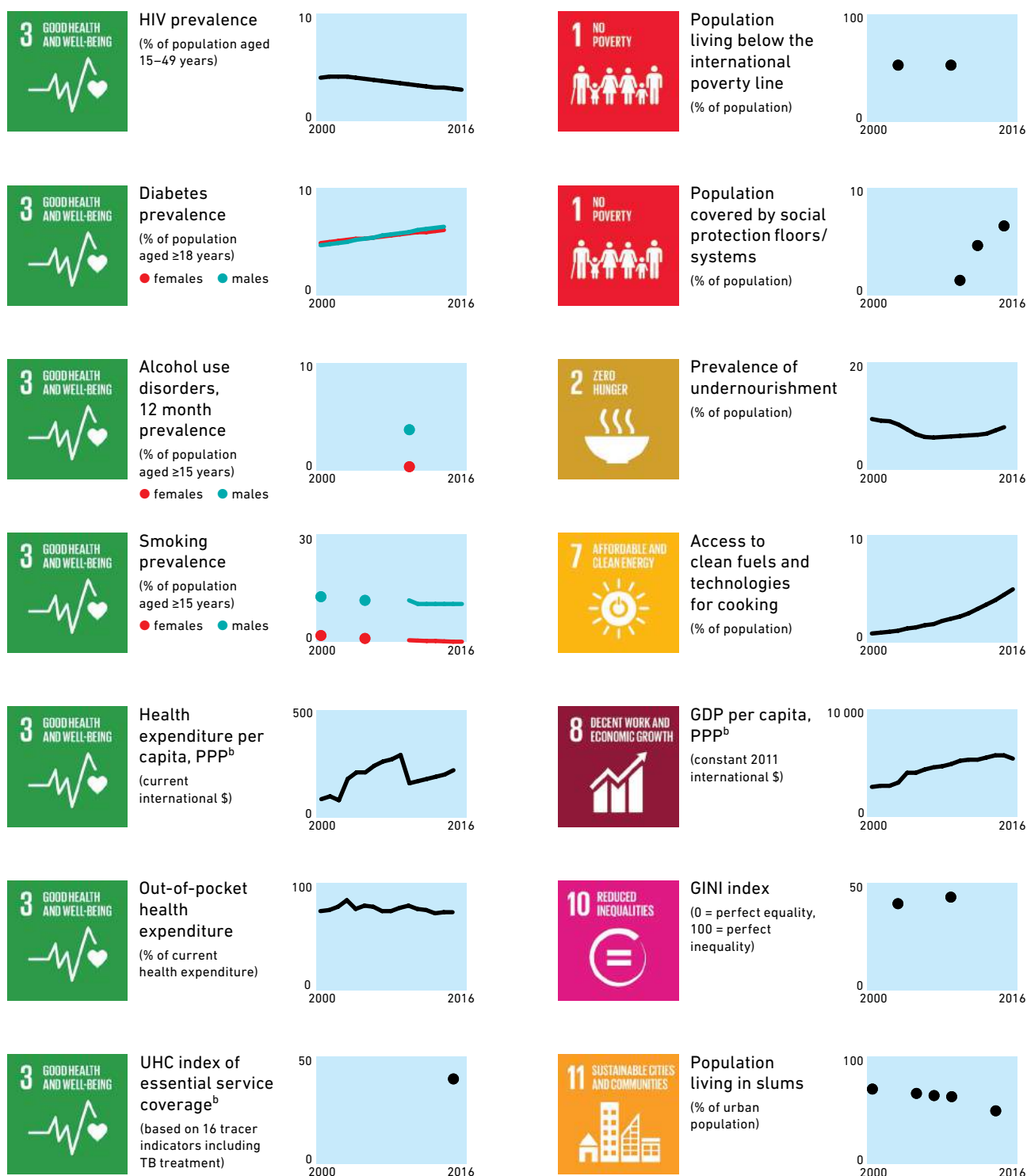
^c Includes cases with unknown previous TB treatment history.

^d Includes patients diagnosed before 2017 and patients who were not laboratory-confirmed.

POPULATION 2017 191 MILLION



INDICATORS IN THE SUSTAINABLE DEVELOPMENT GOALS ASSOCIATED WITH TB INCIDENCE^a



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^b GDP = gross domestic product; PPP = purchasing power parity; UHC = universal health coverage

Pakistan

POPULATION 2017 **197 MILLION**

ESTIMATES OF TB BURDEN,^a 2017

	NUMBER (THOUSANDS)	RATE (PER 100 000 POPULATION)
Mortality (excludes HIV+TB)	54 (42–67)	27 (21–34)
Mortality (HIV+TB only)	2.2 (1.1–3.8)	1.1 (0.56–1.9)
Incidence (includes HIV+TB)	525 (373–704)	267 (189–357)
Incidence (HIV+TB only)	7.3 (3.6–12)	3.7 (1.8–6.2)
Incidence (MDR/RR-TB) ^b	27 (17–39)	14 (8.8–20)

ESTIMATED TB INCIDENCE BY AGE AND SEX (THOUSANDS),^a 2017

	0–14 YEARS	> 14 YEARS	TOTAL
Females	27 (25–29)	207 (166–248)	235 (185–284)
Males	30 (28–32)	261 (203–319)	291 (223–359)
Total	57 (51–63)	468 (329–607)	525 (373–704)

TB CASE NOTIFICATIONS, 2017

Total cases notified	368 897
Total new and relapse	359 224
— % tested with rapid diagnostics at time of diagnosis	3%
— % with known HIV status	7%
— % pulmonary	80%
— % bacteriologically confirmed among pulmonary	48%

UNIVERSAL HEALTH COVERAGE AND SOCIAL PROTECTION

TB treatment coverage (notified/estimated incidence), 2017	68% (51–96)
TB patients facing catastrophic total costs	
TB case fatality ratio (estimated mortality/estimated incidence), 2017	0.11 (0.07–0.15)

TB/HIV CARE IN NEW AND RELAPSE TB PATIENTS, 2017

	NUMBER	(%)
Patients with known HIV-status who are HIV-positive	121	<1%
— on antiretroviral therapy	97	80%

DRUG-RESISTANT TB CARE, 2017

	NEW CASES	PREVIOUSLY TREATED CASES	TOTAL NUMBER ^c
Estimated MDR/RR-TB cases among notified pulmonary TB cases			15 000 (12 000–18 000)
Estimated % of TB cases with MDR/RR-TB	4.2% (3.2–5.3)	16% (15–17)	
% notified tested for rifampicin resistance	11%	47%	54 991
MDR/RR-TB cases tested for resistance to second-line drugs			2 887
Laboratory-confirmed cases		MDR/RR-TB: 3 475, XDR-TB: 128	
Patients started on treatment ^d		MDR/RR-TB: 3 016, XDR-TB: 65	

TREATMENT SUCCESS RATE AND COHORT SIZE

	Success	Cohort
New and relapse cases registered in 2016	94%	356 390
Previously treated cases, excluding relapse, registered in 2016	78%	9 671
HIV-positive TB cases registered in 2016		
MDR/RR-TB cases started on second-line treatment in 2015	64%	2 544
XDR-TB cases started on second-line treatment in 2015	29%	77

TB PREVENTIVE TREATMENT, 2017

% of HIV-positive people (newly enrolled in care) on preventive treatment	
% of children (aged < 5) household contacts of bacteriologically-confirmed TB cases on preventive treatment	

TB FINANCING, 2018

National TB budget (US\$ millions)	118
Funding source:	3% domestic, 54% international, 43% unfunded

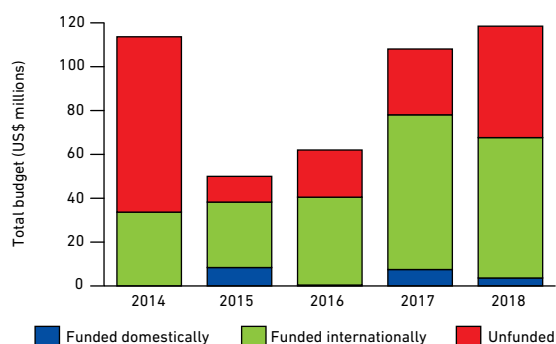
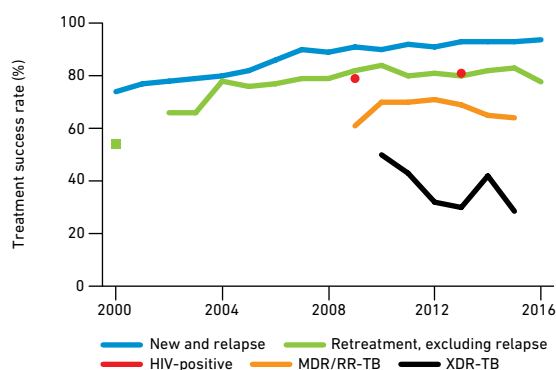
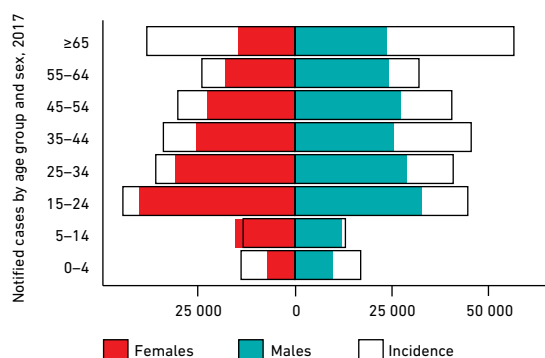
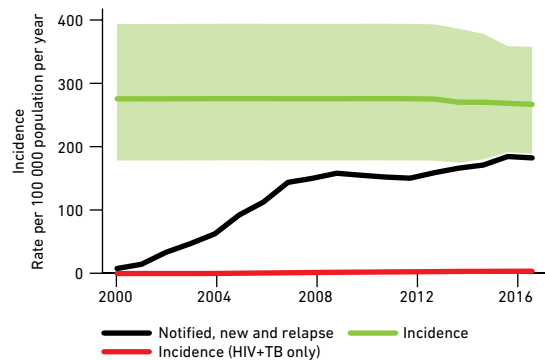
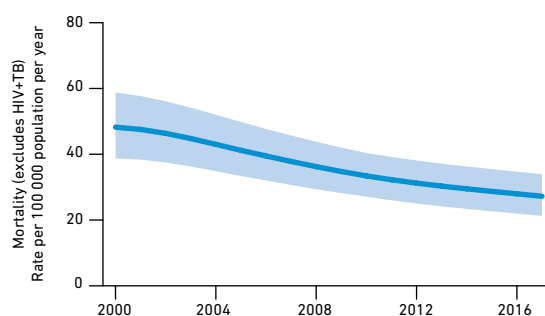
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^a Ranges represent uncertainty intervals.

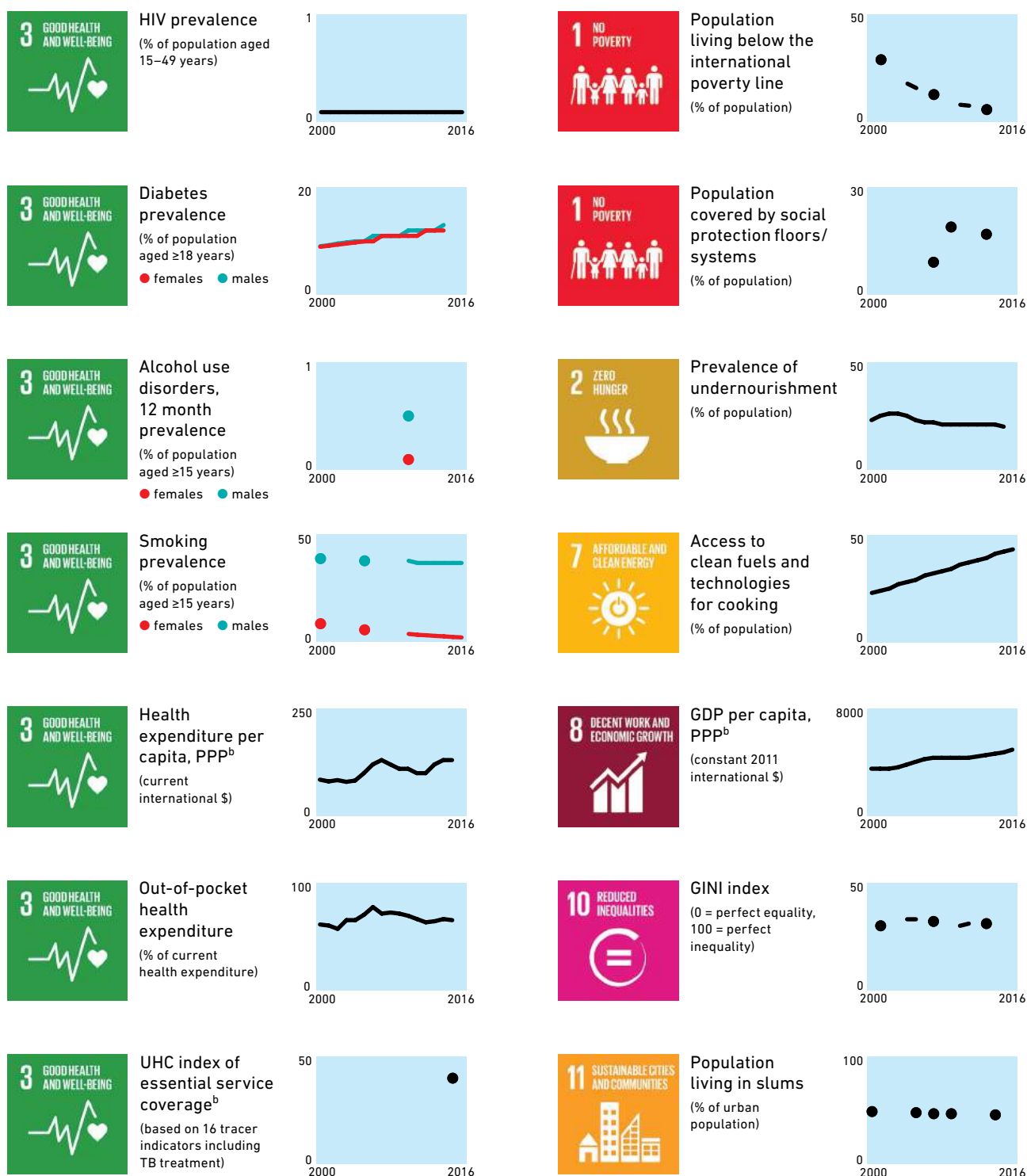
^b MDR is TB resistant to rifampicin and isoniazid; RR is TB resistant to rifampicin.

^c Includes cases with unknown previous TB treatment history.

^d Includes patients diagnosed before 2017 and patients who were not laboratory-confirmed.



INDICATORS IN THE SUSTAINABLE DEVELOPMENT GOALS ASSOCIATED WITH TB INCIDENCE^a



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^b GDP = gross domestic product; PPP = purchasing power parity; UHC = universal health coverage

Philippines

POPULATION 2017 **105 MILLION**

ESTIMATES OF TB BURDEN, ^a 2017

	NUMBER (THOUSANDS)	RATE (PER 100 000 POPULATION)
Mortality (excludes HIV+TB)	26 (23–31)	25 (22–29)
Mortality (HIV+TB only)	0.38 (0–3.3)	0.36 (0–3.1)
Incidence (includes HIV+TB)	581 (326–909)	554 (311–866)
Incidence (HIV+TB only)	7.1 (2.9–13)	6.7 (2.8–12)
Incidence (MDR/RR-TB) ^b	27 (12–47)	26 (12–45)

ESTIMATED TB INCIDENCE BY AGE AND SEX (THOUSANDS), ^a 2017

	0–14 YEARS	> 14 YEARS	TOTAL
Females	34 (30–38)	139 (105–173)	173 (126–220)
Males	37 (33–42)	371 (223–519)	408 (237–580)
Total	71 (59–84)	510 (271–749)	581 (326–909)

TB CASE NOTIFICATIONS, 2017

Total cases notified	328 773
Total new and relapse	317 266
— % tested with rapid diagnostics at time of diagnosis	26%
— % with known HIV status	24%
— % pulmonary	98%
— % bacteriologically confirmed among pulmonary	39%

UNIVERSAL HEALTH COVERAGE AND SOCIAL PROTECTION

TB treatment coverage (notified/estimated incidence), 2017	55% (35–97)
TB patients facing catastrophic total costs, 2017	35%
TB case fatality ratio (estimated mortality/estimated incidence), 2017	0.05 (0.03–0.08)

TB/HIV CARE IN NEW AND RELAPSE TB PATIENTS, 2017

	NUMBER	(%)
Patients with known HIV-status who are HIV-positive	1 335	2%
— on antiretroviral therapy	1 185	89%

DRUG-RESISTANT TB CARE, 2017

	NEW CASES	PREVIOUSLY TREATED CASES	TOTAL NUMBER ^c
Estimated MDR/RR-TB cases among notified pulmonary TB cases			20 000 (18 000–22 000)
Estimated % of TB cases with MDR/RR-TB	2.6% (1.9–3.4)	28% (27–28)	
% notified tested for rifampicin resistance	19%	83%	93 058
MDR/RR-TB cases tested for resistance to second-line drugs			2 401
Laboratory-confirmed cases		MDR/RR-TB: 6 438, XDR-TB: 15	
Patients started on treatment ^d		MDR/RR-TB: 5 623, XDR-TB: 16	

TREATMENT SUCCESS RATE AND COHORT SIZE

	SUCCESS	COHORT
New and relapse cases registered in 2016	91%	332 308
Previously treated cases, excluding relapse, registered in 2016	80%	10 097
HIV-positive TB cases registered in 2016	82%	989
MDR/RR-TB cases started on second-line treatment in 2015	54%	3 851
XDR-TB cases started on second-line treatment in 2015	33%	9

TB PREVENTIVE TREATMENT, 2017

% of HIV-positive people (newly enrolled in care) on preventive treatment	57%
% of children (aged < 5) household contacts of bacteriologically-confirmed TB cases on preventive treatment	12% (11–13)

TB FINANCING, 2018

National TB budget (US\$ millions)	160
Funding source:	37% domestic, 21% international, 42% unfunded

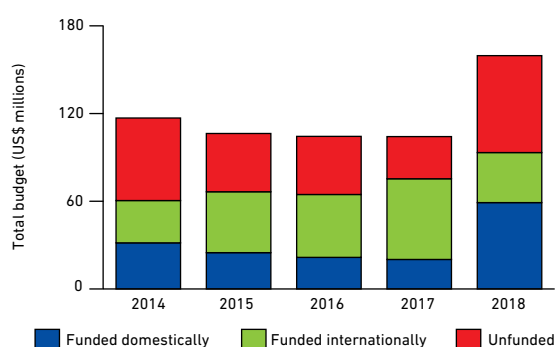
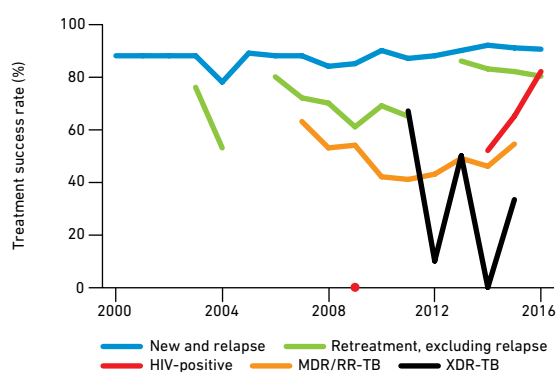
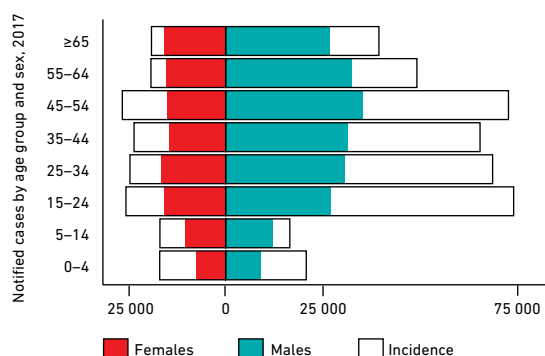
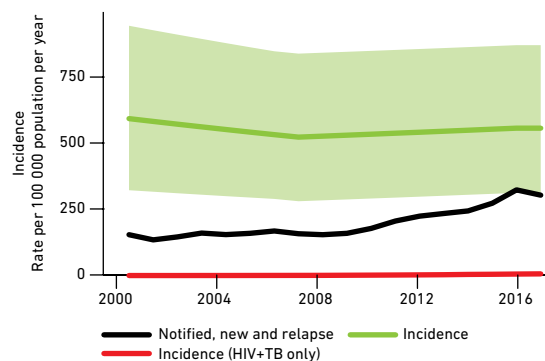
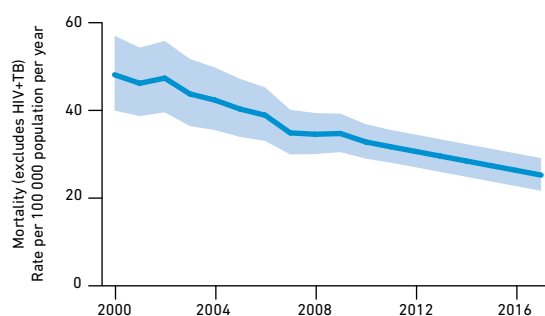
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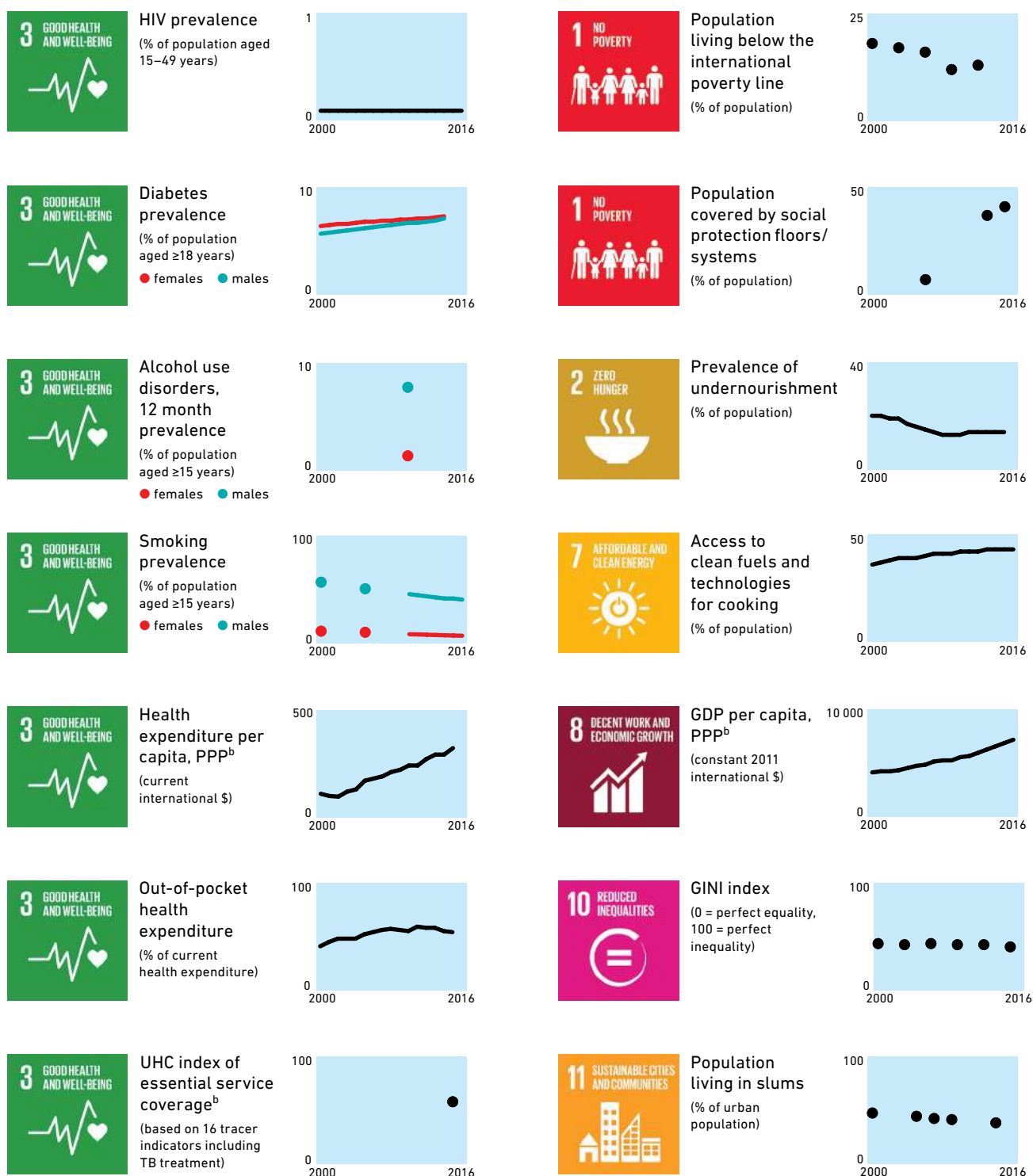
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^d Includes patients diagnosed before 2017 and patients who were not laboratory-confirmed.



