

Monday, 27 Se	Monday, 27 September 2021	
08h00-20h00	Exhibition & Poster Viewing	
08h00-12h00	Continuing Education Courses (CEC)	
08h00-12h00	CEC01 Thyroid hormones, brain development and toxicity testing Chairs: Marta Axelstad, Denmark, Manon Beekhuijzen, Netherlands and Barbara Demeneix, France	
	Thyroid hormone action and disruption during development: pregnancy, brain and rat versus human Barbara Demeneix, UMR 7221 Molecular Physiology and Adaptation (CNRS/MNHN), Paris, France	
	Low thyroid hormone during pregnancy and consequences for child neurological development Peter Taylor, Cardiff University, Cardiff, UK	
	Safeguarding the thyroid system – developing an in vitro testing battery Sharon Munn, European Commission, Joint Research Centre, Ispra, Italy	
	Recommendations for the future: lessons learned from thyroid hormone determinations in OECD/ US EPA guideline studies Abby Li, Exponent Inc., San Francisco, US	
	Searching for an adverse effect endpoint in the developing brain Louise Ramhøj, Technical University of Denmark, Kgs. Lyngby, Denmark	
	Current in vivo guideline testing for endocrine disruption: practical issues and interpretation challenges with focus on thyroid hormones Manon Beekhuijzen, Charles River, Den Bosch, Netherlands	
	Panel Discussion	
08h00-12h00	CEC02 Advances in conducting systematic reviews for chemical assessment: automation, uncertainty assessment and synthesis Chairs: Andrew Rooney, US and Sebastian Hoffmann, Germany	
	LECTURE: Global orientation to systematic review – successes, challenges, and preparing for next generation decision making on mechanistic data	



Elisa Aiassa, Assessment and Methodological Support Unit/ EFSA, Parma, Italy

HANDS ON ACTIVITY 1: PECO, Inclusion/Exclusion

LECTURE: Automated and semi-automated approaches for literature searching, screening, and data extraction for systematic reviews in environmental health

Vickie Walker, National Institute of Environmental Health Sciences, Research Triangle Park, US

HANDS ON ACTIVITY 2: Critical appraisal/Risk of Bias

LECTURE: Synthesis, certainty (GRADE), and qualitative integration of human, animal, and mechanistic data Sebastian Hoffmann, Evidence-based Toxicology Collaboration (EBTC), Paderborn, Germany

LECTURE: Quantitative evidence integration supporting toxicity value development and characterization of uncertainty Daniele Wikoff, ToxStrategies, Asheville, US

HANDS ON ACTIVITY 3: Evidence certainty/GRADE

LECTURE: Conduct and reporting standards for systematic reviews in toxicology and risk assessmentPaul Whaley, Lancaster University, Lancaster Environment Centre, Lancaster, UK

08h00-12h00

CEC03 (subject to updates)

Lessons learned and future directions for toxicology in water safety and security

Chairs: Heidi Foth, Germany and Elaine Faustman, US

Water bodies and frame work for protection

Heidi Foth, Martin Luther University, Institute of Environmental Toxicology, Halle (Saale), Germany

Contamination in groundwater by overuse of fertilizers and implications for human health Speaker tba.

Contamination pattern by pesticides in water

Aristidis Tsatsakis, University of Crete, Greece

Arsenite in drinking water

Louis Schiesari, University of Sao Paolo, Brazil

Ecotoxicological assessment of pharmaceuticals and personal care products using predictive toxicology approaches

Susanne Boutrup and Hans Sanderson DCE (NERI) - Department of Environmental Science, Denmark



	Dissipative use of lead a future risk for groundwater Thomas Schupp, FH Münster – University of Applied Sciences, Steinfurt, Germany Bromate in bathing water – a carcinogenic risk? Speaker tba. Metagenomic approaches for surveillance and testing water Elaine M. Faustman, US
12h00-13h00	Virtual Lunch Break
13h00-14h00	Opening Ceremony incl. EUROTOX Merit Award
14h00-15h00	Bo Holmstedt Memorial Fund Lecture Chair: Heather Wallace, EUROTOX President Context matters: next generation insights on the chemistry of DNA damage and mutation Shana Sturla, ETH Zürich, Zurich, Switzerland
15h00-15h30	Virtual Coffee Break, Exhibition & Poster Viewing/Discussion
15h30–17h30	Session 01 – Symposium The effect of chemicals on the gut microbiota: is it the cause of all problems? Chair: Reinhilde Schoonjans, Italy Gut microbiota and human health through lifespan Anne Salonen, University of Helsinki, Helsinki, Finland Dietary emulsifiers, human microbiota and intestinal inflammation Tom Van de Wiele, Ghent University, Ghent, Belgium Low calorie sweeteners and their impact on the gut microbiota Ian Rowland, Reading University, Reading, UK
	The role of diet on stability, resilience and modulation of the human gut microbiota



