

9:00 - 10:00	Registration & Welcome Coffee
10:00 - 10:15	Welcome by Christian Meusinger, MedPharmPlast Europe President
10:15 - 12:00	<p>THE MEDICAL DEVICE REGULATION & ITS IMPLEMENTATION:</p> <ul style="list-style-type: none"> A state of the art presentation <i>- Adrian Bartlett: Medical Devices, EU Policy Manager, Medicines and Healthcare Products Regulatory Agency (MHRA)</i> The consequences for the relationship between suppliers & converters <i>- Peter Curle: Executive Director, EMEA Advisory Center for Risk/GRC, Ernst & Young AG</i>
12:00 - 12:30	<p>Update on MedPharmPlast Europe regulatory activities</p> <p><i>- Nigel Talboys: Global Director Public Policy and Government Affairs, Terumo BCT / Chairman MPPE Regulatory Task Force</i></p>
12:30 - 13:30	Networking Lunch
13:30 - 14:00	<p>How 3D printing will change the medical landscape</p> <p><i>- Ward Callens: Director of Quality, Materialise</i></p>
14:00 - 14:30	<p>Future Trends In The Medical Devices Industry</p> <p><i>- Antonella Lisella: Consulting Analyst, Frost & Sullivan</i></p>
14:30 - 15:00	Coffee Break
15:00 - 15:30	<p>The changes in testing requirements related to the revision of the USP <661> and comparison to European Pharmacopoeia</p> <p><i>- Frank de Smedt: Department Head Analytical Services, Toxikon Europe NV</i></p>
15:30 - 16:00	<p>Benefits of MedPharmPlast Europe membership and its European network</p> <p><i>- Béatrice Grand Demars: Regulatory & Compliance Manager, Nemera</i></p>
	Closing