INDICATORS IN THE SUSTAINABLE DEVELOPMENT GOALS ASSOCIATED WITH TB INCIDENCE^a



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^b GDP = gross domestic product; PPP = purchasing power parity; UHC = universal health coverage

Russian Federation

POPULATION 2017^a 144 MILLION

ESTIMATES OF TB BURDEN, ^b 2017

	NUMBER (THOUSANDS)	RATE (PER 100 000 POPULATION)
Mortality (excludes HIV+TB)	10 (9.4–12)	7.3 (6.6–8)
Mortality (HIV+TB only)	1.7 (0.85–2.8)	1.2 (0.59–1.9)
Incidence (includes HIV+TB)	86 (56–123)	60 (39–85)
Incidence (HIV+TB only)	18 (12–26)	13 (8.3–18)
Incidence (MDR/RR-TB) ^c	56 (36–82)	39 (25–57)

ESTIMATED TB INCIDENCE BY AGE AND SEX (THOUSANDS), ^b 2017

	0–14 YEARS	> 14 YEARS	TOTAL
Females	1.4 (1.3–1.4)	25 (20–30)	26 (21–32)
Males	1.2 (1.1–1.3)	59 (40–78)	60 (40–79)
Total	2.6 (2.4–2.7)	84 (51–116)	86 (56–123)

TB CASE NOTIFICATIONS, 2017

Total cases notified	114 187
Total new and relapse	84 510
— % tested with rapid diagnostics at time of diagnosis	78%
— % with known HIV status	96%
— % pulmonary	92%
— % bacteriologically confirmed among pulmonary	52%

UNIVERSAL HEALTH COVERAGE AND SOCIAL PROTECTION

TB treatment coverage (notified/estimated incidence), 2017	98% (69–150)
TB patients facing catastrophic total costs	
TB case fatality ratio (estimated mortality/estimated incidence), 2017	0.15 (0.09–0.21)

TB/HIV CARE IN NEW AND RELAPSE TB PATIENTS, 2017

	NUMBER	(%)
Patients with known HIV-status who are HIV-positive	15 695	19%
— on antiretroviral therapy	9 748	62%

DRUG-RESISTANT TB CARE, 2017

	NEW CASES	PREVIOUSLY TREATED CASES	TOTAL NUMBER ^d
Estimated MDR/RR-TB cases among notified pulmonary TB cases			49 000 (49 000–50 000)
Estimated % of TB cases with MDR/RR-TB	32% (31–33)	67% (66–67)	
% notified tested for rifampicin resistance	42%	58%	55 344
MDR/RR-TB cases tested for resistance to second-line drugs			23 098
Laboratory-confirmed cases		MDR/RR-TB: 26 602, XDR-TB: 3 661	
Patients started on treatment ^e		MDR/RR-TB: 26 457, XDR-TB: 2 770	

TREATMENT SUCCESS RATE AND COHORT SIZE

	SUCCESS	COHORT
New and relapse cases registered in 2016	72%	73 137
Previously treated cases, excluding relapse, registered in 2016	49%	9 245
HIV-positive TB cases registered in 2016	53%	1 328
MDR/RR-TB cases started on second-line treatment in 2015	54%	22 607
XDR-TB cases started on second-line treatment in 2015	33%	2 882

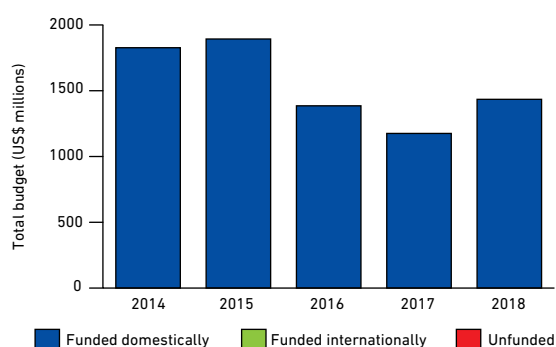
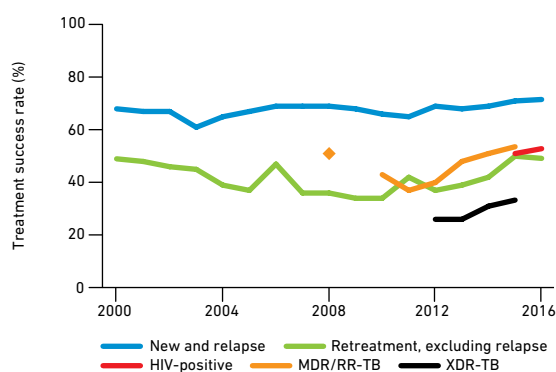
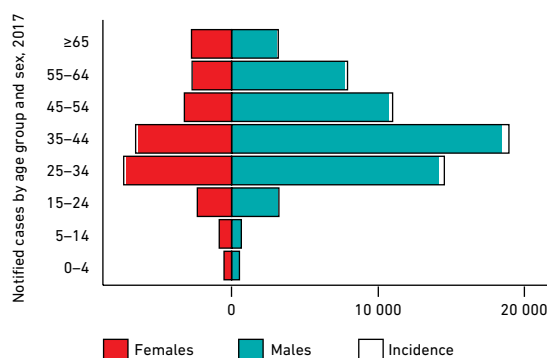
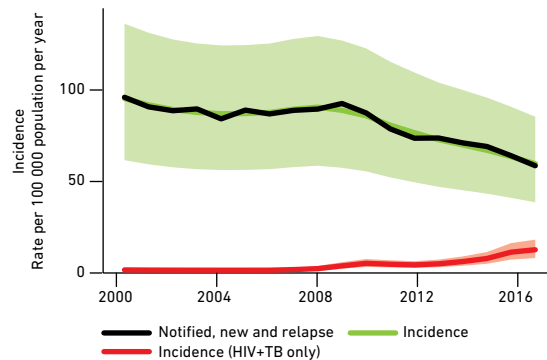
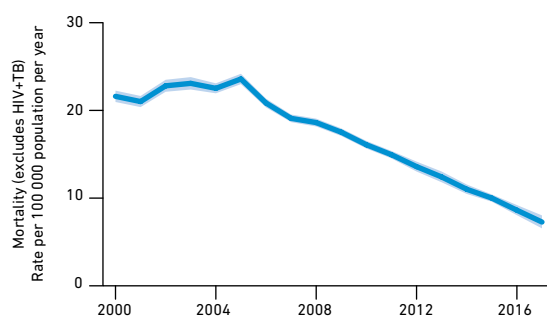
TB PREVENTIVE TREATMENT, 2017

% of HIV-positive people (newly enrolled in care) on preventive treatment	97%
% of children (aged < 5) household contacts of bacteriologically-confirmed TB cases on preventive treatment ^f	> 100%

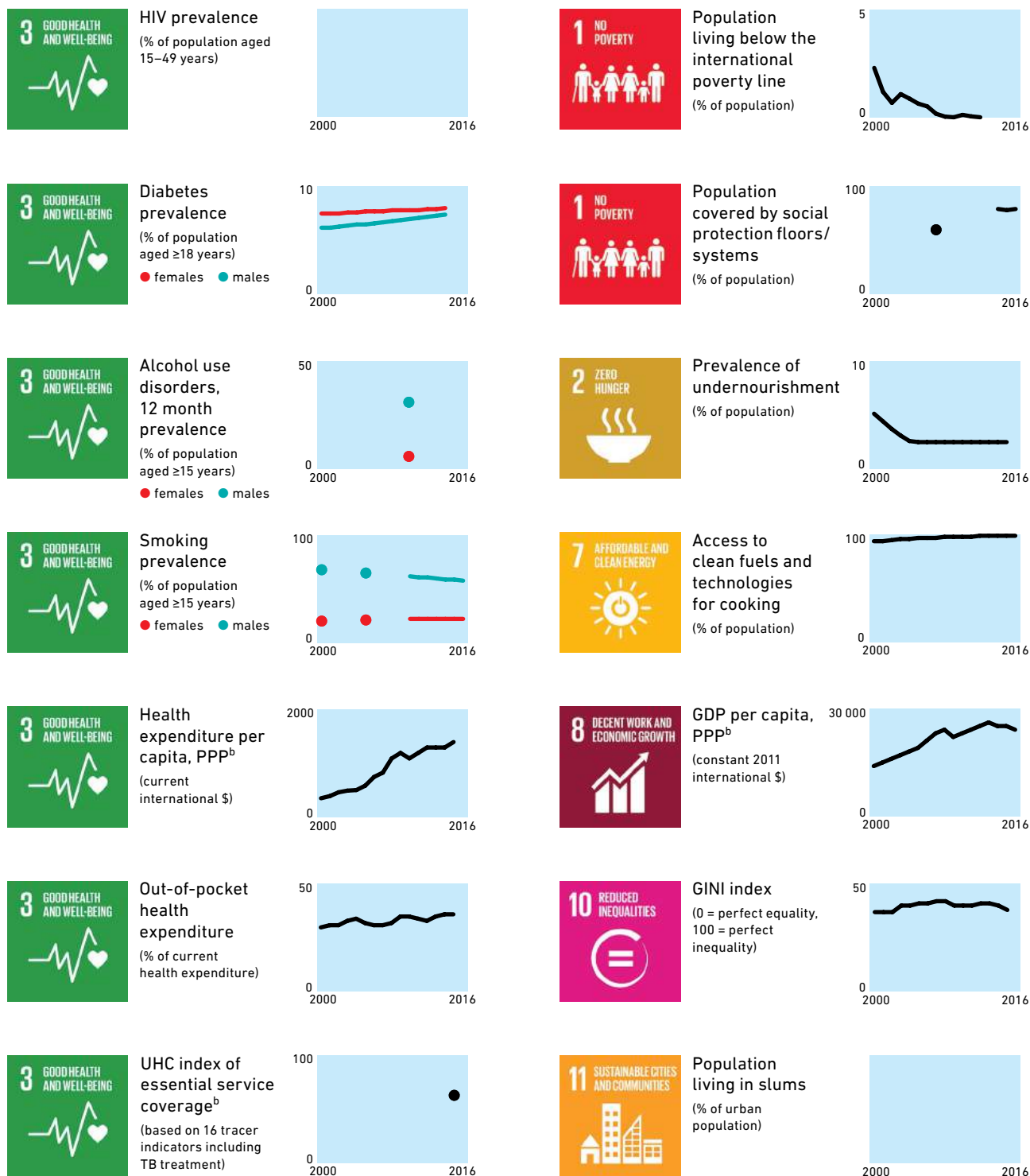
TB FINANCING, 2018

National TB budget (US\$ millions)	1 435
Funding source:	100% domestic, 0% international, 0% unfunded

Data are as reported to WHO. Estimates of TB and MDR/RR-TB burden are produced by WHO in consultation with countries. Estimates are rounded and totals are computed prior to rounding.
^a UN Population Division estimates are lower than the population registered by the Federal State Statistics Service of the Russian Federation.
^b Ranges represent uncertainty intervals.
^c MDR is TB resistant to rifampicin and isoniazid; RR is TB resistant to rifampicin.
^d Includes cases with unknown previous TB treatment history.
^e Includes patients diagnosed before 2017 and patients who were not laboratory-confirmed.
^f Reasons for higher than expected coverage might be that the numerator did not exclude non-household contacts or children of five years and older.



INDICATORS IN THE SUSTAINABLE DEVELOPMENT GOALS ASSOCIATED WITH TB INCIDENCE^a



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^a Data sources: SDG indicators database, The World Bank, World Health Organization. Missing values and empty boxes indicate data not available in these data sources.

^b GDP = gross domestic product; PPP = purchasing power parity; UHC = universal health coverage

South Africa

POPULATION 2017 **57 MILLION**

ESTIMATES OF TB BURDEN,^a 2017

	NUMBER (THOUSANDS)	RATE (PER 100 000 POPULATION)
Mortality (excludes HIV+TB)	22 (20–24)	39 (35–43)
Mortality (HIV+TB only)	56 (39–77)	99 (68–135)
Incidence (includes HIV+TB)	322 (230–428)	567 (406–754)
Incidence (HIV+TB only)	193 (137–258)	340 (241–455)
Incidence (MDR/RR-TB) ^b	14 (8.9–20)	25 (16–36)

ESTIMATED TB INCIDENCE BY AGE AND SEX (THOUSANDS),^a 2017

	0–14 YEARS	> 14 YEARS	TOTAL
Females	18 (17–19)	117 (95–138)	135 (108–161)
Males	20 (18–21)	167 (130–204)	187 (143–231)
Total	38 (34–42)	284 (202–366)	322 (230–428)

TB CASE NOTIFICATIONS, 2017

Total cases notified	227 224
Total new and relapse	220 163
— % tested with rapid diagnostics at time of diagnosis	66%
— % with known HIV status	94%
— % pulmonary	89%
— % bacteriologically confirmed among pulmonary	65%

UNIVERSAL HEALTH COVERAGE AND SOCIAL PROTECTION

TB treatment coverage (notified/estimated incidence), 2017	68% (51–96)
TB patients facing catastrophic total costs	
TB case fatality ratio (estimated mortality/estimated incidence), 2017	0.25 (0.16–0.35)

TB/HIV CARE IN NEW AND RELAPSE TB PATIENTS, 2017

	NUMBER	(%)
Patients with known HIV-status who are HIV-positive	123 148	60%
— on antiretroviral therapy	109 799	89%

DRUG-RESISTANT TB CARE, 2017

	NEW CASES	PREVIOUSLY TREATED CASES	TOTAL NUMBER ^c
Estimated MDR/RR-TB cases among notified pulmonary TB cases			7 700 (6 000–9 400)
Estimated % of TB cases with MDR/RR-TB	3.4% (2.5–4.3)	7.1% (4.8–9.5)	
% notified tested for rifampicin resistance	64%	68%	150 548
MDR/RR-TB cases tested for resistance to second-line drugs			
Laboratory-confirmed cases		MDR/RR-TB: 15 986, XDR-TB: 747	
Patients started on treatment ^d		MDR/RR-TB: 10 259, XDR-TB: 463	

TREATMENT SUCCESS RATE AND COHORT SIZE

	SUCCESS	COHORT
New and relapse cases registered in 2016	82%	236 702
Previously treated cases, excluding relapse, registered in 2016	62%	5 129
HIV-positive TB cases registered in 2016	80%	133 710
MDR/RR-TB cases started on second-line treatment in 2015	55%	9 750
XDR-TB cases started on second-line treatment in 2015	48%	427

TB PREVENTIVE TREATMENT, 2017

% of HIV-positive people (newly enrolled in care) on preventive treatment	53%
% of children (aged < 5) household contacts of bacteriologically-confirmed TB cases on preventive treatment	79% (72–86)

TB FINANCING, 2018

National TB budget (US\$ millions)	271
Funding source:	91% domestic, 8% international, <1% unfunded

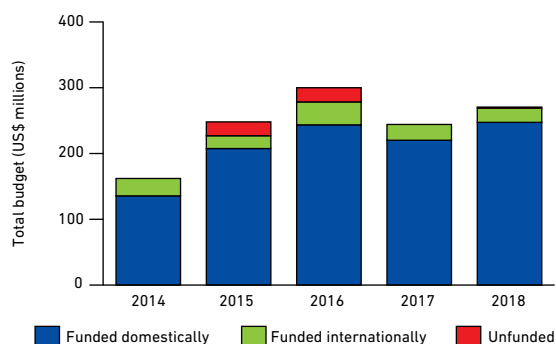
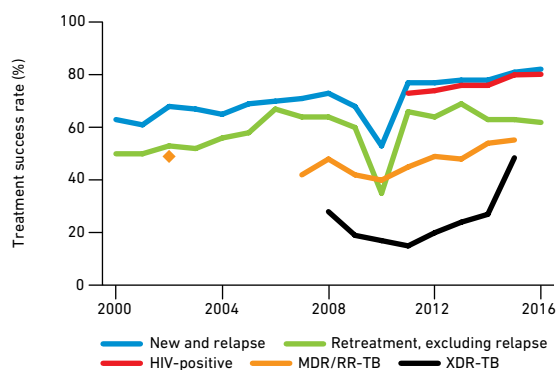
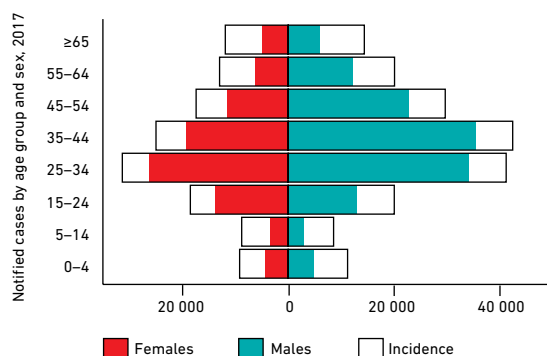
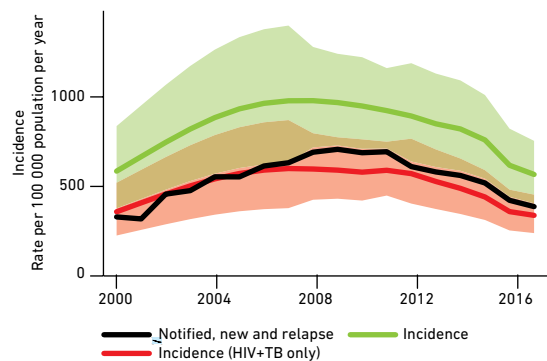
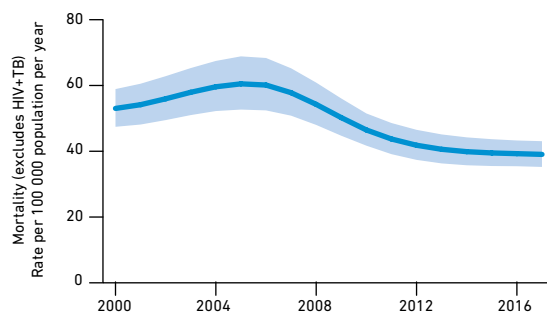
Data are as reported to WHO. Estimates of TB and MDR/RR-TB burden are produced by WHO in consultation with countries. Estimates are rounded and totals are computed prior to rounding.

^a Ranges represent uncertainty intervals. Estimates of TB incidence and mortality for South Africa will be reviewed after final results from their national TB prevalence survey are available in 2019.

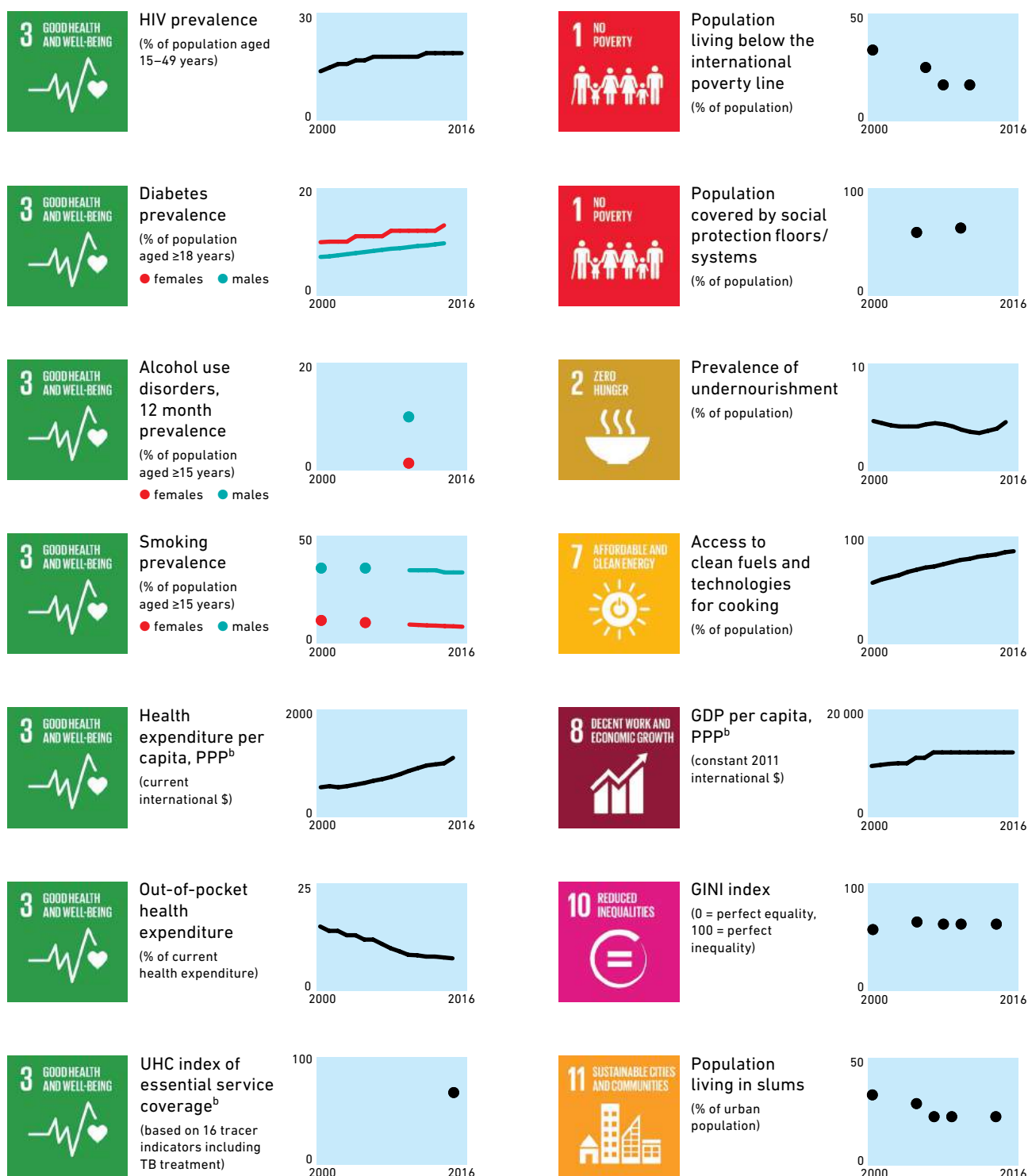
^b MDR is TB resistant to rifampicin and isoniazid; RR is TB resistant to rifampicin.

^c Includes cases with unknown previous TB treatment history.

^d Includes patients diagnosed before 2017 and patients who were not laboratory-confirmed.



