

MedPharmPlast Europe Summer Conference 2018 04-05 July 2018 - Lyon, MARRIOTT Lyon Cite Internationale

Material Characterization & Risk Assessment

in collaboration with **WNelson Labs**. EUROPE FORMERLY TOXIKON EUROPE

PROGRAMME

Wednesday, 4 July

10:00 – 10:30	Registration & Welcome Coffee
10:30 – 10:45	Welcome - James Stern, President MedPharmPlast Europe
10:45 – 11:45	Changes to the ISO 10993 - Thor Rollins, ISO 10993 Committee
11:45 – 12:00	Short Coffee Break
12:00 – 12:45	Selection of Raw Material for Risk Reduction - Part 1 Launch of new guideline (VDI) for medical grade plastics - Stefan Roth, VDI Committee, Schmalkalden University of Applied Sciences
12:45 – 13:45	Networking Lunch
13:45 – 14:45	 Selection of Raw Material for Risk Reduction – Part 2 Managing Risk and Consistency in the Raw Material Supply Chain – a case study - Nicholas Berendt, Sealed Air Regulatory Updates on Developments in EUP/ USP - Frank De Smedt, Nelson Labs
14:45– 15:15	Networking Coffee
15:15 – 16:15	Toxicological Assessment of Extractables & Leachables o General Principles - Aurélien Bignon, Biom Advice (tbc) o Case Study - Pascale Van Hoydonck, Terumo
16:15 – 16 :45	MedPharmPlast Europe - Your Voice in Brussels - James Stern
16:45 – 17:00	Wrap-up & Closing
17:00 – 18:30	Networking Drinks
	Media Partner







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Thursday, 5 July

08:30 – 09:15	Registration & Welcome Coffee
09:15 – 10:00	Biocompatibility - Thor Rollins, ISO 10993 Committee
10:00 – 10:30	Development of an in-vitro testing battery to assess biocompatibility of medical devices - Elisabeth Mertl, OFI Technologie & Innovation GmbH
10:30 – 11:00	Networking Coffee
11:00 – 11:30	Implementation of the Medical Device Regulation: View of the Medical Device Producers - Nathalie Buijs, MedTech Europe (tbc)
11:30 – 12:15	Sustainability Issues in the Plastics Industry o EU Plastics Strategy/Toxic Free Environment Strategy - Silvia Freni Sterrantino, EuPC o View from the industry - Christian Meusinger, Nemera
12:15 – 12:30	Wrap-up & Closing