	Carmen Peláez, Autonomous University of Madrid, Madrid, Spain
15h30–17h30	Session 02 – Symposium In vitro organotypic models for predicting the toxicity of chemicals or drugs Chairs: Saadia Kerdine-Römer, France and Lisbeth Knudsen, Denmark
	Predictive (diseased) 3D lung models to assess effects of aerosolized nanomaterials and nanodrugs Barbara Rothen Rutishauser, Université de Fribourg, Fribourg, Switzerland
	Advanced in vitro models for nephrotoxicity testing: as complex as possible, but simple in use Rosaline Masereeuw, Utrecht Institute for Pharmaceutical Sciences, Utrecht, Netherlands
	Human 3D brain model to study developmental neurotoxicity David Pamies, Université de Lausanne, Lausanne, Switzerland
	Mini-gut organoids for therapeutic testing Nathalie Vergnolle, IRSD, Toulouse, France
15h30–17h30	Session 03 – Symposium Artificial intelligence and machine learning in chemical risk assessment Chairs: João Barroso, Italy and Anne Marie Vinggaard, Denmark
	Systematic reviews and chemical risk assessment: current challenges, and the need for AI in overcoming them Paul Whaley, Lancaster University, Lancaster, UK
	Use of chemical informatics, quantum chemistry modelling and artificial intelligence algorithms to predict molecular initiating events Tim Allen, St. John's College, Cambridge, UK
	Machine learning in silico models in chemical hazard identification Eva Bay Wedeby, Technical University of Denmark, Kgs. Lyngby, Denmark
	Virtual physiological human Geris Liesbet, University of Liège, Belgium



15h30–17h30	Session 03 A – Symposium EAPCCT symposium on COVID-19 and the toxicity of therapeutic drugs and vaccines Chair: Martin Wilks, Switzerland and Paul Dargan, UK Specific treatments in the critically ill COVID-19 patients: benefits and toxicities Bruno Mégarbane, University of Paris, France Understanding vaccine-induced immune thrombocytopenia and thrombosis Beverly Hunt, Guy's and St Thomas' NHS Foundation Trust and King's College London, UK Impact of the COVID crisis on European poison centres Davide Lonati, Poison Control Centre and National Toxicology Information Centre - Toxicology Unit, Pavia, Italy Rapid Development of the FDA ACMT COVID-19 ToxIC (FACT) Pharmacovigilance Pilot Project to Monitor Adverse Events Reported in Association with COVID-19 Therapeutics Paul Wax, American College of Medical Toxicology, Phoenix, AZ, US
17h30-18h00	Virtual Coffee Break, Exhibition & Poster Viewing/Discussion
18h00-20h00	Session 04 – Symposium Personalized nano-immunotoxicology for the workplace Chairs: Martin Himly, Austria and Paola Italiani, Italy Does immunotoxicity of nanomaterials depend on the individual pre-existing conditions? A need for a personalised testing strategy Paola Italiani, National Research Council, Naples, Italy In vitro immuno-nanotoxicological methods that take pre-existing conditions into account Martin Himly, Salzburg University, Salzburg, Austria Understanding the biological impact of industrial engineered nanomaterials upon the alveolar epithelial barrier in vitro using occupational and in vivo relevant concentrations
	Martin Clift, Swansea University, Swansea, UK Impact of nanomaterials on haemodynamic parameters in normal and disease conditions Julie Laloy, Université de Namur, Namur, Belgium



18h00-20h00	Session 05 – Symposium The use of minipigs in juvenile studies in an evolving regulatory landscape Chairs: Andrew Makin, Denmark and Lars Friis Mikkelsen, Denmark Use of juvenile minipigs in testing drugs and foodstuffs to demonstrate safety for human children – practical issues Andrew Makin, Andrew Makin Preclinical Consulting, Kokkedal, Denmark The minipig – a rising star for nonclinical safety testing in support of development of paediatric medicines? Georg Schmitt, Roche Pharma, Basel, Switzerland The juvenile Göttingen Minipig: role of organ development in view of food and drug safety in neonates, infants and toddlers Steven Van Cruchten, University of Antwerp, Antwerp, Belgium Early life nutrition and later life cardiometabolic health in Göttingen Minipigs
18h00-20h00	Sietse Jan Koopmans, Wageningen UR Livestock Research, Wageningen, Netherlands Session 06 – Symposium Human induced pluripotent stem cell (iPSC)-based test systems for future mechanism-based chemical safety testing Chairs: Catherine Verfaillie, Belgium and Marcel Leist, Germany
	iPSC-derived neurospheres for chemical safety assessment Andras Dinnyes, Biotalentum, Gödöllő, Hungary Multicellular 3D liver models based on hiPSC-derived liver cells Catherine Verfaillie, Leuven University, Leuven, Belgium
	Dissecting lineage specific oxidative stress response dynamics using high content imaging of the hiPSC HMOX1 fluorescent reporter line Marije Niemeijer, Leiden Academic Centre for Drug Research, Leiden, Netherlands Nephrotoxic liability assessment using hiPSC-derived renal glomerular and proximal tubular epithelial cells Anja Wilmes, Free University Amsterdam, Amsterdam, Netherlands
18h00-19h00	Industry Symposium I