INDICATORS IN THE SUSTAINABLE DEVELOPMENT GOALS ASSOCIATED WITH TB INCIDENCE^a



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^b GDP = gross domestic product; PPP = purchasing power parity; UHC = universal health coverage

Thailand

POPULATION 2017 **69 MILLION**

ESTIMATES OF TB BURDEN,^a 2017

	NUMBER (THOUSANDS)	RATE (PER 100 000 POPULATION)
Mortality (excludes HIV+TB)	9.3 (7–12)	13 (10–17)
Mortality (HIV+TB only)	2.9 (2.1–3.8)	4.2 (3.1–5.6)
Incidence (includes HIV+TB)	108 (82–138)	156 (119–199)
Incidence (HIV+TB only)	11 (8.5–15)	16 (12–21)
Incidence (MDR/RR-TB)**	3.9 (2.5–5.7)	5.7 (3.6–8.2)

ESTIMATED TB INCIDENCE BY AGE AND SEX (THOUSANDS),^a 2017

	0–14 YEARS	> 14 YEARS	TOTAL
Females	4.1 (3.9–4.3)	33 (28–37)	37 (31–42)
Males	4.4 (4.2–4.7)	67 (53–81)	71 (56–86)
Total	8.5 (7.9–9.1)	100 (75–124)	108 (82–138)

TB CASE NOTIFICATIONS, 2017

Total cases notified	82 008
Total new and relapse	80 160
— % tested with rapid diagnostics at time of diagnosis	12%
— % with known HIV status	82%
— % pulmonary	83%
— % bacteriologically confirmed among pulmonary	55%

UNIVERSAL HEALTH COVERAGE AND SOCIAL PROTECTION

TB treatment coverage (notified/estimated incidence), 2017	74% (58–98)
TB patients facing catastrophic total costs	
TB case fatality ratio (estimated mortality/estimated incidence), 2017	0.11 (0.08–0.16)

TB/HIV CARE IN NEW AND RELAPSE TB PATIENTS, 2017

	NUMBER	(%)
Patients with known HIV-status who are HIV-positive	7 130	11%
— on antiretroviral therapy	4 577	64%

DRUG-RESISTANT TB CARE, 2017

	NEW CASES	PREVIOUSLY TREATED CASES	TOTAL NUMBER ^e
Estimated MDR/RR-TB cases among notified pulmonary TB cases			2 700 (2 100–3 300)
Estimated % of TB cases with MDR/RR-TB	2.2% (1.5–3)	24% (18–31)	
% notified tested for rifampicin resistance	24%	37%	24 470
MDR/RR-TB cases tested for resistance to second-line drugs			272
Laboratory-confirmed cases			MDR/RR-TB: 1 339, XDR-TB: 7
Patients started on treatment ^d			MDR/RR-TB: 851, XDR-TB: 8

TREATMENT SUCCESS RATE AND COHORT SIZE

	SUCCESS	COHORT
New and relapse cases registered in 2016	83%	68 146
Previously treated cases, excluding relapse, registered in 2016	71%	3 806
HIV-positive TB cases registered in 2016	72%	6 552
MDR/RR-TB cases started on second-line treatment in 2015	60%	352
XDR-TB cases started on second-line treatment in 2015		

TB PREVENTIVE TREATMENT, 2017

% of HIV-positive people (newly enrolled in care) on preventive treatment	
% of children (aged < 5) household contacts of bacteriologically-confirmed TB cases on preventive treatment	5% (4.6–5.5)

TB FINANCING, 2018

National TB budget (US\$ millions)	26
Funding source:	87% domestic, 13% international, 0% unfunded

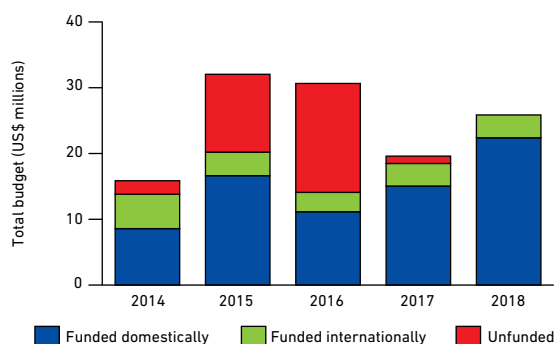
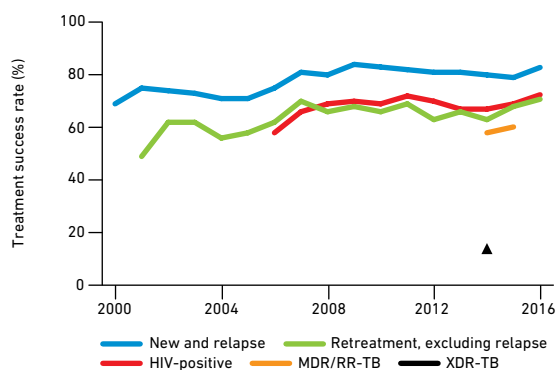
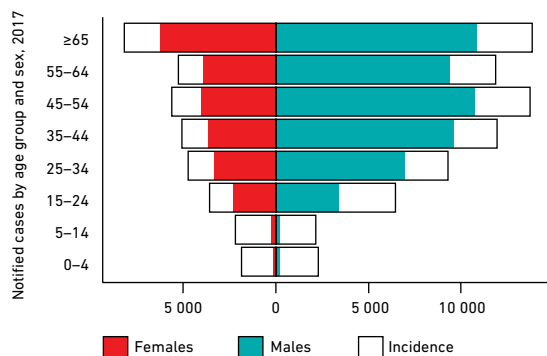
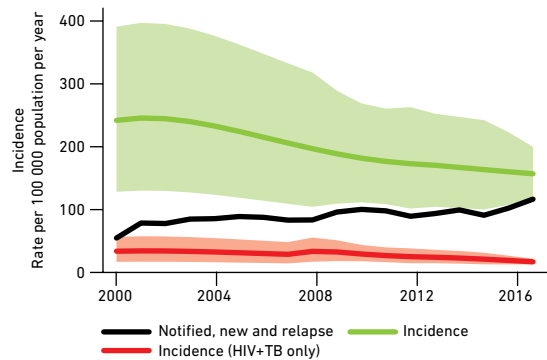
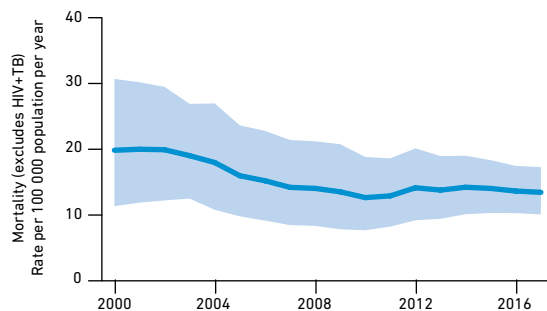
Data are as reported to WHO. Estimates of TB and MDR/RR-TB burden are produced by WHO in consultation with countries. Estimates are rounded and totals are computed prior to rounding.

^a Ranges represent uncertainty intervals.

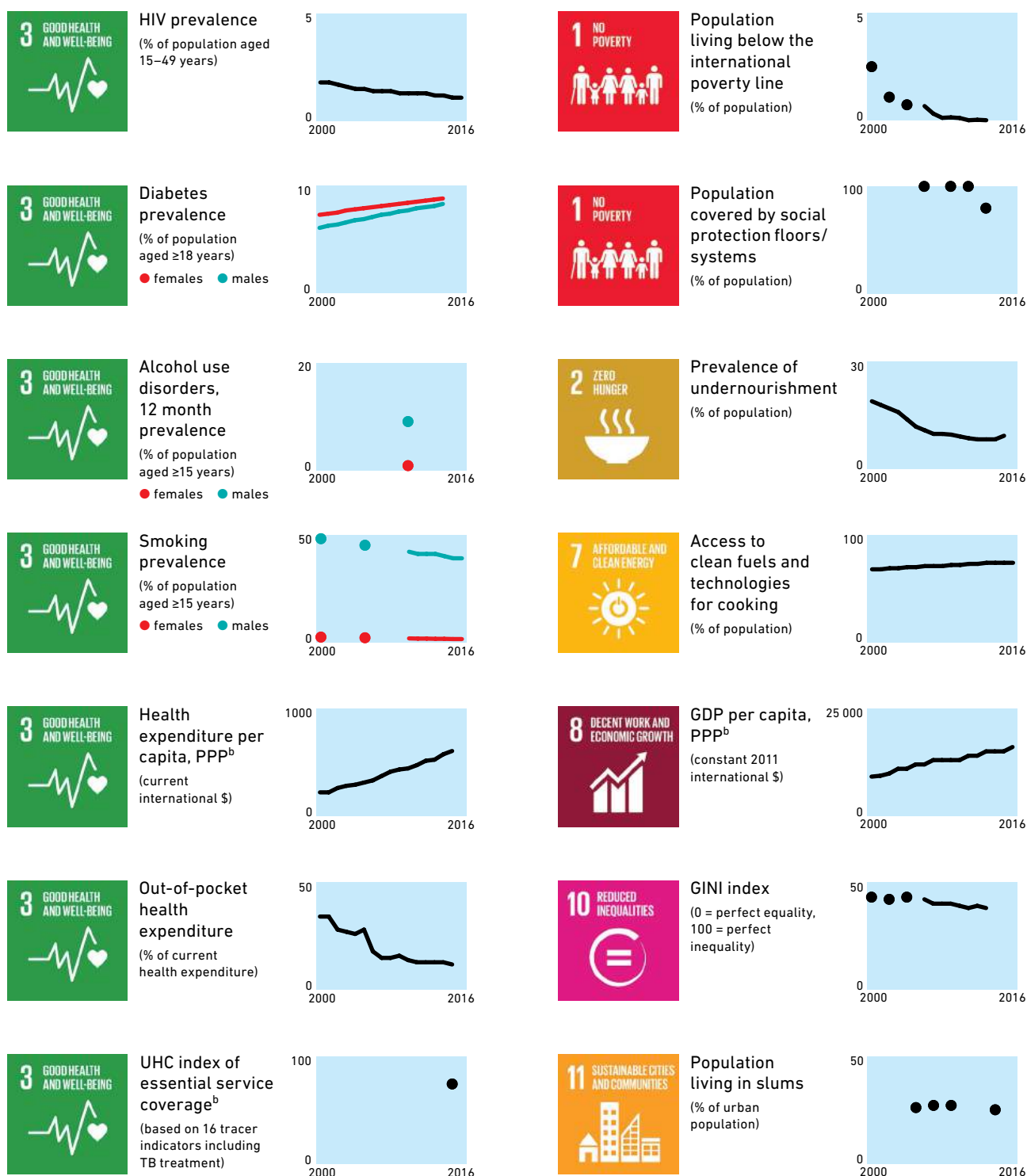
^b MDR is TB resistant to rifampicin and isoniazid; RR is TB resistant to rifampicin.

^c Includes cases with unknown previous TB treatment history.

^d Includes patients diagnosed before 2017 and patients who were not laboratory-confirmed.



INDICATORS IN THE SUSTAINABLE DEVELOPMENT GOALS ASSOCIATED WITH TB INCIDENCE^a



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^b GDP = gross domestic product; PPP = purchasing power parity; UHC = universal health coverage

United Republic of Tanzania

POPULATION 2017 **57 MILLION**

ESTIMATES OF TB BURDEN,^a 2017

	NUMBER (THOUSANDS)	RATE (PER 100 000 POPULATION)
Mortality (excludes HIV+TB)	27 (12–48)	47 (21–83)
Mortality (HIV+TB only)	22 (10–38)	39 (18–67)
Incidence (includes HIV+TB)	154 (73–266)	269 (127–464)
Incidence (HIV+TB only)	48 (31–69)	84 (54–120)
Incidence (MDR/RR-TB)**	1.7 (0.52–3.5)	2.9 (0.91–6)

ESTIMATED TB INCIDENCE BY AGE AND SEX (THOUSANDS),^a 2017

	0–14 YEARS	> 14 YEARS	TOTAL
Females	4.1 (3.7–4.5)	46 (30–62)	50 (32–68)
Males	4.6 (4.1–5)	100 (50–150)	104 (51–158)
Total	8.7 (7.4–9.9)	146 (57–234)	154 (73–266)

TB CASE NOTIFICATIONS, 2017

Total cases notified	69 818
Total new and relapse	68 473
— % tested with rapid diagnostics at time of diagnosis	14%
— % with known HIV status	98%
— % pulmonary	80%
— % bacteriologically confirmed among pulmonary	52%

UNIVERSAL HEALTH COVERAGE AND SOCIAL PROTECTION

TB treatment coverage (notified/estimated incidence), 2017	44% (26–94)
TB patients facing catastrophic total costs	
TB case fatality ratio (estimated mortality/estimated incidence), 2017	0.35 (0.13–0.62)

TB/HIV CARE IN NEW AND RELAPSE TB PATIENTS, 2017

	NUMBER	(%)
Patients with known HIV-status who are HIV-positive	21 449	31%
— on antiretroviral therapy	20 314	95%

DRUG-RESISTANT TB CARE, 2017

	NEW CASES	PREVIOUSLY TREATED CASES	TOTAL NUMBER ^e
Estimated MDR/RR-TB cases among notified pulmonary TB cases			640 (290–980)
Estimated % of TB cases with MDR/RR-TB	0.9% (0.3–1.5)	4.7% (0.7–8.6)	
% notified tested for rifampicin resistance	14%	56%	11 769
MDR/RR-TB cases tested for resistance to second-line drugs			44
Laboratory-confirmed cases			MDR/RR-TB: 200, XDR-TB: 0
Patients started on treatment ^d			MDR/RR-TB: 167, XDR-TB: 0

TREATMENT SUCCESS RATE AND COHORT SIZE

	SUCCESS	COHORT
New and relapse cases registered in 2016	90%	64 609
Previously treated cases, excluding relapse, registered in 2016	81%	1 334
HIV-positive TB cases registered in 2016	85%	22 642
MDR/RR-TB cases started on second-line treatment in 2015	74%	119
XDR-TB cases started on second-line treatment in 2015	100%	1

TB PREVENTIVE TREATMENT, 2017

% of HIV-positive people (newly enrolled in care) on preventive treatment	
% of children (aged < 5) household contacts of bacteriologically-confirmed TB cases on preventive treatment	35% (32–38)

TB FINANCING, 2018

National TB budget (US\$ millions)	64
Funding source:	3% domestic, 46% international, 52% unfunded

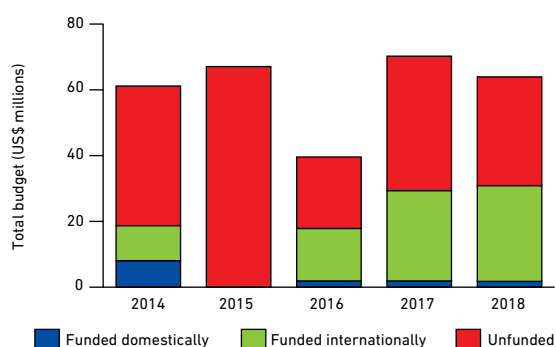
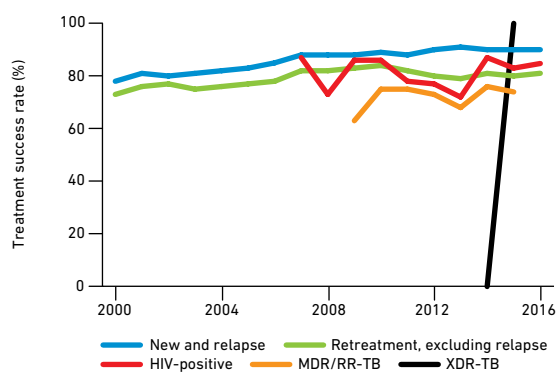
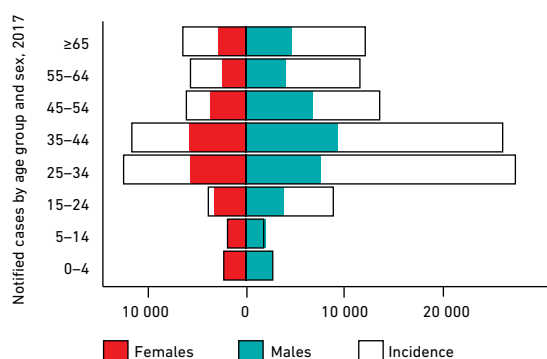
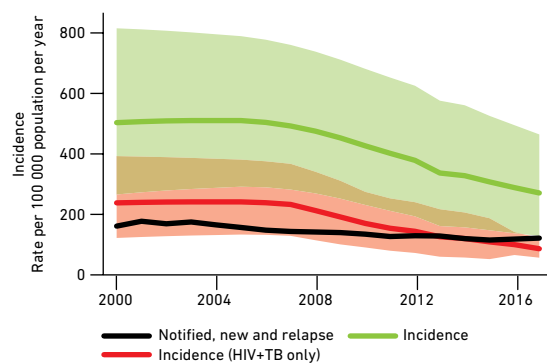
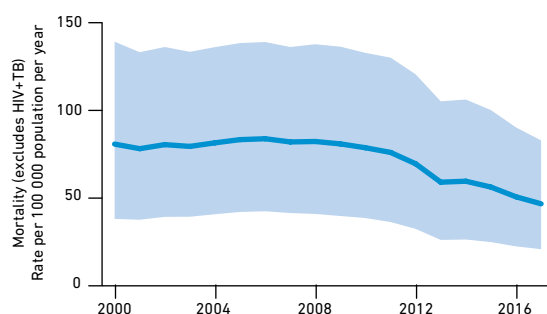
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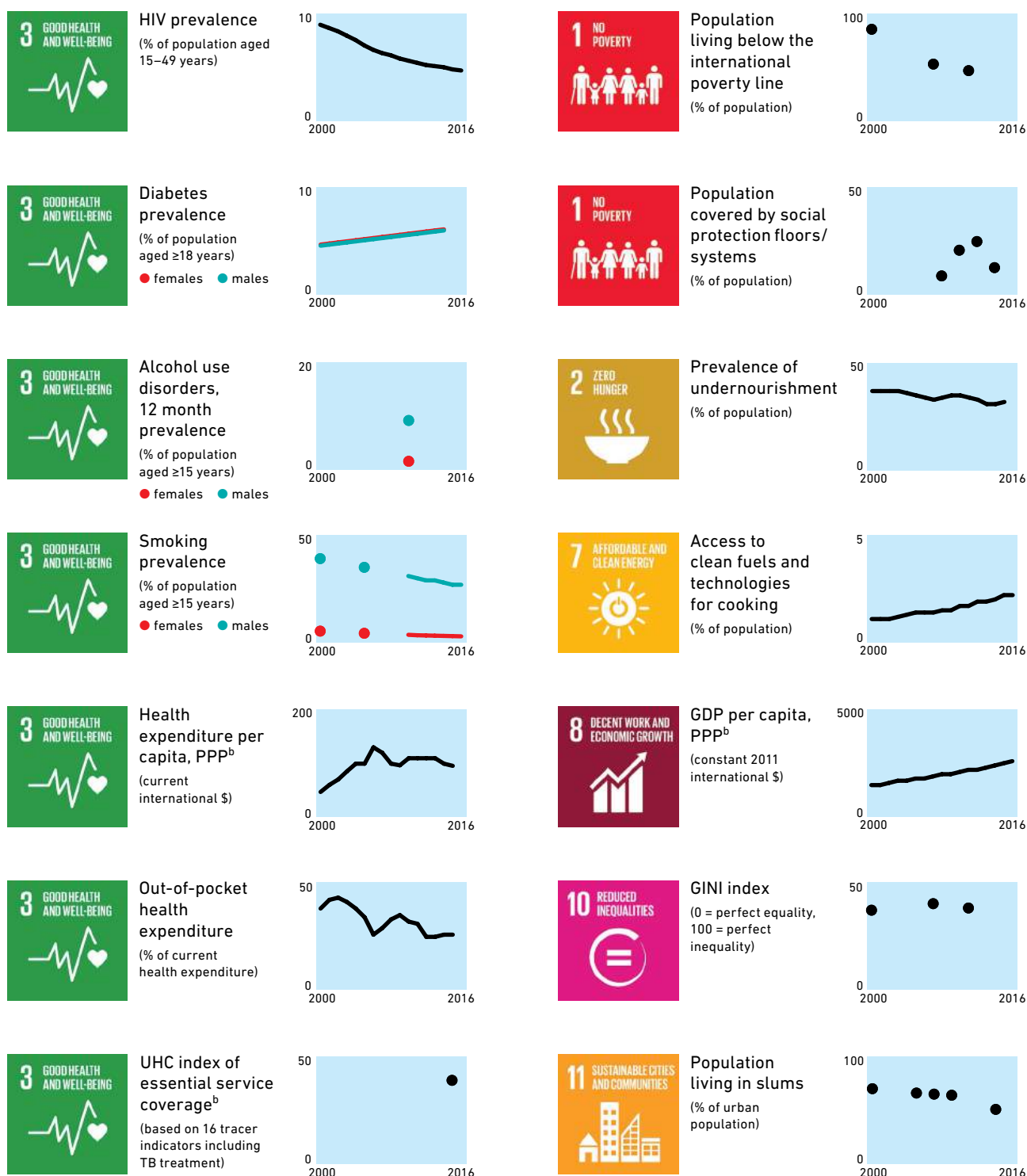
^b MDR is TB resistant to rifampicin and isoniazid; RR is TB resistant to rifampicin.

^c Includes cases with unknown previous TB treatment history.

^d Includes patients diagnosed before 2017 and patients who were not laboratory-confirmed.



