

# Introduction to Chemical Compliance for Medical Devices

## Program

08.30 - 09.00	Registration and breakfast
09.00 - 09.10	Welcome by Medicoindustrien
09.10 - 09.25	Presentation and introduction
09.25 - 10.10	Regulatory requirements - MDR/ISO
10.10 - 10.30	Coffee break
10.30 - 11.00	Test strategy for chemical characterisation – ISO 10993-18
11.00 - 11.50	How to perform chemical characterization – ISO 10993-18
11.50 - 12.00	Questions /discussions
12.00 - 13.00	Lunch
13.00 - 13.40	Short introduction to toxicology
13.40 - 14.20	Introduction of ISO framework for safety assessment of medical devices
14.20 - 14.40	Coffee break
14.40 - 15.20	Safety assessment of the extracted chemicals, new ISO standard
15.20 - 15.40	How to collect relevant toxicological data for safety assessment
15.40 - 15.50	Questions/discussion
15.50 - 16.00	Wrap up and conclusions