19h00-20h00

Industry Symposium II

Tuesday, 28 Se	Tuesday, 28 September 2021	
08h00-18h30	Exhibition & Poster Viewing	
08h00-12h00	CEC04 Inflammation as a mediator of toxic responses Chairs: Emanuela Corsini, Italy and Ron Tjalkens, US Inflammation as a mediator of toxic responses	
	Marie Cumberbatch, Immune Insight, Alderley Park, UK The multiple facets of skin inflammation: from direct toxic insult to specific immune responses Marc Pallardy, Université Paris-Saclay, Châtenay-Malabry, France Innate immune inflammatory signaling in glial cells modulates chemical neurotoxicity Ron Tjalkens, Colorado Stage University, Fort Collins, US Evaluating cytokines in immunotoxicity testing	
	Emanuela Corsini, University of Milan, Milan, Italy	
08h00-12h00	CEC05 Nanotoxicology Chair: Ulla Vogel, Denmark Genotoxicity of nanomaterials	
	Julia Catalán Rodríguez, Finnish Institute of Occupational Health, Helsinki, Finland Nanomaterial-induced inflammation, acute phase response and risk of cardiovascular disease Ulla Vogel, National Research Center of the Working Environment, Copenhagen, Denmark Toxicity of nanomaterial in the user phase	
	Anne Saber, National Research Center of the Working Environment, Copenhagen, Denmark	



	In vitro-based high-throughput screening and toxicogenomics to support effective safety evaluation of engineered nanomaterials Penny Nymark, Karolinska Institutet, Institute of Environmental Medicine, Stockholm, Sweden
08h00-12h00	CEC06 Toxicity assessment in drug development Chair: Stine Bartelt, Måløv, Denmark Challenging early target safety assessment strategies
	Phototoxicity of small molecules – from initial assessment to in vivo studies Allan Dahl Rasmussen, Lundbeck A/S, Valby, Denmark Effects of an FGF21 analogue on the female reproductive system Stine Bartelt, Novo Nordisk A/S, Måløv, Denmark PEGylated coagulation factor IX: The road to regulatory approval
12h00-13h00	Hanne Offenberg, Novo Nordisk A/S, Måløv, Denmark Virtual Coffee Break, Exhibition & Poster Viewing/Discussion
13h00-14h00	EUROTOX–SOT Debate Chairs: Félix Carvalho, EUROTOX President-Elect and Michael Aschner, SOT Vice President Individualized toxicity Is the future of risk assessment Alan Boobis (EUROTOX debater) and Syril Pettit (SOT debater)
14h00-16h00	Session 07 – Roundtable Setting the European Environment and Health Research Agenda, 2020-2030: the HERA project Chairs: Robert Barouki, France and Manolis Kogevinas, Spain Identifying research gaps in environment and health research Roel Vermeulen, Utrecht University, Utrecht, Netherlands and Annette Peters, Helmholtz Zentrum, Munich, Germany