INDICATORS IN THE SUSTAINABLE DEVELOPMENT GOALS ASSOCIATED WITH TB INCIDENCE^a



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^b GDP = gross domestic product; PPP = purchasing power parity; UHC = universal health coverage

ESTIMATES OF TB BURDEN, ^a 2017

	NUMBER (THOUSANDS)	RATE (PER 100 000 POPULATION)
Mortality (excludes HIV+TB)	12 (7.5–17)	12 (7.8–17)
Mortality (HIV+TB only)	0.84 (0.61–1.1)	0.88 (0.64–1.2)
Incidence (includes HIV+TB)	124 (101–148)	129 (106–155)
Incidence (HIV+TB only)	4.5 (3.7–5.4)	4.7 (3.8–5.7)
Incidence (MDR/RR-TB)**	7.1 (4.6–10)	7.4 (4.8–11)

ESTIMATED TB INCIDENCE BY AGE AND SEX (THOUSANDS), ^a 2017

	0–14 YEARS	> 14 YEARS	TOTAL
Females	6.6 (6.3–6.9)	30 (27–33)	37 (33–40)
Males	8.5 (8.1–8.9)	79 (67–90)	87 (73–101)
Total	15 (14–16)	109 (89–128)	124 (101–148)

TB CASE NOTIFICATIONS, 2017

Total cases notified	105 733
Total new and relapse	102 725
— % tested with rapid diagnostics at time of diagnosis	26%
— % with known HIV status	85%
— % pulmonary	80%
— % bacteriologically confirmed among pulmonary	70%

UNIVERSAL HEALTH COVERAGE AND SOCIAL PROTECTION

TB treatment coverage (notified/estimated incidence), 2017	83% (69–100)
TB patients facing catastrophic total costs, 2016	63% (58–67)
TB case fatality ratio (estimated mortality/estimated incidence), 2017	0.1 (0.06–0.15)

TB/HIV CARE IN NEW AND RELAPSE TB PATIENTS, 2017

	NUMBER	(%)
Patients with known HIV-status who are HIV-positive	3 239	4%
— on antiretroviral therapy	3 054	94%

DRUG-RESISTANT TB CARE, 2017

	NEW CASES	PREVIOUSLY TREATED CASES	TOTAL NUMBER ^e
Estimated MDR/RR-TB cases among notified pulmonary TB cases			4 900 (3 800–6 000)
Estimated % of TB cases with MDR/RR-TB	4.1% (2.7–5.7)	17% (17–18)	
% notified tested for rifampicin resistance	32%	67%	41 595
MDR/RR-TB cases tested for resistance to second-line drugs			1 484
Laboratory-confirmed cases		MDR/RR-TB: 3 071, XDR-TB: 50	
Patients started on treatment ^d		MDR/RR-TB: 2 694, XDR-TB: 12	

TREATMENT SUCCESS RATE AND COHORT SIZE

	SUCCESS	COHORT
New and relapse cases registered in 2016	92%	99 869
Previously treated cases, excluding relapse, registered in 2016	86%	4 430
HIV-positive TB cases registered in 2016	49%	2 669
MDR/RR-TB cases started on second-line treatment in 2015	74%	2 045
XDR-TB cases started on second-line treatment in 2015		

TB PREVENTIVE TREATMENT, 2017

% of HIV-positive people (newly enrolled in care) on preventive treatment	31%
% of children (aged < 5) household contacts of bacteriologically-confirmed TB cases on preventive treatment	26% (24–29)

TB FINANCING, 2018

National TB budget (US\$ millions)	70
Funding source:	11% domestic, 28% international, 61% unfunded

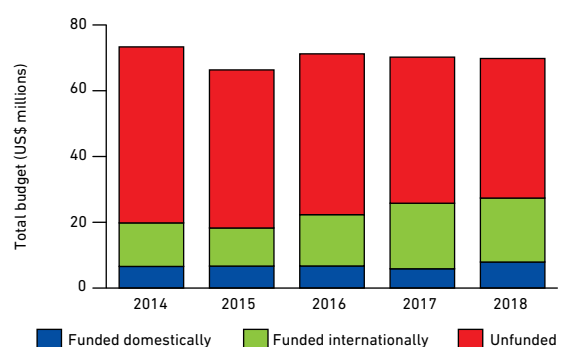
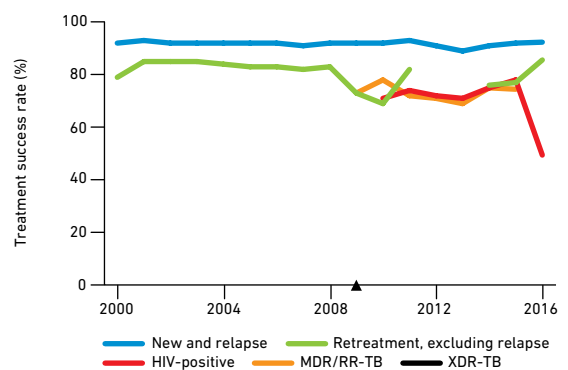
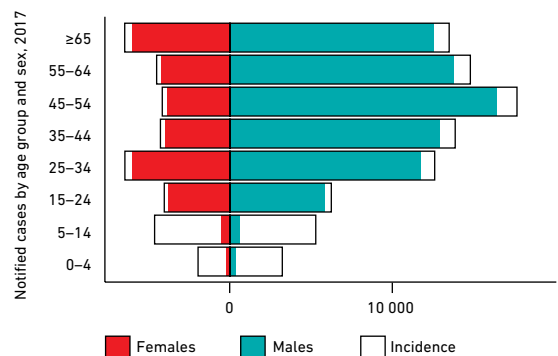
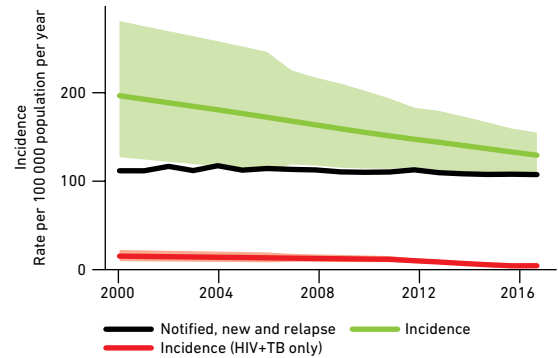
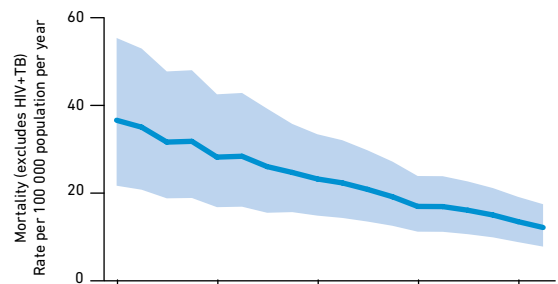
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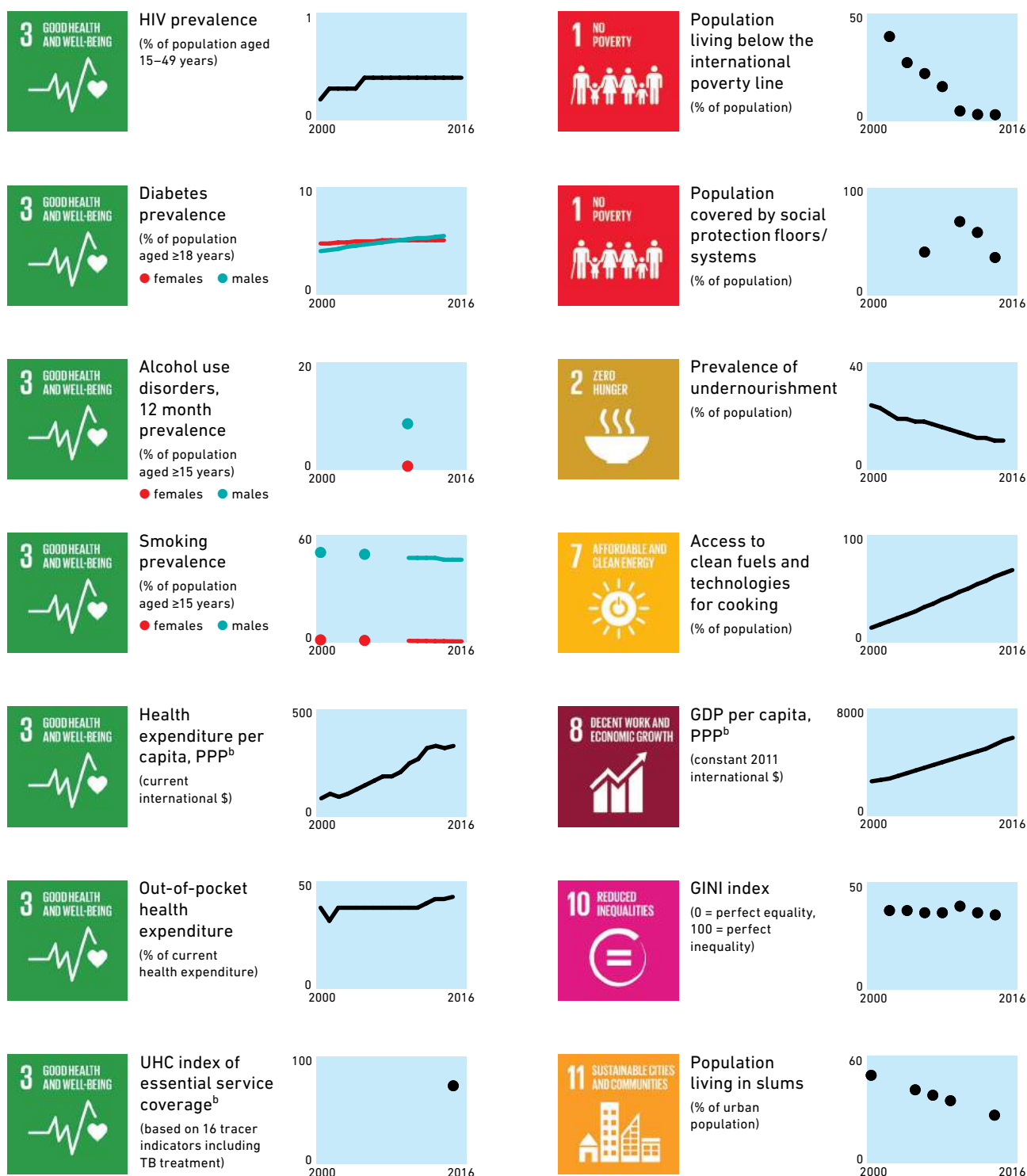
^b MDR is TB resistant to rifampicin and isoniazid; RR is TB resistant to rifampicin.

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^d Includes patients diagnosed before 2017 and patients who were not laboratory-confirmed.



INDICATORS IN THE SUSTAINABLE DEVELOPMENT GOALS ASSOCIATED WITH TB INCIDENCE^a



Targets for reductions in TB incidence and TB deaths set in WHO's End TB Strategy and the United Nations' Sustainable Development Goals (SDGs) are ambitious. Achieving them requires progress in reducing health-related risk factors for TB infection and disease, as well as broader social and economic determinants of TB infection and disease. WHO has developed a TB-SDG monitoring framework that comprises 14 indicators under seven SDGs for which there is evidence of an association with TB incidence. Further details are provided in [Chapter 2](#).

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^b GDP = gross domestic product; PPP = purchasing power parity; UHC = universal health coverage

Cambodia

POPULATION 2017 **16 MILLION**

ESTIMATES OF TB BURDEN, ^a 2017

	NUMBER (THOUSANDS)	RATE (PER 100 000 POPULATION)
Mortality (excludes HIV+TB)	3.1 (2–4.3)	19 (13–27)
Mortality (HIV+TB only)	0.41 (0.27–0.57)	2.6 (1.7–3.6)
Incidence (includes HIV+TB)	52 (36–72)	326 (224–447)
Incidence (HIV+TB only)	1.3 (0.89–1.8)	8.2 (5.6–11)
Incidence (MDR/RR-TB)**	1.2 (0.52–2.1)	7.2 (3.2–13)

ESTIMATED TB INCIDENCE BY AGE AND SEX (THOUSANDS), ^a 2017

	0–14 YEARS	> 14 YEARS	TOTAL
Females	3.1 (2.9–3.4)	19 (15–22)	22 (17–27)
Males	3.4 (3.1–3.7)	27 (20–34)	30 (22–38)
Total	6.6 (5.8–7.4)	46 (31–60)	52 (36–72)

TB CASE NOTIFICATIONS, 2017

Total cases notified	34 467
Total new and relapse	34 238
— % tested with rapid diagnostics at time of diagnosis	
— % with known HIV status	87%
— % pulmonary	66%
— % bacteriologically confirmed among pulmonary	54%

UNIVERSAL HEALTH COVERAGE AND SOCIAL PROTECTION

TB treatment coverage (notified/estimated incidence), 2017	66% (48–96)
TB patients facing catastrophic total costs	
TB case fatality ratio (estimated mortality/estimated incidence), 2017	0.07 (0.04–0.1)

TB/HIV CARE IN NEW AND RELAPSE TB PATIENTS, 2017

	NUMBER	(%)
Patients with known HIV-status who are HIV-positive	748	3%
— on antiretroviral therapy	698	93%

DRUG-RESISTANT TB CARE, 2017

	NEW CASES	PREVIOUSLY TREATED CASES	TOTAL NUMBER ^e
Estimated MDR/RR-TB cases among notified pulmonary TB cases			560 (290–840)
Estimated % of TB cases with MDR/RR-TB	1.8% (0.9–3)	11% (3.2–22)	
% notified tested for rifampicin resistance		33%	1 313
MDR/RR-TB cases tested for resistance to second-line drugs			100
Laboratory-confirmed cases		MDR/RR-TB: 136, XDR-TB: 1	
Patients started on treatment ^d		MDR/RR-TB: 143, XDR-TB: 1	

TREATMENT SUCCESS RATE AND COHORT SIZE

	SUCCESS	COHORT
New and relapse cases registered in 2016	94%	32 478
Previously treated cases, excluding relapse, registered in 2016	92%	38
HIV-positive TB cases registered in 2016		
MDR/RR-TB cases started on second-line treatment in 2015	71%	75
XDR-TB cases started on second-line treatment in 2015		0

TB PREVENTIVE TREATMENT, 2017

% of HIV-positive people (newly enrolled in care) on preventive treatment	21%
% of children (aged < 5) household contacts of bacteriologically-confirmed TB cases on preventive treatment	44% (40–48)

TB FINANCING, 2018

National TB budget (US\$ millions)	37
Funding source:	12% domestic, 24% international, 64% unfunded

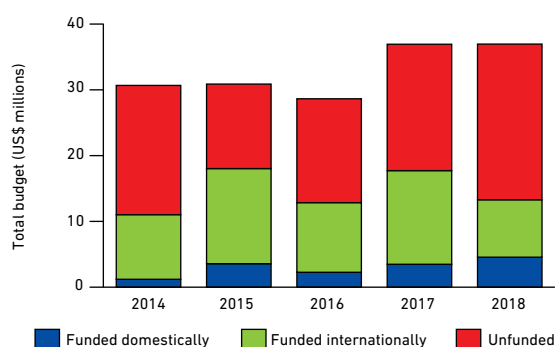
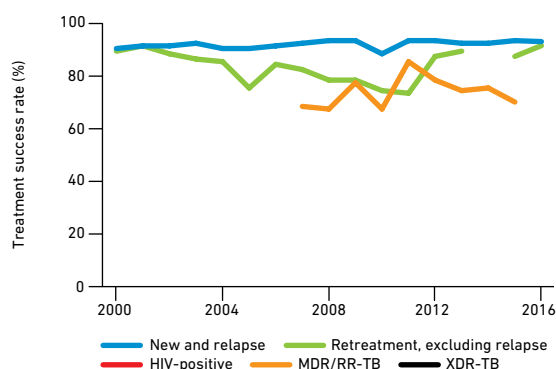
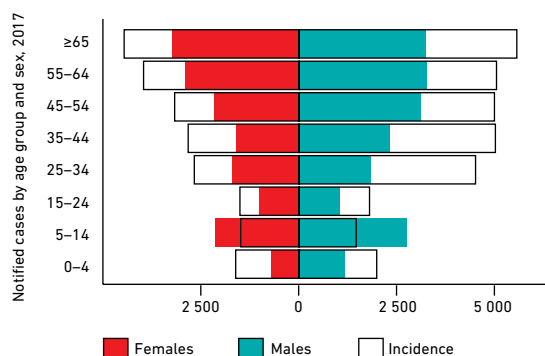
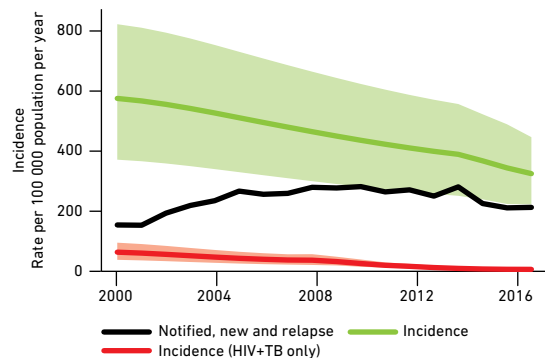
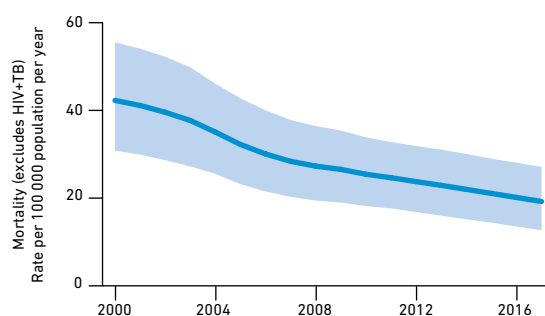
Data are as reported to WHO. Estimates of TB and MDR/RR-TB burden are produced by WHO in consultation with countries. Estimates are rounded and totals are computed prior to rounding.

^a Ranges represent uncertainty intervals.

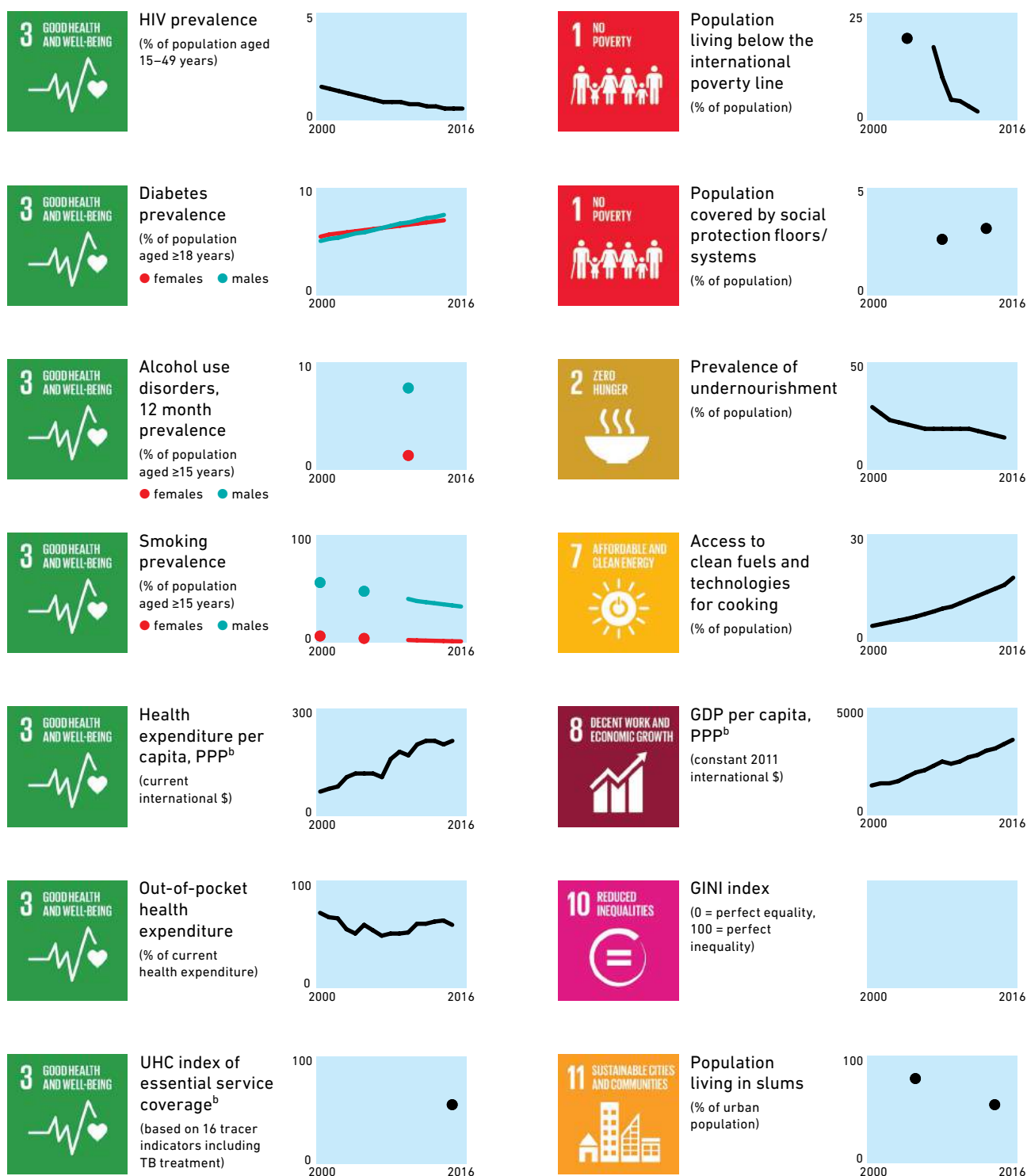
^b MDR is TB resistant to rifampicin and isoniazid; RR is TB resistant to rifampicin.

^c Includes cases with unknown previous TB treatment history.

^d Includes patients diagnosed before 2017 and patients who were not laboratory-confirmed.



INDICATORS IN THE SUSTAINABLE DEVELOPMENT GOALS ASSOCIATED WITH TB INCIDENCE^a



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^b GDP = gross domestic product; PPP = purchasing power parity; UHC = universal health coverage

Central African Republic

POPULATION 2017 **4.7 MILLION**

ESTIMATES OF TB BURDEN, ^a 2017

	NUMBER (THOUSANDS)	RATE (PER 100 000 POPULATION)
Mortality (excludes HIV+TB)	3.2 (1.8–4.9)	68 (38–105)
Mortality (HIV+TB only)	2.7 (1.4–4.4)	58 (31–94)
Incidence (includes HIV+TB)	20 (13–28)	423 (274–604)
Incidence (HIV+TB only)	6.2 (3.3–10)	134 (72–214)
Incidence (MDR/RR-TB) ^b	0.15 (0.084–0.23)	3.2 (1.8–4.9)

ESTIMATED TB INCIDENCE BY AGE AND SEX (THOUSANDS), ^a 2017

	0–14 YEARS	> 14 YEARS	TOTAL
Females	1.3 (1.2–1.4)	6.3 (4.9–7.6)	7.5 (5.7–9.4)
Males	1.4 (1.3–1.6)	11 (7.6–14)	12 (8.4–16)
Total	2.7 (2.3–3.1)	17 (11–23)	20 (13–28)

TB CASE NOTIFICATIONS, 2017

Total cases notified	9 819
Total new and relapse	9 664
— % tested with rapid diagnostics at time of diagnosis	3%
— % with known HIV status	77%
— % pulmonary	81%
— % bacteriologically confirmed among pulmonary	66%

UNIVERSAL HEALTH COVERAGE AND SOCIAL PROTECTION

TB treatment coverage (notified/estimated incidence), 2017	49% (34–76)
TB patients facing catastrophic total costs	
TB case fatality ratio (estimated mortality/estimated incidence), 2017	0.31 (0.16–0.48)

TB/HIV CARE IN NEW AND RELAPSE TB PATIENTS, 2017

	NUMBER	(%)
Patients with known HIV-status who are HIV-positive	2 098	28%
— on antiretroviral therapy	1 541	73%

DRUG-RESISTANT TB CARE, 2017

	NEW CASES	PREVIOUSLY TREATED CASES	TOTAL NUMBER ^c
Estimated MDR/RR-TB cases among notified pulmonary TB cases			91 (0–190)
Estimated % of TB cases with MDR/RR-TB	0.4% (0–2.2)	14% (10–18)	
% notified tested for rifampicin resistance	0%	62%	268
MDR/RR-TB cases tested for resistance to second-line drugs			2
Laboratory-confirmed cases			MDR/RR-TB: 93, XDR-TB: 2
Patients started on treatment ^d			MDR/RR-TB: 86, XDR-TB: 0

TREATMENT SUCCESS RATE AND COHORT SIZE

	SUCCESS	COHORT
New and relapse cases registered in 2016	78%	5 312
Previously treated cases, excluding relapse, registered in 2016	79%	186
HIV-positive TB cases registered in 2016	73%	2 053
MDR/RR-TB cases started on second-line treatment in 2015	88%	41
XDR-TB cases started on second-line treatment in 2015		0

