

**EUROTOX 2025 workshop: “Advancing Risk Assessment and Compliance with European Regulations” –
Timetable for online lectures (Central European Time; pm)**

May 5		May 7		May 8			
6.00 – 6.45	Emanuela Corsini, Marc Pallardy, Anja Haveric <i>Introduction to the workshop Ice breaker</i>	6.00 – 6.45	Yalcın Duydu <i>Integrated testing strategies; decision tree; Threshold of Toxicological Concern (TTC)</i>	6.00 – 6.50	Corrado Lodovico Galli <i>Genotoxicity – a landmark in cosmetic safety assessment</i>		
6.50 – 7.35	Corrado Lodovico Galli <i>Risk analysis paradigm</i>	6.50 – 7.35	Danijela Djukic Cosic <i>In-silico method with toxicogenomic data mining</i>	7.00 – 7.50	Alicia Painsi <i>In vitro to in vivo extrapolation (IVIVE), quantitative IVIVE and route-to-route extrapolation to advance chemical risk assessment</i>		
7.40 – 8.25	Emanuela Corsini <i>Alternative methods in immunotoxicology</i>	7.40 – 8.25	Nursen Ayse Basaran <i>Using in vitro methods in hazard and risk assessment</i>	7.40 – 8.25			
May 12		May 14		May 15		May 16	
6.00 – 6.55	Tzveta Georgieva <i>New approach methodologies (NAMs) in toxicology</i>	6.00 – 6.55	Marijana Curcic <i>Dose-response analyses; derivation and use of health-based guidance values (e.g. RfD, ADI, AOEL, DNEL etc.); benchmark dose modelling; margin of exposure</i>	6.00 – 6.55	Corrado Lodovico Galli <i>Safety assessment of mixtures</i>	6.00 – 7.30	Thomas Hartung <i>Toxicology – the transformative potential of AI in toxicology</i>
7.00 – 7.55	Gonca Cakmak <i>Risk management and risk communication in the regulation of occupational agents</i>	7.00 – 7.55	Marijana Curcic <i>International classification schemes (e.g. CLP)</i>	7.00 – 7.55	Danijela Djukic Cosic <i>Application of regulations, guidelines and risk assessment for cosmetics</i>		
May 19		May 21		May 22			
6.00 – 6.55	Doris Marko <i>Challenges in risk assessment of mycotoxins</i>	6.00 – 6.55	Biljana Antonijevic <i>Guidelines for the safety assessment of pharmaceutical impurities</i>	6.00 – 6.55	Zorica Bulat <i>Toxicological risks of consumer products - application of regulations and guidelines</i>		
7.00 – 7.55	Maria Dusinska <i>Risk assessment of nano and advanced materials</i>	7.00 – 7.55	Nursen Ayse Basaran <i>Application of regulations, guidelines and risk assessment for pesticides</i>	7.00 – 7.55	Marc Pallardy <i>Regulatory requirements for non-clinical development to support clinical studies</i>		
May 26		May 28		May 29			
6.00 – 6.55	Dominique Brees <i>Translational safety assessment, bridging the gap between animal and human studies</i>	6.00 – 7.55	Michelle Embry & Sandrine Déglin <i>Risk assessment case studies</i>	6.00 – 7.55	Michelle Embry & Sandrine Déglin <i>Risk assessment case studies</i>		
7.00 – 7.55	Martin F. Wilks <i>Mode of action and Adverse Outcome Pathway (AOP) frameworks in risk assessment – the same or different?</i>						