	Stakeholder approach for identification of research needs of policy and practice in environment, climate and health Brigit Staatsen, RIVM, Bilthoven, Netherlands
	Major environmental stressors and their effect on health: a global perspective Julia Nowacki, WHO Regional Center, Bonn, Germany
	Infrastructure needs in the field of environment and health Jana Klánová, Recetox, Brno, Czech Republic
14h00–16h00	Session 08 – Symposium Back-translation from clinical outcomes, how did investigative toxicology, modelling and simulation actually perform? Chairs: Harrie C.M. Boonen, Denmark and François Pognan, Switzerland
	Mathematical modelling combined with new in-vitro technologies to enable quantitative translation between preclinical and clinical safety
	Carmen Pin, AstraZeneca, Cambridge, UK
	A case study of a retrospective review of risk factors for DILI based on in vitro and in vivo data Thomas Steger-Hartmann, Bayer AG, Berlin, Germany
	Moving from detection of cardiovascular liabilities to quantitative mechanistic translational understanding: challenges and opportunities
	Amy Pointon, AstraZeneca, Cambridge, UK
	eTRANSAFE's Rosetta stone – a new approach to overcome safety translational hurdles Jan Kors, Erasmus University Medical Center, Rotterdam, Netherlands
14h00-16h00	Session 09 – Workshop Increasing confidence in non-animal approaches for regulatory decision-making Chairs: Suzanne Fitzpatrick, US and Fiona Sewell, UK
	Acceptance of in silico methods for regulatory purposes Glenn Myatt, Leadscope, Columbus, US
	Strengthening a grouping/read-across case using omics-derived molecular mechanistic evidence from an invertebrate model
	Tomasz Sobanski, ECHA, Helsinki, Finland & Mark Viant, University of Birmingham, Birmingham, UK



	A new path for pesticide assessment: using the AOP framework as a tool in risk assessment Susanne Hougaard Bennekou, Technical University of Denmark, Kgs. Lyngby, Denmark
14h00-16h00	Session 28 – Symposium Preclinical immune-safety evaluation of immuno-oncology therapies Chair: Curtis Maier, US
	Current nonclinical evaluation of immune-related safety risks for IO biopharmaceuticals N. N.
	Current nonclinical evaluation of immune-related safety risks for engineered T cell therapies Hervé Lebrec, AMGEN, US
	Clinical toxicology of immune checkpoint blockers Nathalie Chaput-Gras, University Paris-Sud Institut Gustave Roussy, Châtenay-Malabry & Villejuif, France
	Regulatory considerations and establishing FIH dose across immunomodulators Gabriele Reichmann, Paul-Ehrlich-Institut, Langen, Germany
16h00–16h30	Virtual Coffee Break, Exhibition & Poster Viewing/Discussion
16h30-18h30	Session 10 – Symposium Computational modeling of AOP networks to assist risk assessment of chemicals Chairs: Frederic Bois, UK and Joost Beltman, Netherlands
	Quantitative Bayesian networks analyses of mitochondrial toxicity Frederic Bois, CERTARA Inc., Sheffield, UK
	Logic modeling of toxicology pathways Attila Gabor, EMBL, Heidelberg, Germany
	Data-driven computational modeling of the DNA damage response and liver toxicity Joost Beltman, Leiden University, Leiden, Netherlands



	Spatial-temporal multiscale-multilevel modeling of APAP damage and its consequence on ammonia detoxification in a virtual liver lobule: steps towards a virtual liver Dirk Drasdo, INRIA & University of Leipzig, Paris & Leipzig, France & Germany
16h30-18h30	Session 11 – Symposium Emerging tools for the investigation and prediction of liver toxicity Chairs: Mathieu Vinken, Belgium and Magnus Ingelman-Sundberg, Sweden
	ULA 3D spheroids as a tool for studying normal and diseased liver function and for prediction of drug pharmacokinetics and hepatotoxicity Magnus Ingelman-Sundberg, Karolinska Institutet, Stockholm, Sweden
	Functional imaging of hepatotoxicity Jan Hengstler, Leibniz Research Center (IfADo), Dortmund, Germany
	Dynamic imaging of stress response pathway activation for quantitative systems liver toxicity approaches Bob van de Water, Leiden University, Leiden, Netherlands
	Using real time sensors to illuminate human-relevant mechanisms of action Yaakov Nahmias, Silberman Institute of Life Sciences, Jerusalem, Israel
16h30-18h30	Session 12 – Symposium Application of high throughput transcriptomics in mechanism-based chemical safety assessment Chair: Hennicke Kamp, Germany
	High throughput transcriptomics for determining chemical-induced perturbations to predict adverse renal outcomes Paul Jennings, Free University Amsterdam, Netherlands
	Early prediction of late adverse outcome using benchmark dose modelling of high throughput transcriptomics data Scott Auerbach, U.S. NIEHS/National Toxicology Program, Durham, US
	Transcriptomic profiling of the inter-individual variability of chemical-induced cellular stress response activation in primary human hepatocytes Marije Niemeijer, Leiden University, Leiden, Netherlands