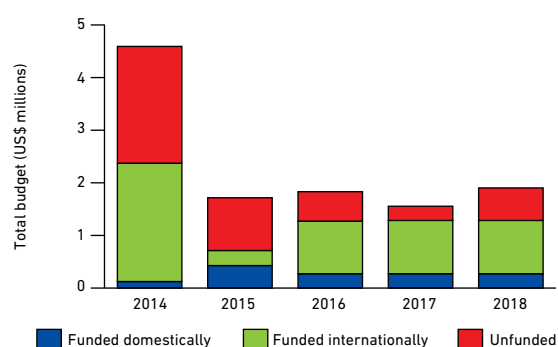
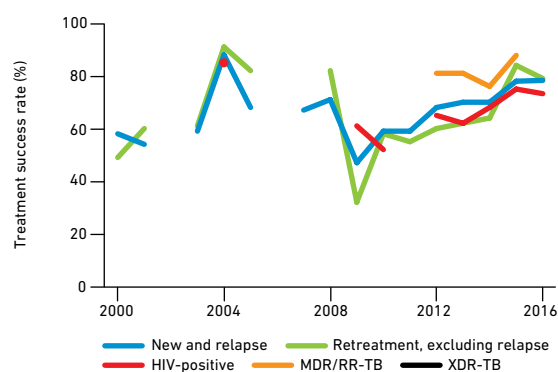
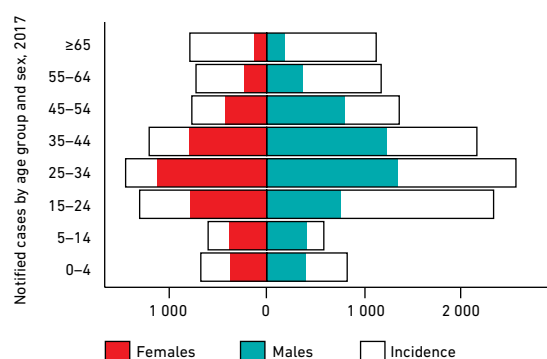
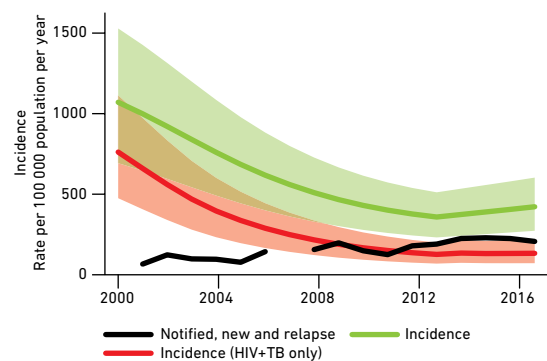
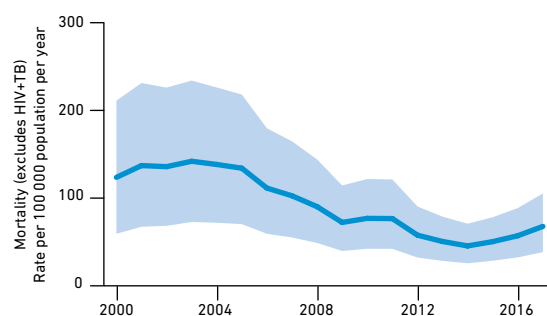
TB PREVENTIVE TREATMENT, 2017

% of HIV-positive people (newly enrolled in care) on preventive treatment	0%
% of children (aged < 5) household contacts of bacteriologically-confirmed TB cases on preventive treatment	

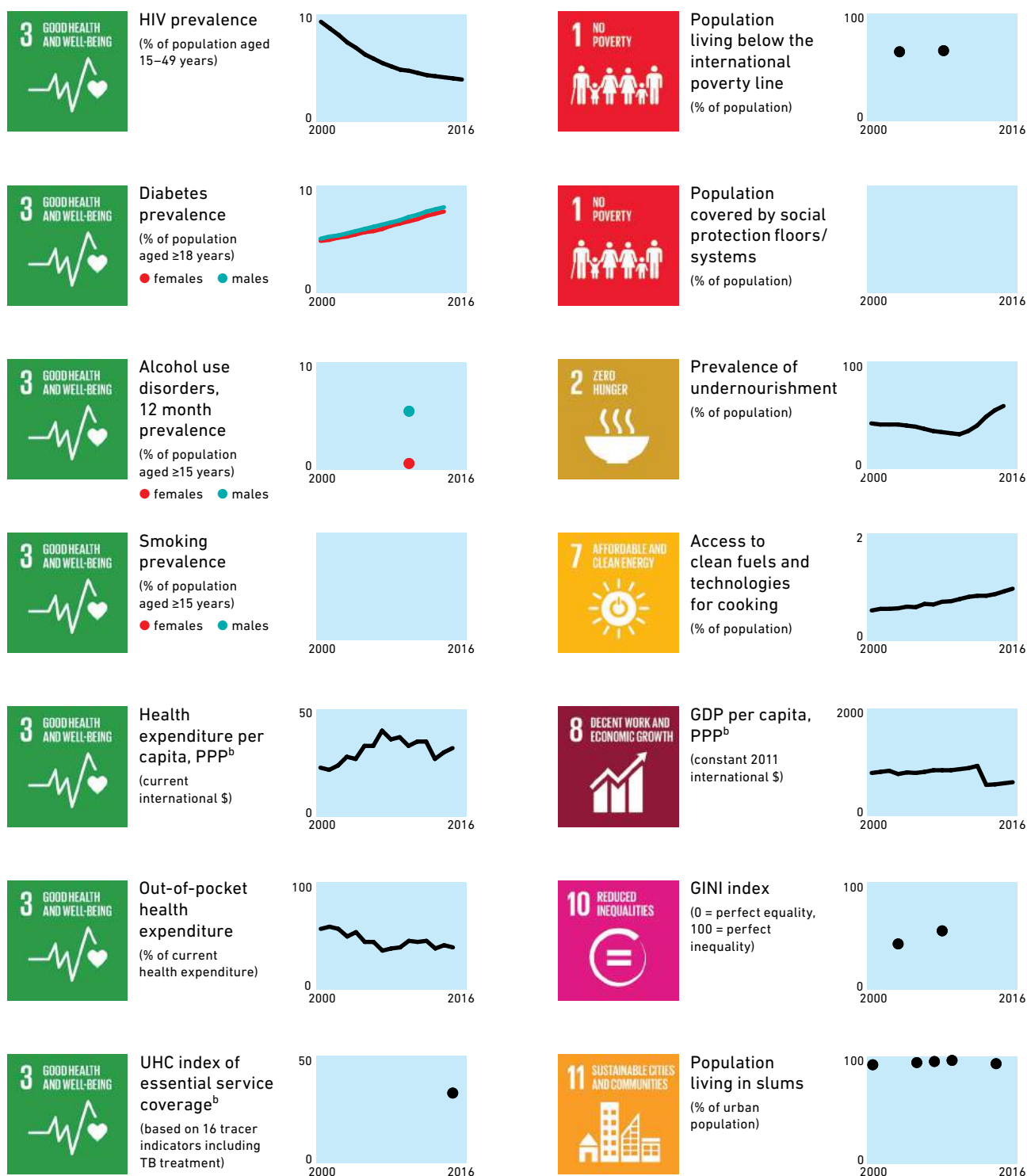
TB FINANCING, 2018

National TB budget (US\$ millions)	1.9
Funding source:	14% domestic, 53% international, 32% unfunded

Data are as reported to WHO. Estimates of TB and MDR/RR-TB burden are produced by WHO in consultation with countries. Estimates are rounded and totals are computed prior to rounding.
^a Ranges represent uncertainty intervals.
^b MDR is TB resistant to rifampicin and isoniazid; RR is TB resistant to rifampicin.
^c Includes cases with unknown previous TB treatment history.
^d Includes patients diagnosed before 2017 and patients who were not laboratory-confirmed.



INDICATORS IN THE SUSTAINABLE DEVELOPMENT GOALS ASSOCIATED WITH TB INCIDENCE^a



Targets for reductions in TB incidence and TB deaths set in WHO's End TB Strategy and the United Nations' Sustainable Development Goals (SDGs) are ambitious. Achieving them requires progress in reducing health-related risk factors for TB infection and disease, as well as broader social and economic determinants of TB infection and disease. WHO has developed a TB-SDG monitoring framework that comprises 14 indicators under seven SDGs for which there is evidence of an association with TB incidence. Further details are provided in [Chapter 2](#).

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^b GDP = gross domestic product; PPP = purchasing power parity; UHC = universal health coverage

Congo

POPULATION 2017 **5.3 MILLION**

ESTIMATES OF TB BURDEN, ^a 2017

	NUMBER (THOUSANDS)	RATE (PER 100 000 POPULATION)
Mortality (excludes HIV+TB)	3.3 (1.9–5.2)	63 (36–98)
Mortality (HIV+TB only)	2.3 (1.2–3.7)	43 (22–71)
Incidence (includes HIV+TB)	20 (13–29)	376 (239–545)
Incidence (HIV+TB only)	5.3 (2.7–8.6)	100 (52–164)
Incidence (MDR/RR-TB) ^b	0.61 (0.25–1.1)	12 (4.8–21)

ESTIMATED TB INCIDENCE BY AGE AND SEX (THOUSANDS), ^a 2017

	0–14 YEARS	> 14 YEARS	TOTAL
Females	1.3 (1.2–1.4)	6.3 (4.8–7.7)	7.6 (5.7–9.5)
Males	1.5 (1.3–1.6)	11 (7.6–14)	12 (8.3–16)
Total	2.8 (2.3–3.2)	17 (11–23)	20 (13–29)

TB CASE NOTIFICATIONS, 2017

Total cases notified	10 263
Total new and relapse	10 005
— % tested with rapid diagnostics at time of diagnosis	5%
— % with known HIV status	13%
— % pulmonary	78%
— % bacteriologically confirmed among pulmonary	51%

UNIVERSAL HEALTH COVERAGE AND SOCIAL PROTECTION

TB treatment coverage (notified/estimated incidence), 2017	51% (35–80)
TB patients facing catastrophic total costs	
TB case fatality ratio (estimated mortality/estimated incidence), 2017	0.29 (0.15–0.46)

TB/HIV CARE IN NEW AND RELAPSE TB PATIENTS, 2017

	NUMBER	(%)
Patients with known HIV-status who are HIV-positive	374	30%
— on antiretroviral therapy	272	73%

DRUG-RESISTANT TB CARE, 2017

	NEW CASES	PREVIOUSLY TREATED CASES	TOTAL NUMBER ^c
Estimated MDR/RR-TB cases among notified pulmonary TB cases			290 (170–420)
Estimated % of TB cases with MDR/RR-TB	2.5% (1.1–4.3)	21% (16–27)	
% notified tested for rifampicin resistance	<1%	89%	532
MDR/RR-TB cases tested for resistance to second-line drugs			0
Laboratory-confirmed cases		MDR/RR-TB: 58, XDR-TB: 0	
Patients started on treatment ^d		MDR/RR-TB: 28, XDR-TB: 0	

TREATMENT SUCCESS RATE AND COHORT SIZE

	SUCCESS	COHORT
New and relapse cases registered in 2016	77%	10 656
Previously treated cases, excluding relapse, registered in 2016	53%	232
HIV-positive TB cases registered in 2016		0
MDR/RR-TB cases started on second-line treatment in 2015	62%	13
XDR-TB cases started on second-line treatment in 2015		0

TB PREVENTIVE TREATMENT, 2017

% of HIV-positive people (newly enrolled in care) on preventive treatment	
% of children (aged < 5) household contacts of bacteriologically-confirmed TB cases on preventive treatment	4.1% (3.7–4.5)

TB FINANCING, 2018

National TB budget (US\$ millions)	1.5
Funding source:	72% domestic, 28% international, 0% unfunded

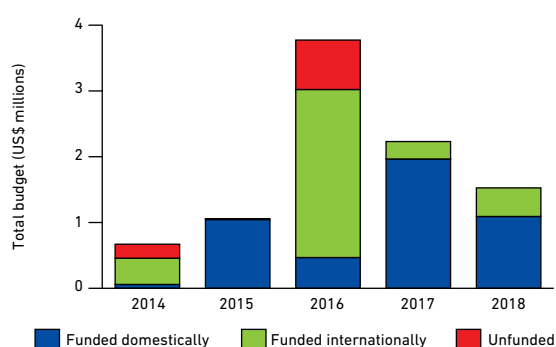
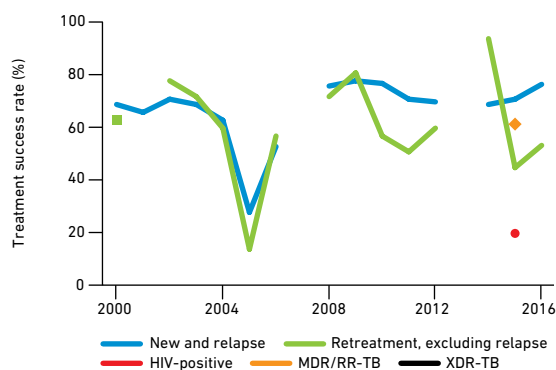
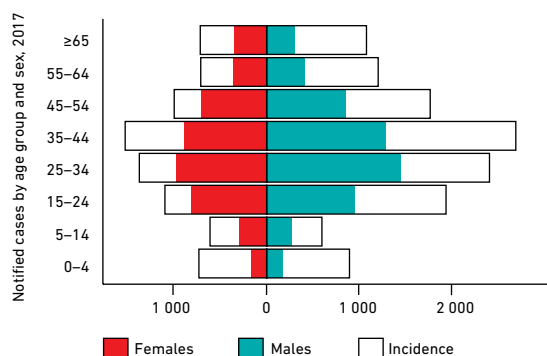
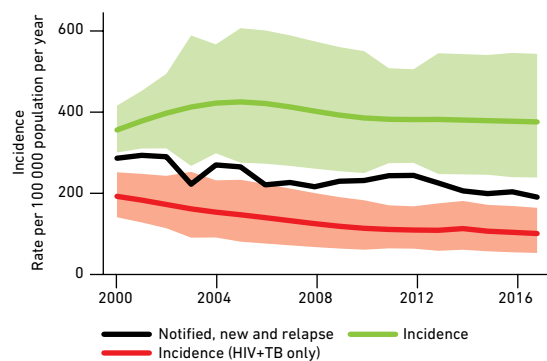
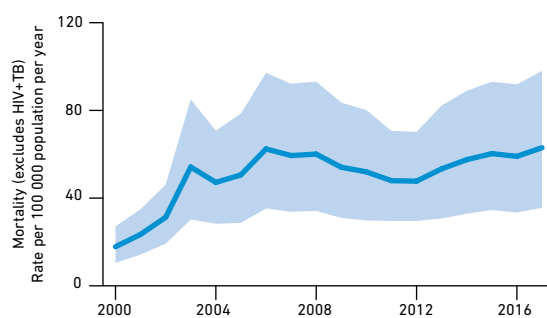
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^a Ranges represent uncertainty intervals.

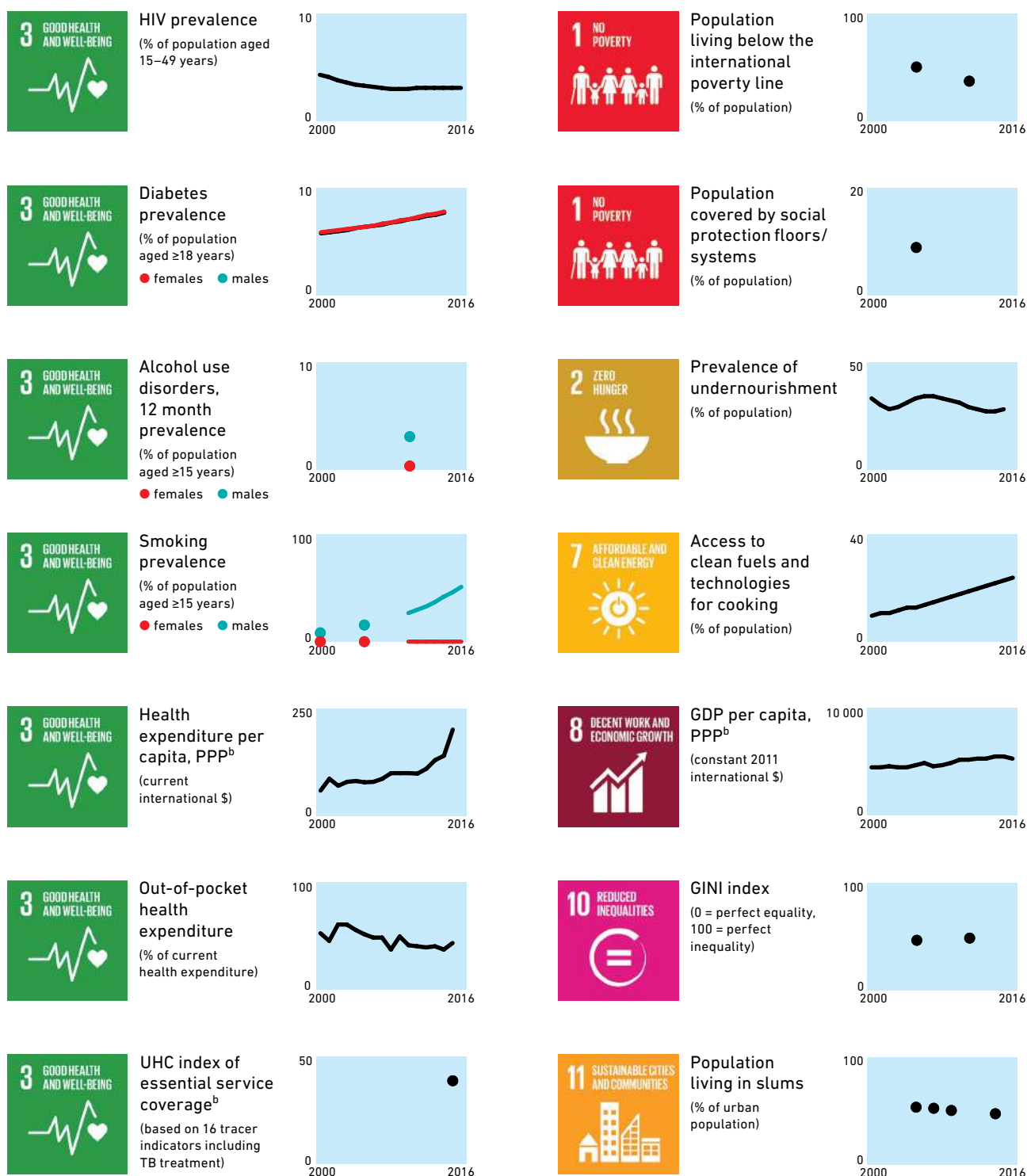
^b MDR is TB resistant to rifampicin and isoniazid; RR is TB resistant to rifampicin.

^c Includes cases with unknown previous TB treatment history.

^d Includes patients diagnosed before 2017 and patients who were not laboratory-confirmed.



INDICATORS IN THE SUSTAINABLE DEVELOPMENT GOALS ASSOCIATED WITH TB INCIDENCE^a



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^b GDP = gross domestic product; PPP = purchasing power parity; UHC = universal health coverage

Lesotho

ESTIMATES OF TB BURDEN, ^a 2017

	NUMBER (THOUSANDS)	RATE (PER 100 000 POPULATION)
Mortality (excludes HIV+TB)	1 (0.55–1.7)	46 (25–75)
Mortality (HIV+TB only)	4.6 (2.9–6.7)	206 (128–302)
Incidence (includes HIV+TB)	15 (9.6–21)	665 (430–949)
Incidence (HIV+TB only)	11 (6.7–15)	470 (298–680)
Incidence (MDR/RR-TB) ^b	1.1 (0.61–1.7)	50 (27–78)

ESTIMATED TB INCIDENCE BY AGE AND SEX (THOUSANDS), ^a 2017

	0–14 YEARS	> 14 YEARS	TOTAL
Females	1 (0.93–1.1)	4.6 (3.6–5.7)	5.7 (4.3–7.1)
Males	1.1 (1–1.3)	8 (5.7–10)	9.2 (6.4–12)
Total	2.2 (1.8–2.5)	13 (8.1–17)	15 (9.6–21)

TB CASE NOTIFICATIONS, 2017

Total cases notified	7 271
Total new and relapse	7 117
— % tested with rapid diagnostics at time of diagnosis	
— % with known HIV status	92%
— % pulmonary	89%
— % bacteriologically confirmed among pulmonary	58%

UNIVERSAL HEALTH COVERAGE AND SOCIAL PROTECTION

TB treatment coverage (notified/estimated incidence), 2017	48% (34–74)
TB patients facing catastrophic total costs	
TB case fatality ratio (estimated mortality/estimated incidence), 2017	0.39 (0.2–0.6)

TB/HIV CARE IN NEW AND RELAPSE TB PATIENTS, 2017

	NUMBER	(%)
Patients with known HIV-status who are HIV-positive	4 690	70%
— on antiretroviral therapy	4 310	92%

DRUG-RESISTANT TB CARE, 2017

	NEW CASES	PREVIOUSLY TREATED CASES	TOTAL NUMBER ^c
Estimated MDR/RR-TB cases among notified pulmonary TB cases			410 (330–490)
Estimated % of TB cases with MDR/RR-TB	4.8% (3.7–6)	14% (9.5–18)	
% notified tested for rifampicin resistance	63%		3 839
MDR/RR-TB cases tested for resistance to second-line drugs			
Laboratory-confirmed cases			MDR/RR-TB: 351, XDR-TB:
Patients started on treatment ^d			MDR/RR-TB: 151, XDR-TB:

TREATMENT SUCCESS RATE AND COHORT SIZE

	SUCCESS	COHORT
New and relapse cases registered in 2016	77%	7 291
Previously treated cases, excluding relapse, registered in 2016	68%	228
HIV-positive TB cases registered in 2016	74%	5 085
MDR/RR-TB cases started on second-line treatment in 2015	66%	210
XDR-TB cases started on second-line treatment in 2015		

TB PREVENTIVE TREATMENT, 2017

% of HIV-positive people (newly enrolled in care) on preventive treatment	
% of children (aged < 5) household contacts of bacteriologically-confirmed TB cases on preventive treatment	

TB FINANCING, 2018

National TB budget (US\$ millions)	3.3
Funding source:	30% domestic, 55% international, 15% unfunded

Data are as reported to WHO. Estimates of TB and MDR/RR-TB burden are produced by WHO in consultation with countries. Estimates are rounded and totals are computed prior to rounding.

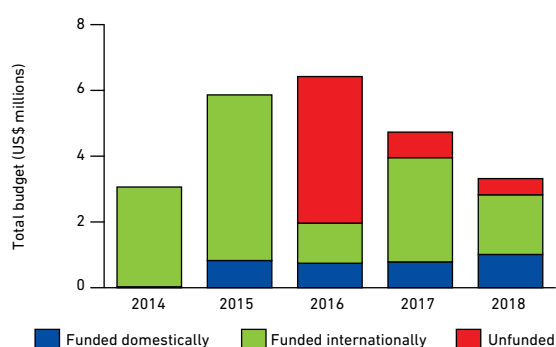
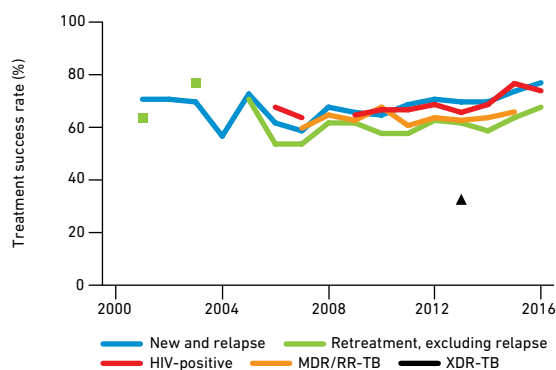
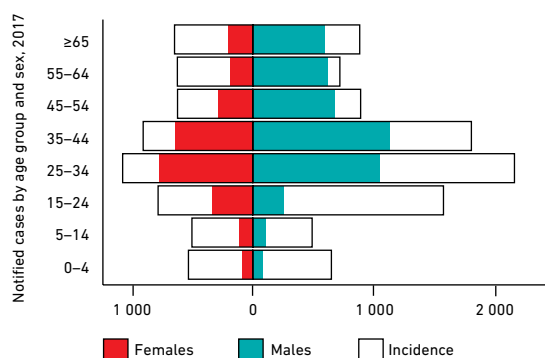
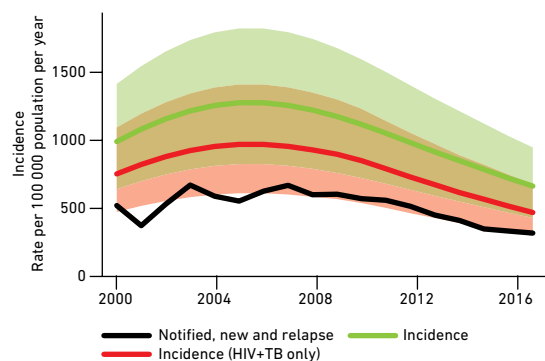
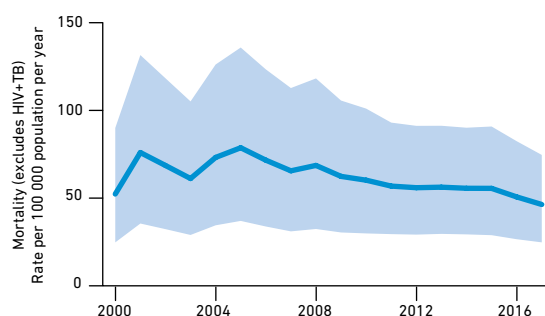
^a Ranges represent uncertainty intervals.

^b MDR is TB resistant to rifampicin and isoniazid; RR is TB resistant to rifampicin.

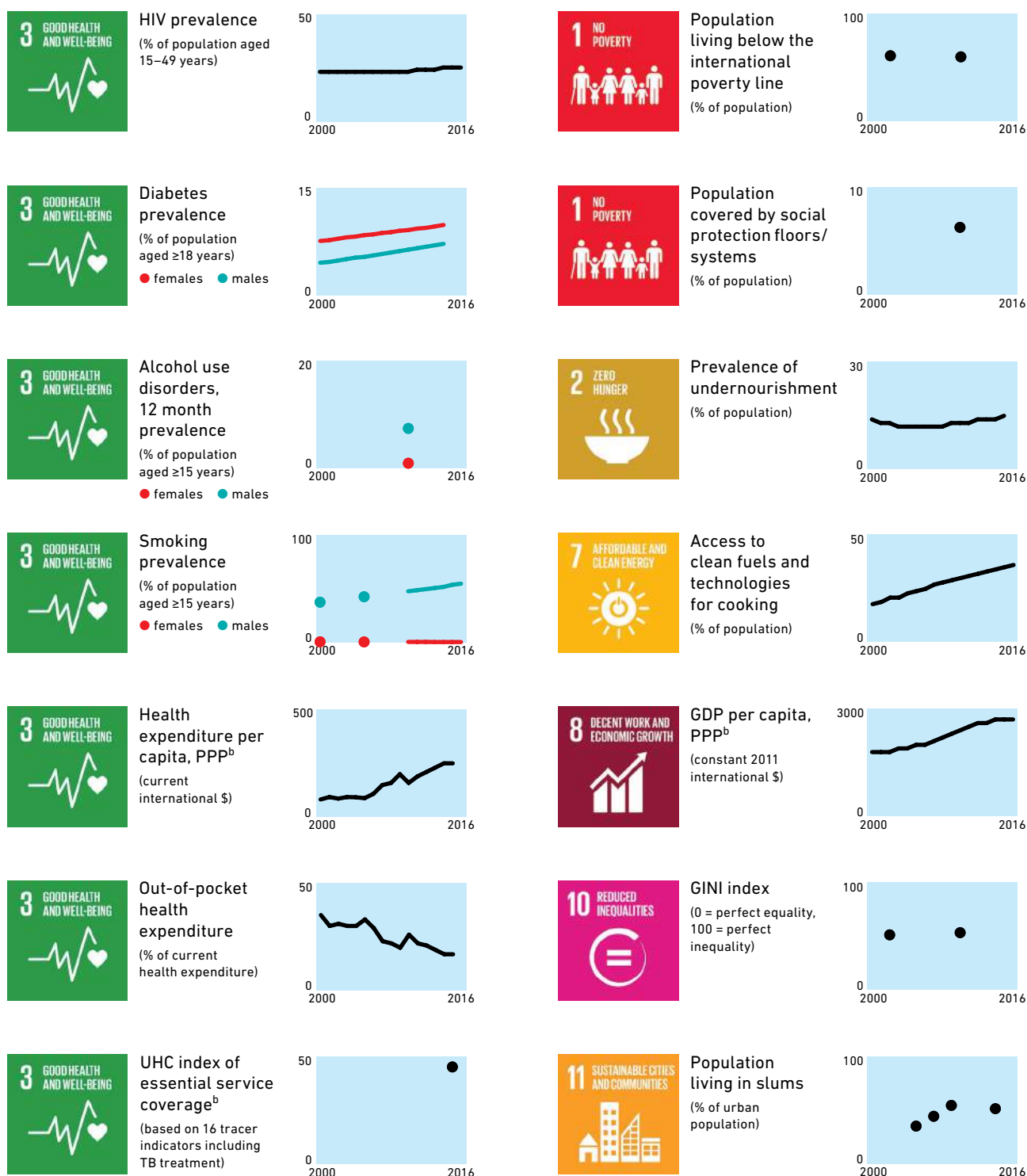
^c Includes cases with unknown previous TB treatment history.

^d Includes patients diagnosed before 2017 and patients who were not laboratory-confirmed.

POPULATION 2017 2.2 MILLION



INDICATORS IN THE SUSTAINABLE DEVELOPMENT GOALS ASSOCIATED WITH TB INCIDENCE^a



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^b GDP = gross domestic product; PPP = purchasing power parity; UHC = universal health coverage

Liberia

POPULATION 2017 **4.7 MILLION**

ESTIMATES OF TB BURDEN, ^a 2017

	NUMBER (THOUSANDS)	RATE (PER 100 000 POPULATION)
Mortality (excludes HIV+TB)	2.7 (1.6–4.1)	57 (34–86)
Mortality (HIV+TB only)	0.91 (0.57–1.3)	19 (12–28)
Incidence (includes HIV+TB)	15 (9.4–21)	308 (199–440)
Incidence (HIV+TB only)	2.2 (1.4–3.2)	47 (30–68)
Incidence (MDR/RR-TB)**	0.39 (0.15–0.74)	8.3 (3.2–16)

ESTIMATED TB INCIDENCE BY AGE AND SEX (THOUSANDS), ^a 2017

	0–14 YEARS	> 14 YEARS	TOTAL
Females	0.91 (0.82–0.99)	4.6 (3.6–5.7)	5.5 (4.2–6.9)
Males	1 (0.9–1.1)	8 (5.7–10)	9 (6.3–12)
Total	1.9 (1.6–2.2)	13 (8.1–17)	15 (9.4–21)

TB CASE NOTIFICATIONS, 2017

Total cases notified	7 728
Total new and relapse	7 728
— % tested with rapid diagnostics at time of diagnosis	17%
— % with known HIV status	70%
— % pulmonary	68%
— % bacteriologically confirmed among pulmonary	64%

UNIVERSAL HEALTH COVERAGE AND SOCIAL PROTECTION

TB treatment coverage (notified/estimated incidence), 2017	53% (37–82)
TB patients facing catastrophic total costs	
TB case fatality ratio (estimated mortality/estimated incidence), 2017	0.26 (0.14–0.4)

TB/HIV CARE IN NEW AND RELAPSE TB PATIENTS, 2017

	NUMBER	(%)
Patients with known HIV-status who are HIV-positive	827	15%
— on antiretroviral therapy	347	42%

DRUG-RESISTANT TB CARE, 2017

	NEW CASES	PREVIOUSLY TREATED CASES	TOTAL NUMBER ^e
Estimated MDR/RR-TB cases among notified pulmonary TB cases			150 (65–230)
Estimated % of TB cases with MDR/RR-TB	2.5% (1.1–4.3)	14% (10–18)	
% notified tested for rifampicin resistance	43%	100%	3 382
MDR/RR-TB cases tested for resistance to second-line drugs			19
Laboratory-confirmed cases			MDR/RR-TB: 88, XDR-TB: 0
Patients started on treatment ^d			MDR/RR-TB: 55, XDR-TB: 0

TREATMENT SUCCESS RATE AND COHORT SIZE

	SUCCESS	COHORT
New and relapse cases registered in 2016	77%	7 374
Previously treated cases, excluding relapse, registered in 2016	50%	127
HIV-positive TB cases registered in 2016		
MDR/RR-TB cases started on second-line treatment in 2015		
XDR-TB cases started on second-line treatment in 2015		

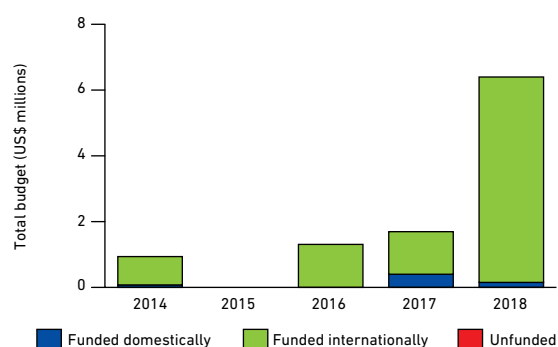
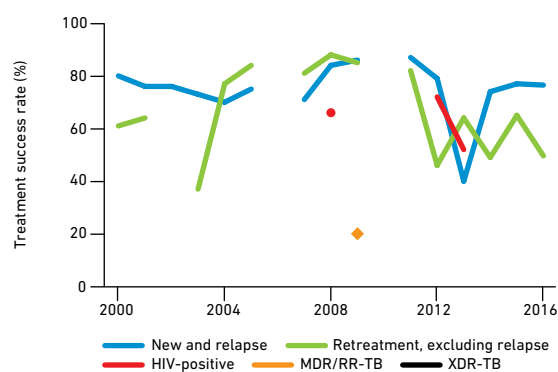
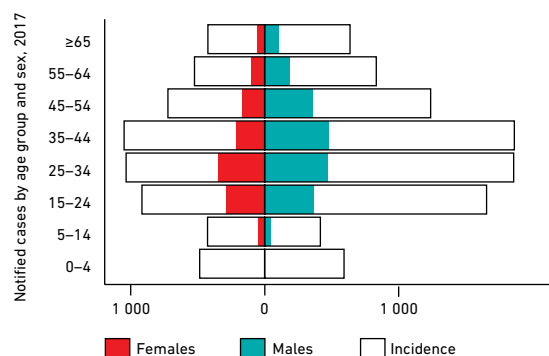
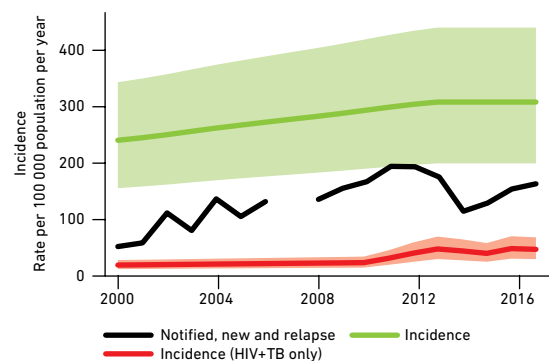
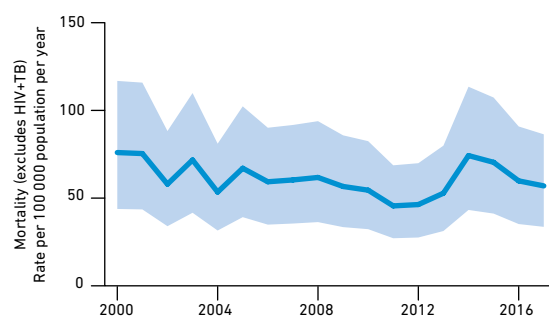
TB PREVENTIVE TREATMENT, 2017

% of HIV-positive people (newly enrolled in care) on preventive treatment	
% of children (aged < 5) household contacts of bacteriologically-confirmed TB cases on preventive treatment	5.8% (5.3–6.3)

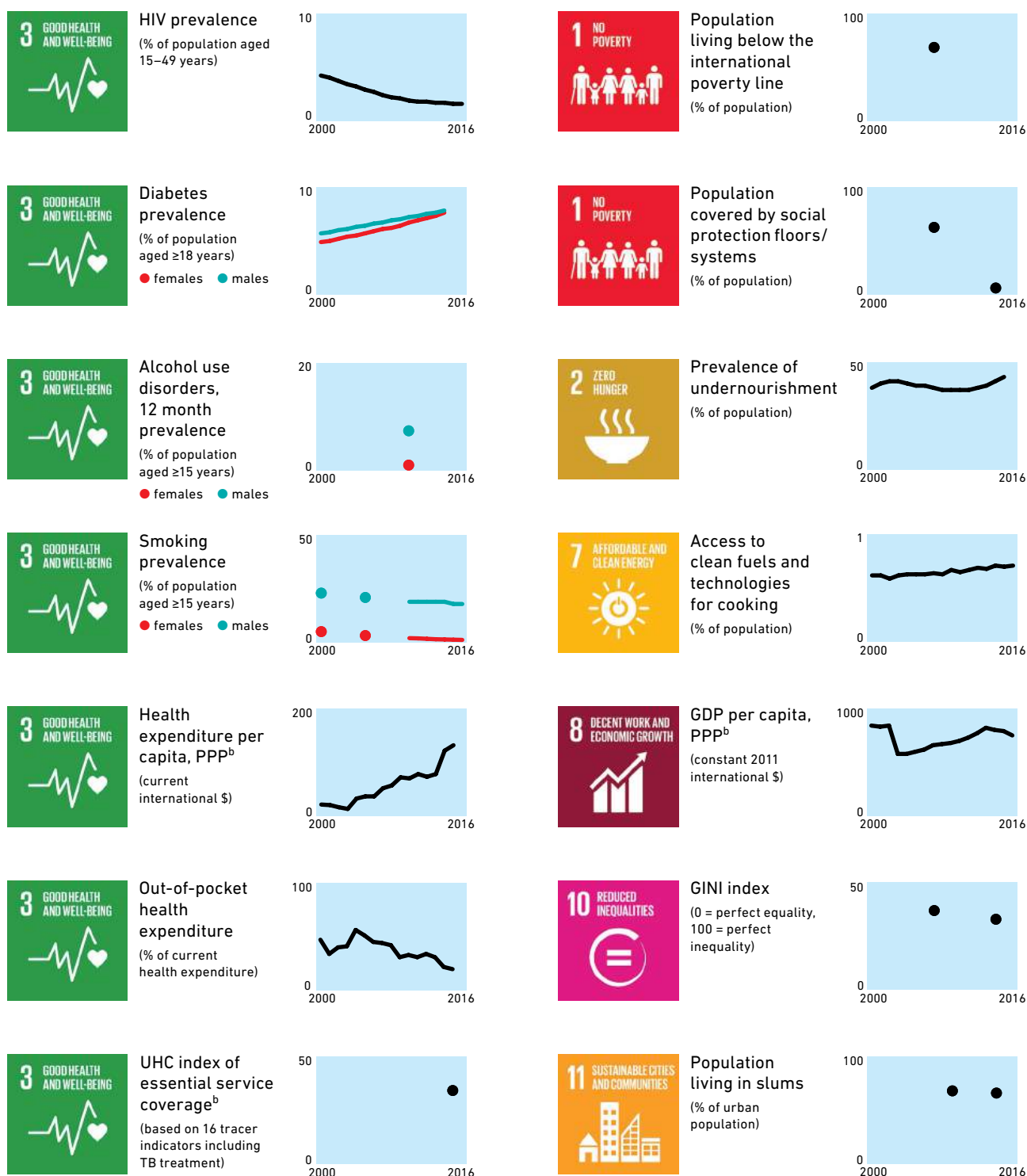
TB FINANCING, 2018

National TB budget (US\$ millions)	6.4
Funding source: 2% domestic, 98% international, 0% unfunded	

Data are as reported to WHO. Estimates of TB and MDR/RR-TB burden are produced by WHO in consultation with countries. Estimates are rounded and totals are computed prior to rounding.
^a Ranges represent uncertainty intervals.
^b MDR is TB resistant to rifampicin and isoniazid; RR is TB resistant to rifampicin.
^c Includes cases with unknown previous TB treatment history.
^d Includes patients diagnosed before 2017 and patients who were not laboratory-confirmed.



INDICATORS IN THE SUSTAINABLE DEVELOPMENT GOALS ASSOCIATED WITH TB INCIDENCE^a



Targets for reductions in TB incidence and TB deaths set in WHO's End TB Strategy and the United Nations' Sustainable Development Goals (SDGs) are ambitious. Achieving them requires progress in reducing health-related risk factors for TB infection and disease, as well as broader social and economic determinants of TB infection and disease. WHO has developed a TB-SDG monitoring framework that comprises 14 indicators under seven SDGs for which there is evidence of an association with TB incidence. Further details are provided in [Chapter 2](#).

^a Data sources: SDG indicators database, The World Bank, World Health Organization. Missing values and empty boxes indicate data not available in these data sources.

^b GDP = gross domestic product; PPP = purchasing power parity; UHC = universal health coverage

Namibia

POPULATION 2017 2.5 MILLION

ESTIMATES OF TB BURDEN, ^a 2017

	NUMBER (THOUSANDS)	RATE (PER 100 000 POPULATION)
Mortality (excludes HIV+TB)	0.75 (0.48–1.1)	30 (19–43)
Mortality (HIV+TB only)	0.8 (0.55–1.1)	31 (22–43)
Incidence (includes HIV+TB)	11 (8.2–14)	423 (324–535)
Incidence (HIV+TB only)	3.9 (2.5–5.5)	153 (99–219)
Incidence (MDR/RR-TB)**	0.95 (0.65–1.3)	37 (26–51)

ESTIMATED TB INCIDENCE BY AGE AND SEX (THOUSANDS), ^a 2017

	0–14 YEARS	> 14 YEARS	TOTAL
Females	0.88 (0.81–0.94)	3.6 (3.1–4.1)	4.4 (3.7–5.2)
Males	0.92 (0.85–0.99)	5.4 (4.4–6.3)	6.3 (5.1–7.5)
Total	1.8 (1.6–2)	8.9 (6.9–11)	11 (8.2–14)

TB CASE NOTIFICATIONS, 2017

Total cases notified	8 855
Total new and relapse	8 575
— % tested with rapid diagnostics at time of diagnosis	100%
— % with known HIV status	98%
— % pulmonary	82%
— % bacteriologically confirmed among pulmonary	83%

UNIVERSAL HEALTH COVERAGE AND SOCIAL PROTECTION

TB treatment coverage (notified/estimated incidence), 2017	80% (63–100)
TB patients facing catastrophic total costs	
TB case fatality ratio (estimated mortality/estimated incidence), 2017	0.15 (0.1–0.2)

TB/HIV CARE IN NEW AND RELAPSE TB PATIENTS, 2017

	NUMBER	(%)
Patients with known HIV-status who are HIV-positive	3 139	36%
— on antiretroviral therapy	3 021	96%

DRUG-RESISTANT TB CARE, 2017

	NEW CASES	PREVIOUSLY TREATED CASES	TOTAL NUMBER ^e
Estimated MDR/RR-TB cases among notified pulmonary TB cases			470 (410–540)
Estimated % of TB cases with MDR/RR-TB	5% (4.1–5.9)	12% (9.4–14)	
% notified tested for rifampicin resistance			
MDR/RR-TB cases tested for resistance to second-line drugs			78
Laboratory-confirmed cases			MDR/RR-TB: 409, XDR-TB: 14
Patients started on treatment ^d			MDR/RR-TB: 410, XDR-TB: 13

TREATMENT SUCCESS RATE AND COHORT SIZE

	SUCCESS	COHORT
New and relapse cases registered in 2016	84%	8 857
Previously treated cases, excluding relapse, registered in 2016	68%	297
HIV-positive TB cases registered in 2016	80%	3 410
MDR/RR-TB cases started on second-line treatment in 2015	74%	288
XDR-TB cases started on second-line treatment in 2015	0%	2

TB PREVENTIVE TREATMENT, 2017

% of HIV-positive people (newly enrolled in care) on preventive treatment	15%
% of children (aged < 5) household contacts of bacteriologically-confirmed TB cases on preventive treatment	34% (31–37)

TB FINANCING, 2018

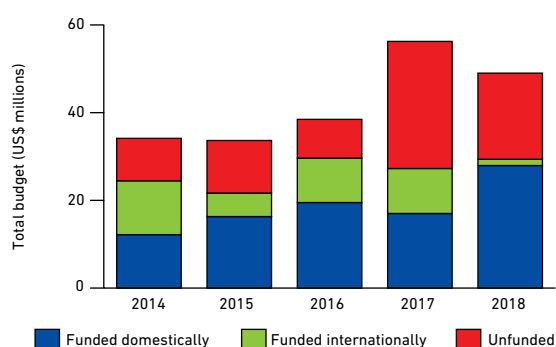
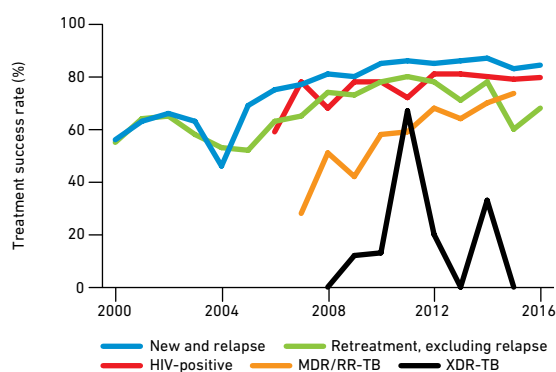
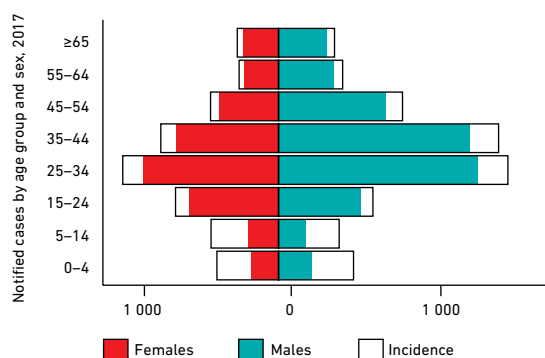
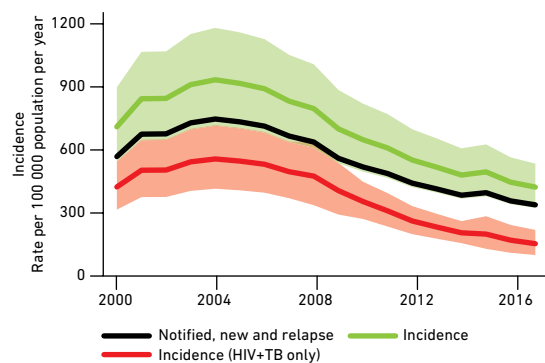
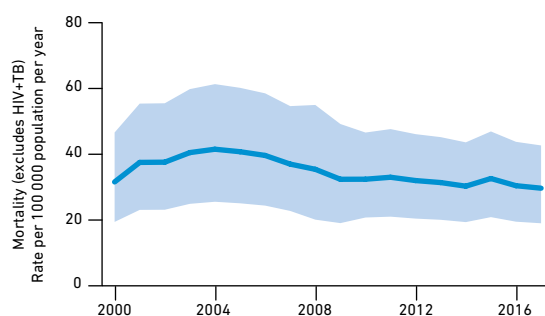
National TB budget (US\$ millions)	49
Funding source:	57% domestic, 3% international, 40% unfunded

Data are as reported to WHO. Estimates of TB and MDR/RR-TB burden are produced by WHO in consultation with countries. Estimates are rounded and totals are computed prior to rounding. ^a Ranges represent uncertainty intervals. Estimates of TB incidence and mortality for Namibia will be reviewed after final results from their national TB prevalence survey are available in 2019.

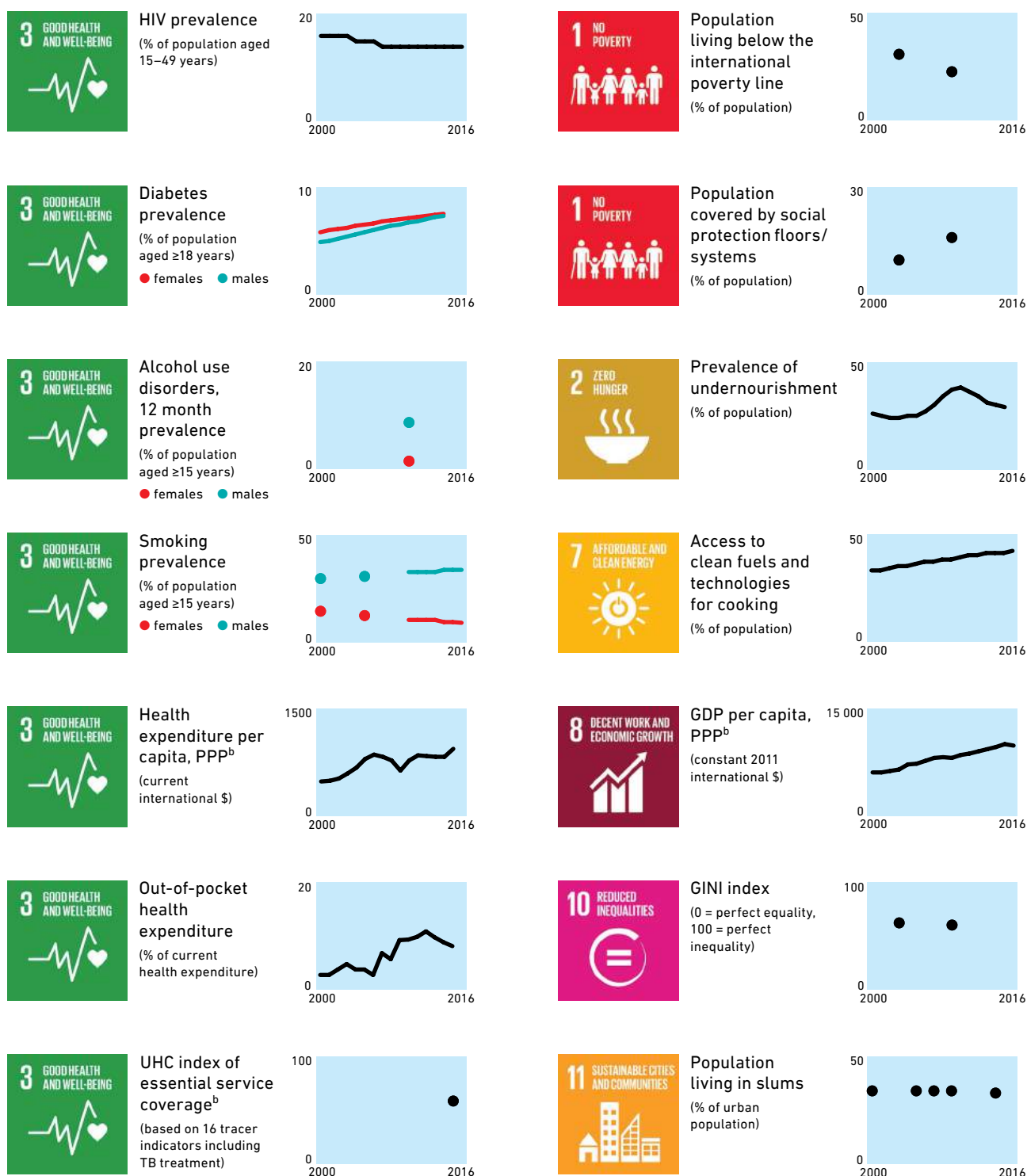
^b MDR is TB resistant to rifampicin and isoniazid; RR is TB resistant to rifampicin.

^c Includes cases with unknown previous TB treatment history.

^d Includes patients diagnosed before 2017 and patients who were not laboratory-confirmed.



INDICATORS IN THE SUSTAINABLE DEVELOPMENT GOALS ASSOCIATED WITH TB INCIDENCE^a



Targets for reductions in TB incidence and TB deaths set in WHO's End TB Strategy and the United Nations' Sustainable Development Goals (SDGs) are ambitious. Achieving them requires progress in reducing health-related risk factors for TB infection and disease, as well as broader social and economic determinants of TB infection and disease. WHO has developed a TB-SDG monitoring framework that comprises 14 indicators under seven SDGs for which there is evidence of an association with TB incidence. Further details are provided in [Chapter 2](#).

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^b GDP = gross domestic product; PPP = purchasing power parity; UHC = universal health coverage

Papua New Guinea

ESTIMATES OF TB BURDEN,^a 2017

	NUMBER (THOUSANDS)	RATE (PER 100 000 POPULATION)
Mortality (excludes HIV+TB)	4.3 (2.9–6)	53 (36–73)
Mortality (HIV+TB only)	0.93 (0.51–1.5)	11 (6.2–18)
Incidence (includes HIV+TB)	36 (29–43)	432 (352–521)
Incidence (HIV+TB only)	3.5 (2–5.5)	43 (24–66)
Incidence (MDR/RR-TB)**	1.9 (1.2–2.8)	23 (15–34)

ESTIMATED TB INCIDENCE BY AGE AND SEX (THOUSANDS),^a 2017

	0–14 YEARS	> 14 YEARS	TOTAL
Females	1.8 (1.7–1.9)	11 (9.7–12)	13 (11–14)
Males	2 (1.9–2.1)	21 (18–24)	23 (19–27)
Total	3.9 (3.6–4.1)	32 (26–38)	36 (29–43)

TB CASE NOTIFICATIONS, 2017

Total cases notified	27 934
Total new and relapse	26 294
— % tested with rapid diagnostics at time of diagnosis	
— % with known HIV status	45%
— % pulmonary	57%
— % bacteriologically confirmed among pulmonary	26%

UNIVERSAL HEALTH COVERAGE AND SOCIAL PROTECTION

TB treatment coverage (notified/estimated incidence), 2017	74% (61–91)
TB patients facing catastrophic total costs	
TB case fatality ratio (estimated mortality/estimated incidence), 2017	0.15 (0.1–0.21)

TB/HIV CARE IN NEW AND RELAPSE TB PATIENTS, 2017

	NUMBER	(%)
Patients with known HIV-status who are HIV-positive	791	7%
— on antiretroviral therapy	753	95%

DRUG-RESISTANT TB CARE, 2017

	NEW CASES	PREVIOUSLY TREATED CASES	TOTAL NUMBER ^e
Estimated MDR/RR-TB cases among notified pulmonary TB cases			960 (650–1 300)
Estimated % of TB cases with MDR/RR-TB	3.4% (1.7–5)	26% (15–36)	
% notified tested for rifampicin resistance	11%	56%	15 090
MDR/RR-TB cases tested for resistance to second-line drugs			161
Laboratory-confirmed cases		MDR/RR-TB: 356, XDR-TB: 13	
Patients started on treatment ^d		MDR/RR-TB: 356, XDR-TB: 15	

TREATMENT SUCCESS RATE AND COHORT SIZE

	SUCCESS	COHORT
New cases registered in 2016	62%	27 294
Previously treated cases registered in 2016	17%	2 457
HIV-positive TB cases registered in 2016		
MDR/RR-TB cases started on second-line treatment in 2015	68%	149
XDR-TB cases started on second-line treatment in 2015	64%	11

TB PREVENTIVE TREATMENT, 2017

% of HIV-positive people (newly enrolled in care) on preventive treatment	16%
% of children (aged < 5) household contacts of bacteriologically-confirmed TB cases on preventive treatment	

TB FINANCING, 2018

National TB budget (US\$ millions)	28
Funding source:	50% domestic, 21% international, 29% unfunded

Data are as reported to WHO. Estimates of TB and MDR/RR-TB burden are produced by WHO in consultation with countries. Estimates are rounded and totals are computed prior to rounding.

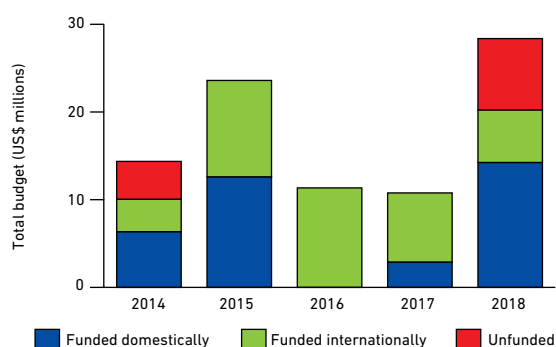
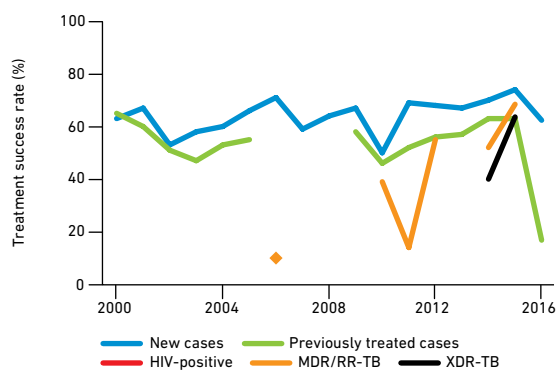
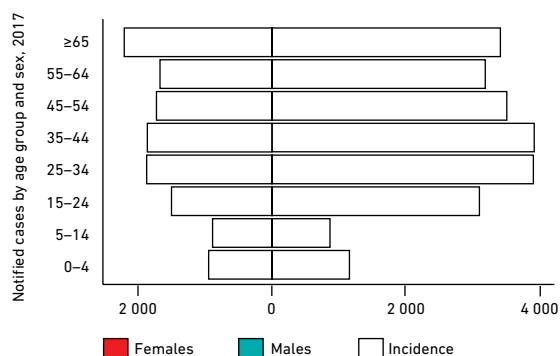
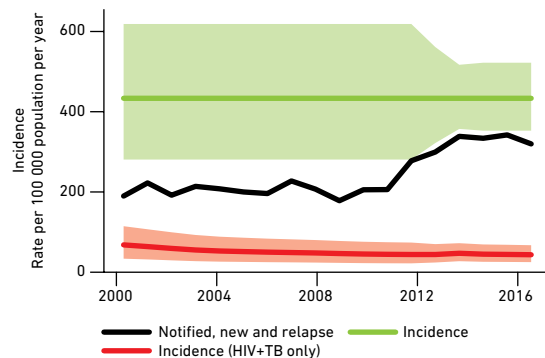
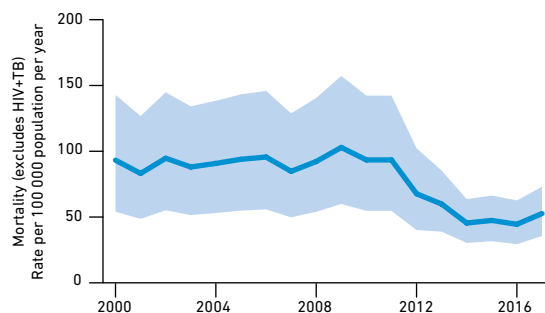
^a Ranges represent uncertainty intervals.

^b MDR is TB resistant to rifampicin and isoniazid; RR is TB resistant to rifampicin.

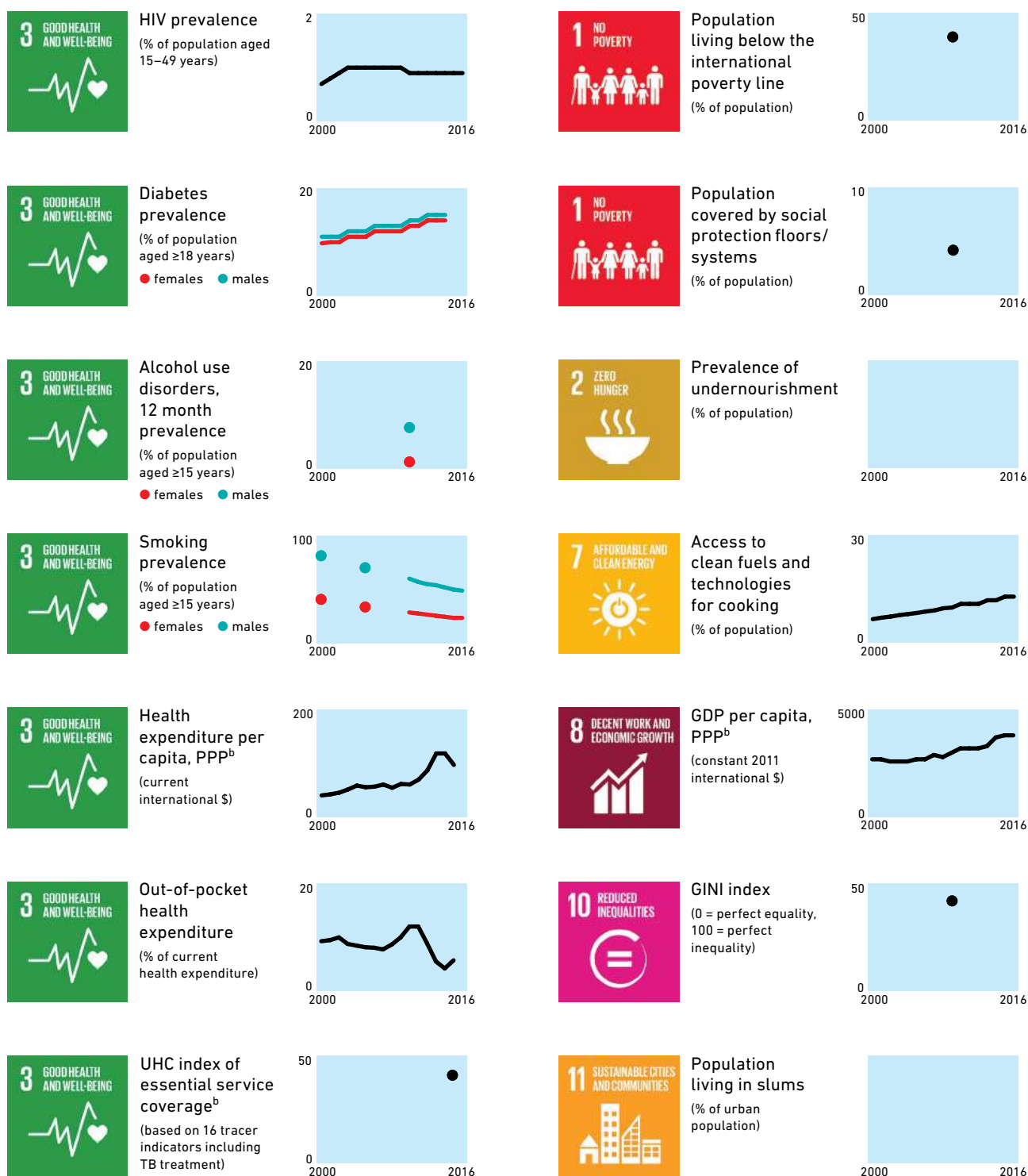
^c Includes cases with unknown previous TB treatment history.

^d Includes patients diagnosed before 2017 and patients who were not laboratory-confirmed.

POPULATION 2017 8.3 MILLION



INDICATORS IN THE SUSTAINABLE DEVELOPMENT GOALS ASSOCIATED WITH TB INCIDENCE^a



Targets for reductions in TB incidence and TB deaths set in WHO's End TB Strategy and the United Nations' Sustainable Development Goals (SDGs) are ambitious. Achieving them requires progress in reducing health-related risk factors for TB infection and disease, as well as broader social and economic determinants of TB infection and disease. WHO has developed a TB-SDG monitoring framework that comprises 14 indicators under seven SDGs for which there is evidence of an association with TB incidence. Further details are provided in [Chapter 2](#).

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^b GDP = gross domestic product; PPP = purchasing power parity; UHC = universal health coverage

Sierra Leone

POPULATION 2017 7.6 MILLION

ESTIMATES OF TB BURDEN, ^a 2017

	NUMBER (THOUSANDS)	RATE (PER 100 000 POPULATION)
Mortality (excludes HIV+TB)	3 (1.8–4.5)	39 (23–59)
Mortality (HIV+TB only)	0.78 (0.49–1.1)	10 (6.5–15)
Incidence (includes HIV+TB)	23 (15–33)	301 (193–431)
Incidence (HIV+TB only)	2.8 (1.8–4)	37 (24–53)
Incidence (MDR/RR-TB)**	0.66 (0.27–1.2)	8.8 (3.6–16)

ESTIMATED TB INCIDENCE BY AGE AND SEX (THOUSANDS), ^a 2017

	0–14 YEARS	> 14 YEARS	TOTAL
Females	1.2 (1.1–1.3)	8.1 (6.2–10)	9.3 (6.9–12)
Males	1.3 (1.2–1.4)	12 (8.6–16)	13 (9.3–17)
Total	2.5 (2.2–2.8)	20 (13–28)	23 (15–33)

TB CASE NOTIFICATIONS, 2017

Total cases notified	16 142
Total new and relapse	15 935
— % tested with rapid diagnostics at time of diagnosis	3%
— % with known HIV status	98%
— % pulmonary	93%
— % bacteriologically confirmed among pulmonary	65%

UNIVERSAL HEALTH COVERAGE AND SOCIAL PROTECTION

TB treatment coverage (notified/estimated incidence), 2017	70% (49–110)
TB patients facing catastrophic total costs	
TB case fatality ratio (estimated mortality/estimated incidence), 2017	0.17 (0.09–0.27)

TB/HIV CARE IN NEW AND RELAPSE TB PATIENTS, 2017

	NUMBER	(%)
Patients with known HIV-status who are HIV-positive	1 970	12%
— on antiretroviral therapy	1 911	97%

DRUG-RESISTANT TB CARE, 2017

	NEW CASES	PREVIOUSLY TREATED CASES	TOTAL NUMBER ^e
Estimated MDR/RR-TB cases among notified pulmonary TB cases			430 (200–660)
Estimated % of TB cases with MDR/RR-TB	2.5% (1.1–4.3)	14% (10–18)	
% notified tested for rifampicin resistance			456
MDR/RR-TB cases tested for resistance to second-line drugs			109
Laboratory-confirmed cases		MDR/RR-TB: 174, XDR-TB: 0	
Patients started on treatment ^d		MDR/RR-TB: 104, XDR-TB: 0	

TREATMENT SUCCESS RATE AND COHORT SIZE

	SUCCESS	COHORT
New and relapse cases registered in 2016	89%	12 497
Previously treated cases, excluding relapse, registered in 2016	72%	239
HIV-positive TB cases registered in 2016		
MDR/RR-TB cases started on second-line treatment in 2015		
XDR-TB cases started on second-line treatment in 2015		

TB PREVENTIVE TREATMENT, 2017

% of HIV-positive people (newly enrolled in care) on preventive treatment	22%
% of children (aged < 5) household contacts of bacteriologically-confirmed TB cases on preventive treatment	

TB FINANCING, 2018

National TB budget (US\$ millions)	16
Funding source:	2% domestic, 76% international, 23% unfunded

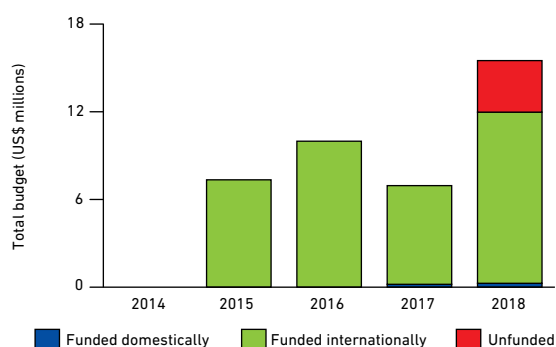
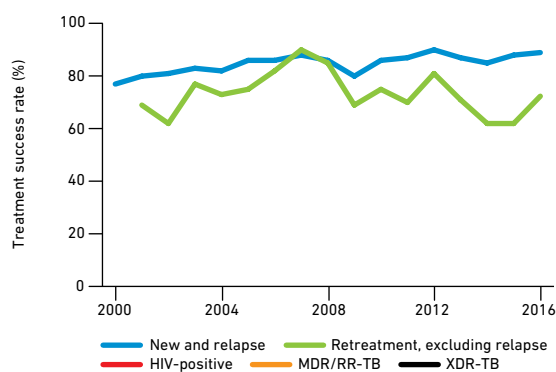
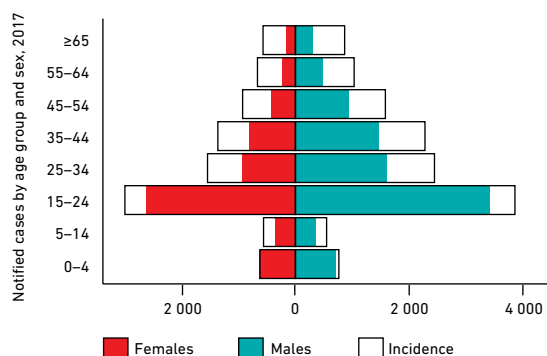
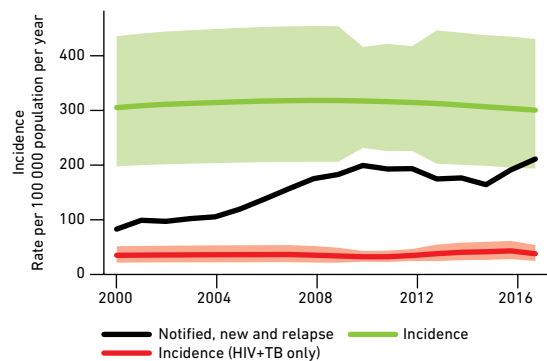
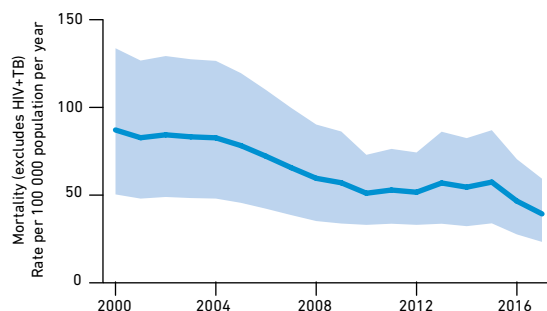
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^a Ranges represent uncertainty intervals.

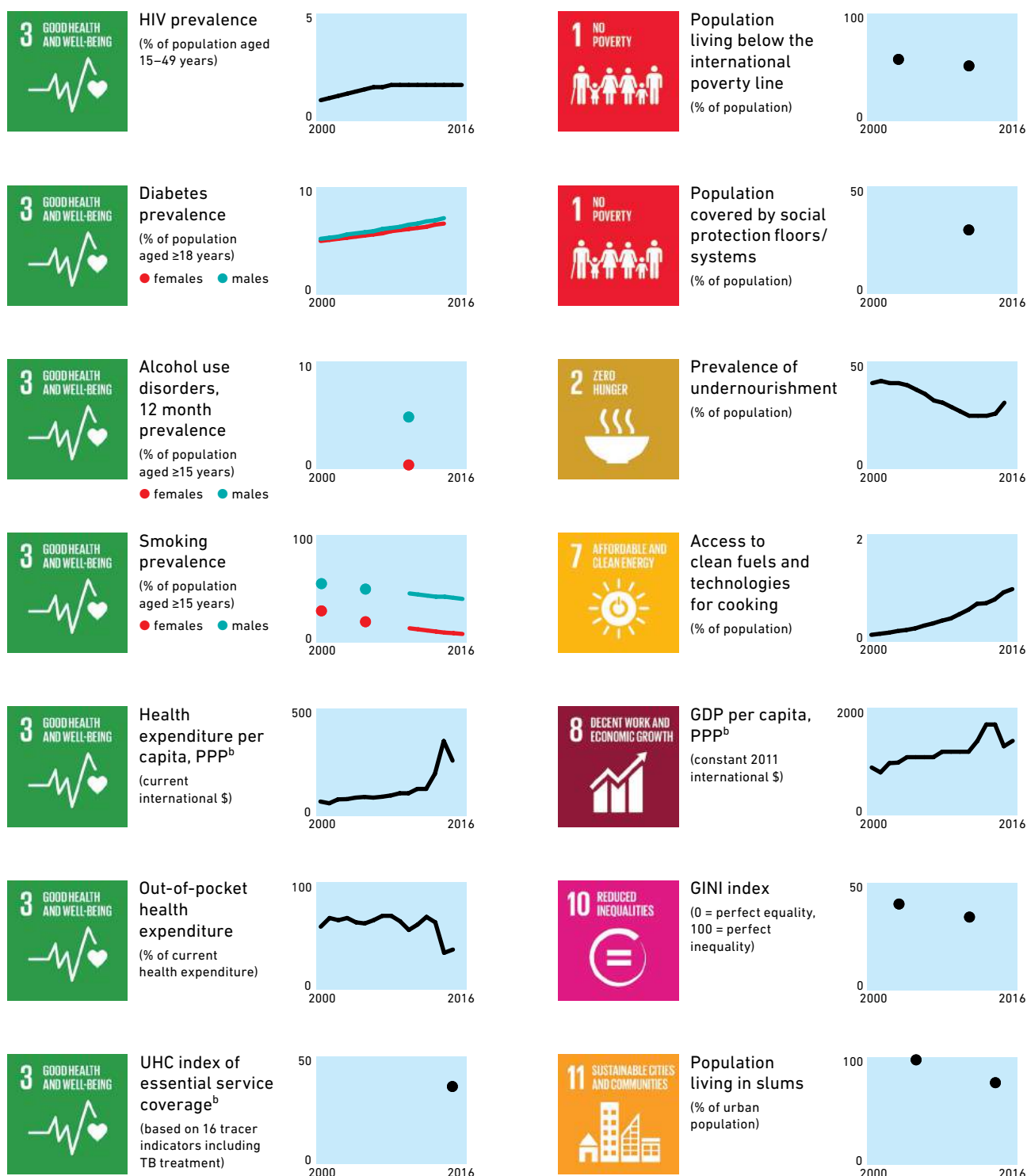
^b MDR is TB resistant to rifampicin and isoniazid; RR is TB resistant to rifampicin.

^c Includes cases with unknown previous TB treatment history.

^d Includes patients diagnosed before 2017 and patients who were not laboratory-confirmed.



INDICATORS IN THE SUSTAINABLE DEVELOPMENT GOALS ASSOCIATED WITH TB INCIDENCE^a



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^b GDP = gross domestic product; PPP = purchasing power parity; UHC = universal health coverage

Zambia

POPULATION 2017 **17 MILLION**

ESTIMATES OF TB BURDEN, ^a 2017

	NUMBER (THOUSANDS)	RATE (PER 100 000 POPULATION)
Mortality (excludes HIV+TB)	5 (2.9–7.7)	30 (17–45)
Mortality (HIV+TB only)	13 (8.2–19)	76 (48–110)
Incidence (includes HIV+TB)	62 (40–88)	361 (234–514)
Incidence (HIV+TB only)	36 (23–52)	210 (135–302)
Incidence (MDR/RR-TB) ^b	1.9 (0.67–3.8)	11 (3.9–22)

ESTIMATED TB INCIDENCE BY AGE AND SEX (THOUSANDS), ^a 2017

	0–14 YEARS	> 14 YEARS	TOTAL
Females	3.6 (3.2–3.9)	20 (16–25)	24 (18–30)
Males	4 (3.6–4.3)	34 (24–44)	38 (26–49)
Total	7.5 (6.5–8.5)	54 (34–74)	62 (40–88)

TB CASE NOTIFICATIONS, 2017

Total cases notified	37 203
Total new and relapse	36 010
— % tested with rapid diagnostics at time of diagnosis	28%
— % with known HIV status	93%
— % pulmonary	84%
— % bacteriologically confirmed among pulmonary	53%

UNIVERSAL HEALTH COVERAGE AND SOCIAL PROTECTION

TB treatment coverage (notified/estimated incidence), 2017	58% (41–90)
TB patients facing catastrophic total costs	
TB case fatality ratio (estimated mortality/estimated incidence), 2017	0.3 (0.17–0.46)

TB/HIV CARE IN NEW AND RELAPSE TB PATIENTS, 2017

	NUMBER	(%)
Patients with known HIV-status who are HIV-positive	20 362	59%
— on antiretroviral therapy	18 341	90%

DRUG-RESISTANT TB CARE, 2017

	NEW CASES	PREVIOUSLY TREATED CASES	TOTAL NUMBER ^c
Estimated MDR/RR-TB cases among notified pulmonary TB cases			1 300 (850–1 800)
Estimated % of TB cases with MDR/RR-TB	1.1% (0.34–2.3)	18% (12–26)	
% notified tested for rifampicin resistance	23%	9%	10 591
MDR/RR-TB cases tested for resistance to second-line drugs			0
Laboratory-confirmed cases		MDR/RR-TB: 546, XDR-TB: 0	
Patients started on treatment ^d		MDR/RR-TB: 270, XDR-TB: 0	

TREATMENT SUCCESS RATE AND COHORT SIZE

	SUCCESS	COHORT
New and relapse cases registered in 2016	88%	38 326
Previously treated cases, excluding relapse, registered in 2016	79%	1 827
HIV-positive TB cases registered in 2016	86%	21 655
MDR/RR-TB cases started on second-line treatment in 2015	60%	99
XDR-TB cases started on second-line treatment in 2015		0

TB PREVENTIVE TREATMENT, 2017

% of HIV-positive people (newly enrolled in care) on preventive treatment	18%
% of children (aged < 5) household contacts of bacteriologically-confirmed TB cases on preventive treatment	3.8% (3.5–4.2)

TB FINANCING, 2018

National TB budget (US\$ millions)	38
Funding source:	28% domestic, 52% international, 20% unfunded

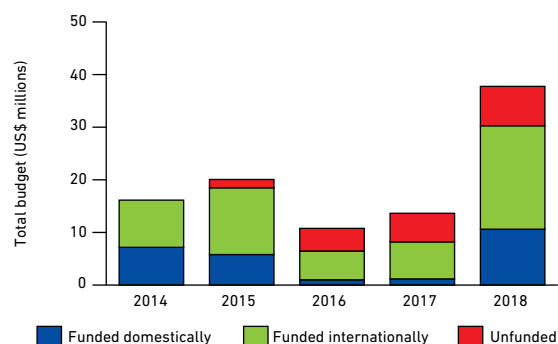
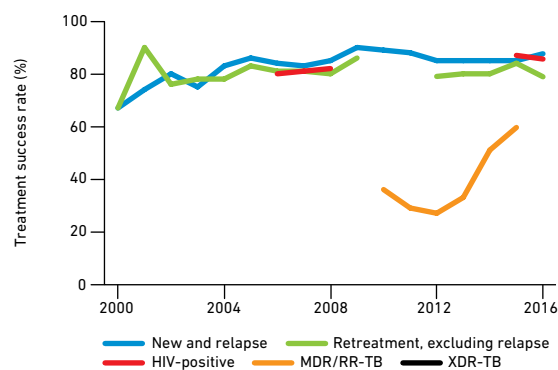
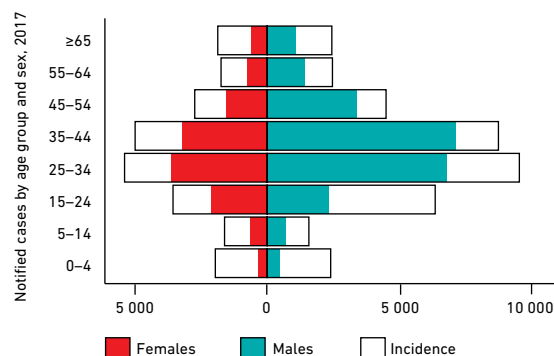
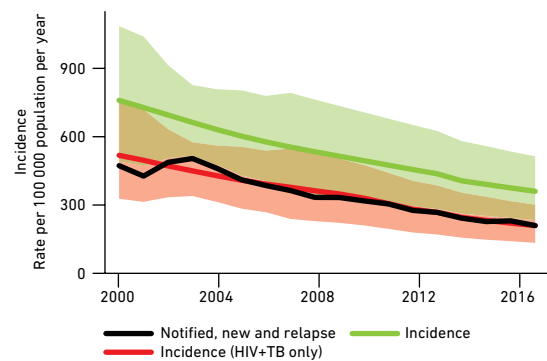
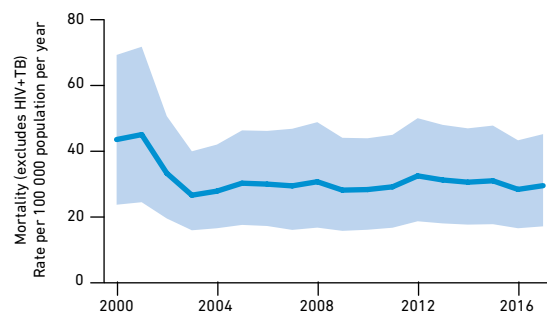
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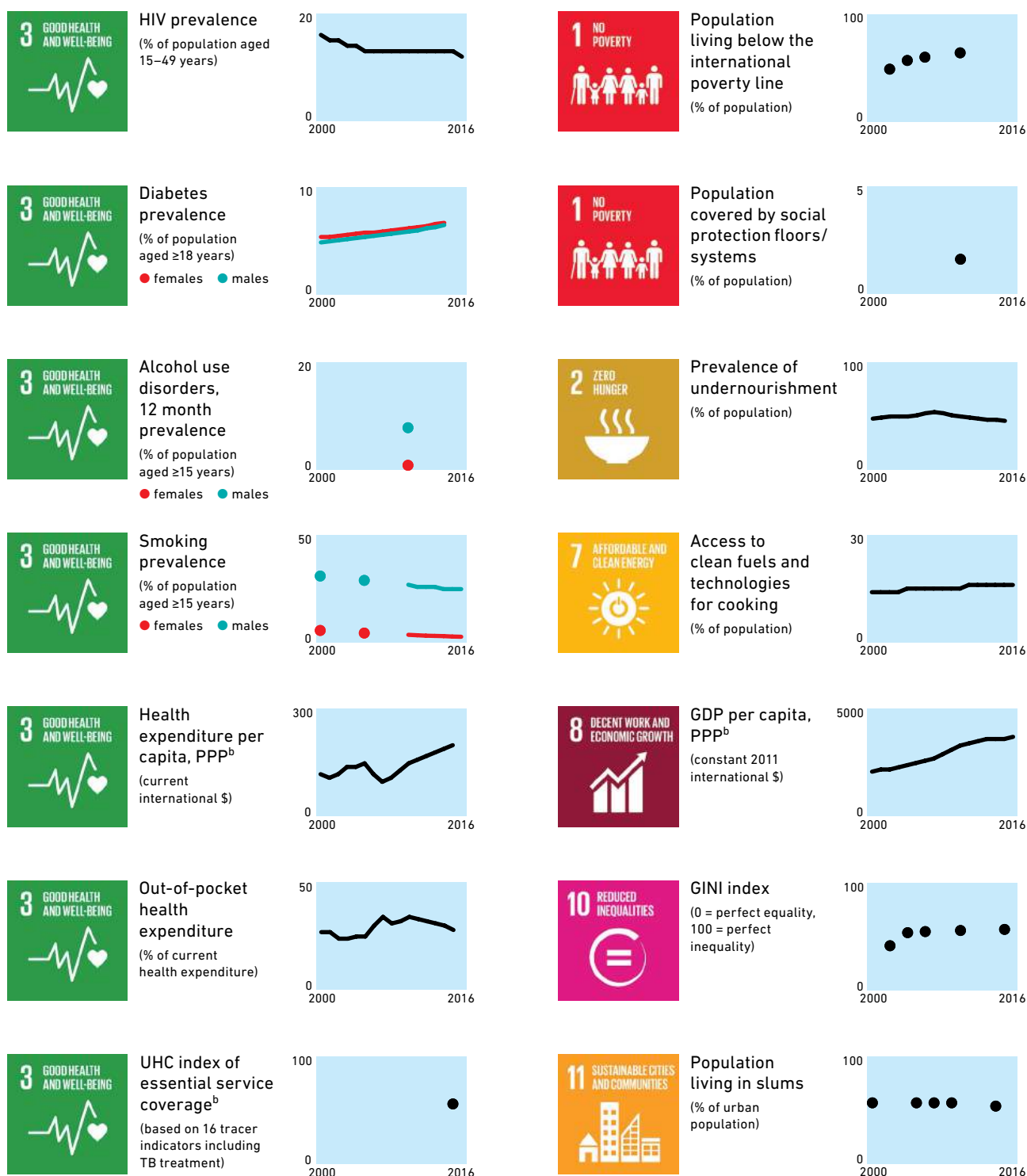
^b MDR is TB resistant to rifampicin and isoniazid; RR is TB resistant to rifampicin.

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^d Includes patients diagnosed before 2017 and patients who were not laboratory-confirmed.



INDICATORS IN THE SUSTAINABLE DEVELOPMENT GOALS ASSOCIATED WITH TB INCIDENCE^a



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^b GDP = gross domestic product; PPP = purchasing power parity; UHC = universal health coverage

Zimbabwe

POPULATION 2017 **17 MILLION**

ESTIMATES OF TB BURDEN, ^a 2017

	NUMBER (THOUSANDS)	RATE (PER 100 000 POPULATION)
Mortality (excludes HIV+TB)	2 (1.3–2.9)	12 (7.7–17)
Mortality (HIV+TB only)	6.3 (4.5–8.5)	38 (27–51)
Incidence (includes HIV+TB)	37 (27–47)	221 (164–287)
Incidence (HIV+TB only)	23 (15–33)	140 (90–199)
Incidence (MDR/RR-TB) ^b	2.3 (1.4–3.5)	14 (8.5–21)

ESTIMATED TB INCIDENCE BY AGE AND SEX (THOUSANDS), ^a 2017

	0–14 YEARS	> 14 YEARS	TOTAL
Females	2.6 (2.4–2.8)	13 (11–15)	15 (12–18)
Males	2.9 (2.7–3.1)	18 (15–22)	21 (17–26)
Total	5.6 (4.9–6.2)	31 (23–39)	37 (27–47)

TB CASE NOTIFICATIONS, 2017

Total cases notified	26 401
Total new and relapse	25 848
— % tested with rapid diagnostics at time of diagnosis	
— % with known HIV status	100%
— % pulmonary	89%
— % bacteriologically confirmed among pulmonary	58%

UNIVERSAL HEALTH COVERAGE AND SOCIAL PROTECTION

TB treatment coverage (notified/estimated incidence), 2017	71% (55–96)
TB patients facing catastrophic total costs	
TB case fatality ratio (estimated mortality/estimated incidence), 2017	0.23 (0.15–0.32)

TB/HIV CARE IN NEW AND RELAPSE TB PATIENTS, 2017

	NUMBER	(%)
Patients with known HIV-status who are HIV-positive	16 602	63%
— on antiretroviral therapy	14 300	86%

DRUG-RESISTANT TB CARE, 2017

	NEW CASES	PREVIOUSLY TREATED CASES	TOTAL NUMBER ^c
Estimated MDR/RR-TB cases among notified pulmonary TB cases			1 300 (920–1 600)
Estimated % of TB cases with MDR/RR-TB	4.6% (3–6.2)	14% (8.9–20)	
% notified tested for rifampicin resistance			7 385
MDR/RR-TB cases tested for resistance to second-line drugs			
Laboratory-confirmed cases		MDR/RR-TB: 474, XDR-TB: 4	
Patients started on treatment ^d		MDR/RR-TB: 439, XDR-TB: 3	

TREATMENT SUCCESS RATE AND COHORT SIZE

	SUCCESS	COHORT
New and relapse cases registered in 2016	81%	26 618
Previously treated cases, excluding relapse, registered in 2016	70%	735
HIV-positive TB cases registered in 2016	78%	16 520
MDR/RR-TB cases started on second-line treatment in 2015	44%	433
XDR-TB cases started on second-line treatment in 2015	80%	5

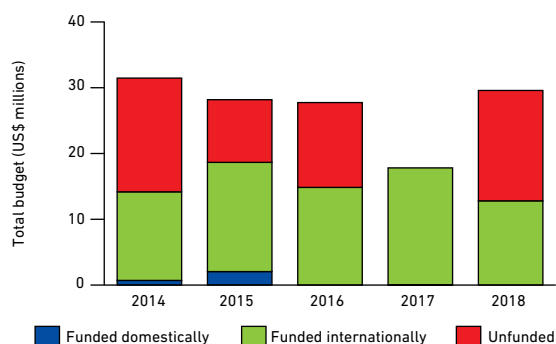
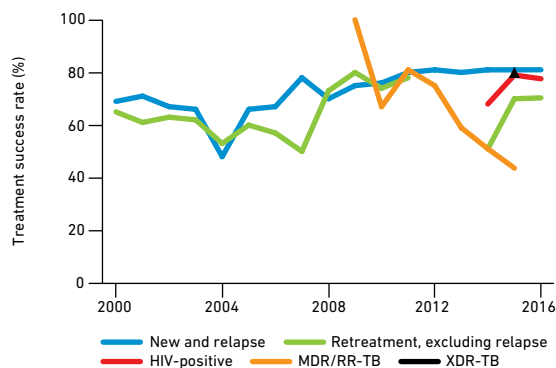
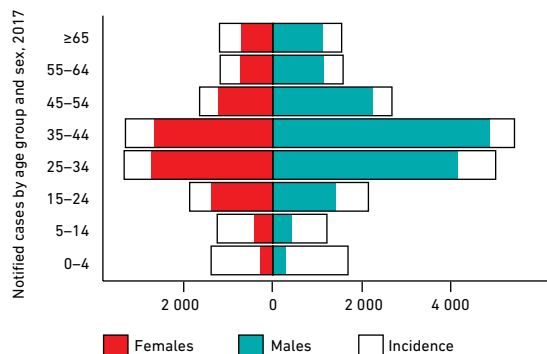
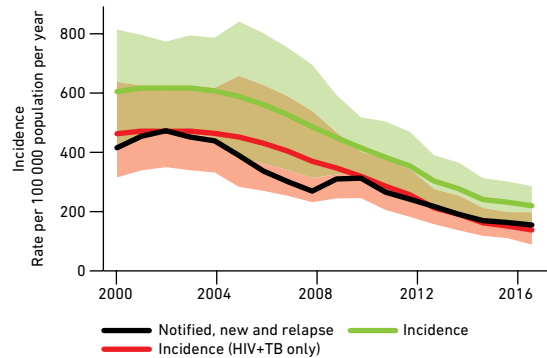
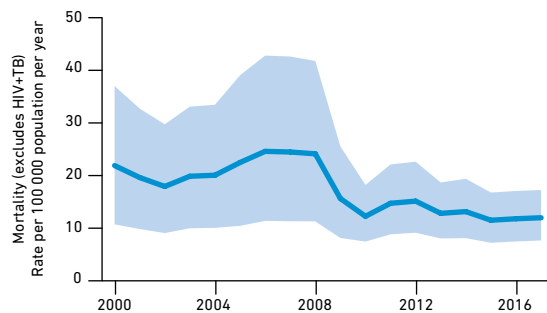
TB PREVENTIVE TREATMENT, 2017

% of HIV-positive people (newly enrolled in care) on preventive treatment ^e	11%
% of children (aged < 5) household contacts of bacteriologically-confirmed TB cases on preventive treatment	24% (22–26)

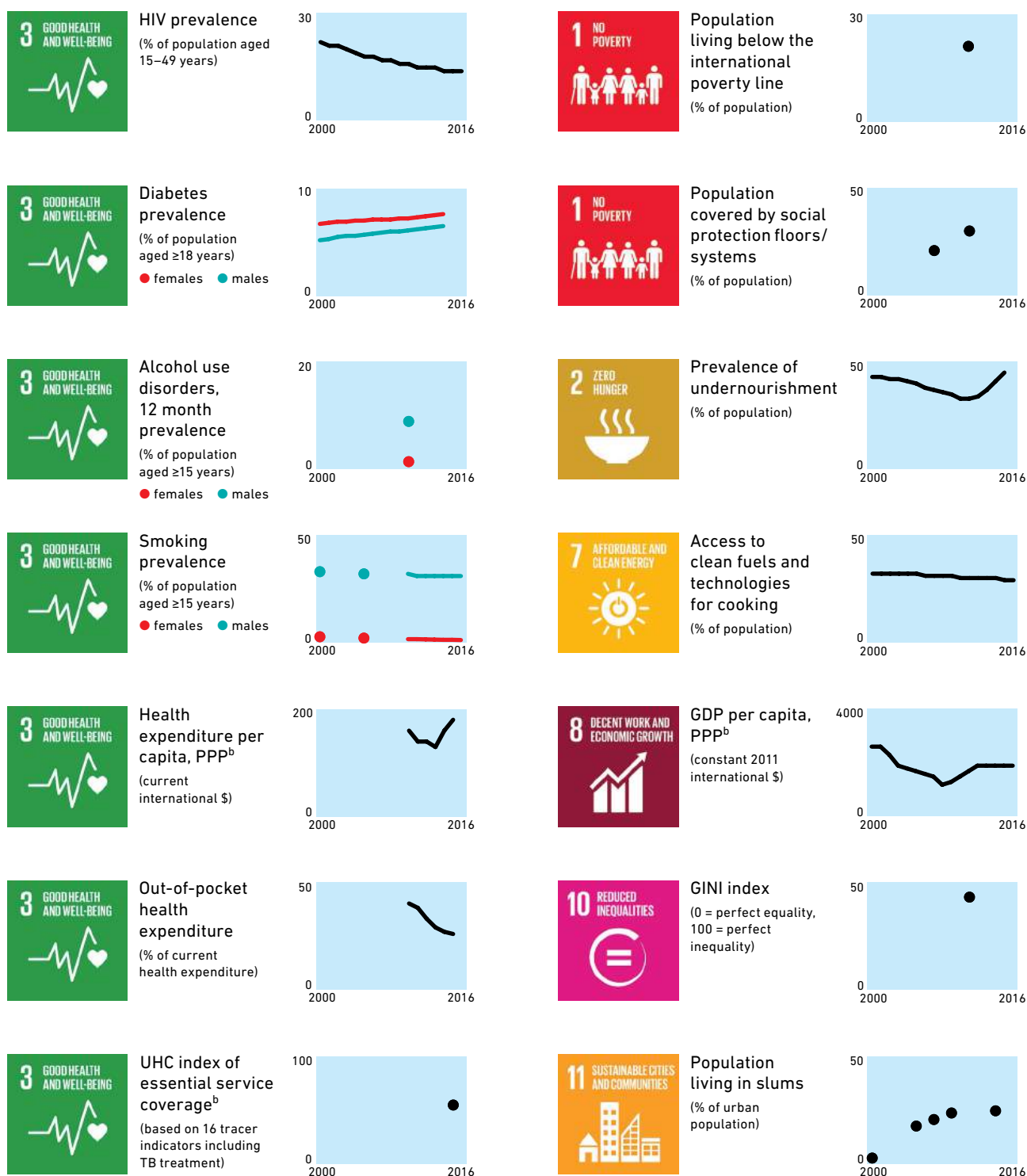
TB FINANCING, 2018

National TB budget (US\$ millions)	30
Funding source:	domestic, 43% international, 57% unfunded

Data are as reported to WHO. Estimates of TB and MDR/RR-TB burden are produced by WHO in consultation with countries. Estimates are rounded and totals are computed prior to rounding.
^a Ranges represent uncertainty intervals.
^b MDR is TB resistant to rifampicin and isoniazid; RR is TB resistant to rifampicin.
^c Includes cases with unknown previous TB treatment history.
^d Includes patients diagnosed before 2017 and patients who were not laboratory-confirmed.
^e The number on preventive treatment was provided for the period July–December 2017 only.



INDICATORS IN THE SUSTAINABLE DEVELOPMENT GOALS ASSOCIATED WITH TB INCIDENCE^a



Targets for reductions in TB incidence and TB deaths set in WHO's End TB Strategy and the United Nations' Sustainable Development Goals (SDGs) are ambitious. Achieving them requires progress in reducing health-related risk factors for TB infection and disease, as well as broader social and economic determinants of TB infection and disease. WHO has developed a TB-SDG monitoring framework that comprises 14 indicators under seven SDGs for which there is evidence of an association with TB incidence. Further details are provided in [Chapter 2](#).

^a Data sources: SDG indicators database, The World Bank, World Health Organization. Missing values and empty boxes indicate data not available in these data sources.

^b GDP = gross domestic product; PPP = purchasing power parity; UHC = universal health coverage



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