	Genomics-based platforms in combination with machine learning algorithms enabling well informed and reliable risk assessments for different toxicological endpoints Andy Forreryd, SenzaGen, Lund, Sweden
16h30-17h30	Industry Symposium III
17h30-18h30	Industry Symposium IV
18h30-20h00	Short Oral Communications I Chairs: tba. SOC01-01 Development of a kidney-on-a-chip model that replicates an antisense oligonucleotide-induced kidney injury biomarker response T. T. Nieskens¹, O. Magnusson¹, M. Persson¹, P. Andersson², M. Söderberg¹, A. Sjögren¹ ¹AstraZeneca, CVRM Safety, Clinical Pharmacology and Safety Sciences, R&D, Gothenburg, Sweden ²AstraZeneca, R&I Safety, Clinical Pharmacology and Safety Sciences, R&D, Gothenburg, Sweden SOC01-02
	Pyrrolizidine alkaloids affect miRNA expression in human HepaRG cells A. M. Enge, H. Sprenger, A. Braeuning, S. Hessel-Pras German Federal Institute for Risk Assessment, Berlin, Berlin, Germany SOC01-03 Novel in vitro model captures drug-induced inflammation as a mechanism contributing to hepatotoxicity F. Tasnim¹, X. Huang¹,², Y. T. Soong¹, H. Yu¹,²,²,³ ¹ Institute of Bioengineering and Nanotechnology, Health and Medical Technologies, Singapore, Singapore ² National University of Singapore, Department of Physiology, Yong Loo Lin School of Medicine, Singapore ³ CAMP IRG, Singapore-MIT Alliance for Research and Technology, Singapore, Singapore SOC01-04 Scientific validity of non-animal-derived antibodies J. Barroso, M. Halder, M. Whelan European Commission, Joint Research Centre, Ispra (VA), Italy SOC01-05
	In vitro toxicity screening of an inclusive panel of engineered nanomaterials using an advanced 3D liver model S. V. Llewellyn ¹ , G. E. Conway ¹ , UK. Shah ¹ , G. J. Jenkins ¹ , M. J. Clift ¹ , S. H. Doak ¹



Swansea University, In Vitro Toxicology Group, Swansea, UK

SOC01-06

Development of a Liver Carcinoma Biomarker Panel in 3D HepG2 Liver Spheroids Following Exposure to Ag and Tio₂ Nanomaterials

<u>G. E. Conway</u>¹, S. V. Llewellyn¹, P. Nymark^{2, 3}, U. B. Vogel⁴, S. Halappanavar⁵, G. J. Jenkins¹, M. J. Clift¹, S. H. Doak¹

¹InVitro Toxicology Group, Swansea University Medical School, Swansea, United Kingdom

²Institute of Environmental Medicine, Karolinska Institute, Stockholm, Sweden

³Misvik Biology Oy, Turku, Finland

⁴National Research Centre for the Working Environment, Copenhagen, Denmark

⁵Environmental Health Science and Research Bureau, Health Canada, Ottawa, Canada

SOC01-07

Combining single cell gene expression analysis and a 3D *in vitro* liver model to investigate cell type-specific responses to profibrotic TGF-B1

C. J. Messner^{1, 2}, L. Babrak¹, G. Titolo¹, M. Caj¹, E. Miho^{1, 3, 4}, L. Suter-Dick^{1, 2}

¹Fachhochschule Nordwestschweiz, Institute for Chemistry and Bioanalytics, Muttenz, Basel-Land, Switzerland

²Swiss Centre for Applied Human Toxicology, Basel, Basel-Stadt, Switzerland

³SIB Swiss Institute of Bioinformatics, Lausanne, Switzerland

⁴aiNET GmbH, Basel, Switzerland

18h30-20h00

Short Oral Communications II

Chairs: tba.

SOC02-01

Towards a personalized drug development using autologous iPSC-derived Multi-Organ-Chips

<u>A. Ramme</u>, L. Koenig, D. Faust, M. Jäschke, N. Nguyen, E.-M. Dehne, U. Marx

TissUse GmbH, Berlin, Berlin, Germany

SOC02-02

Prediction of inotropic effect based on calcium transients in human iPSC-derived cardiomyocytes using novel waveform parameters and a modified random forest algorithm

<u>H. Yang</u>¹, O. Obrezanova², A. Pointon², W. Stebbeds³, J. Francis³, K. A. Beattie⁴, P. Clements⁴, J. S. Harvey⁴, G. F. Smith², A. Bender¹

¹University of Cambridge, Department of Chemistry, Cambridge, United Kingdom

²AstraZeneca, BioPharmaceuticals R&D, Cambridge, United Kingdom

³GlaxoSmithKline, R&D, Stevenage, UK

4GlaxoSmithKline, R&D, Ware, UK



SOC02-03

High-throughput phenotypic profiling within the NAMs-based, tiered hazard evaluation strategy at the United States Environmental Protection Agency

J. Nyffeler^{1, 2}, C. Willis¹, M. Culbreth¹, R. E. Brockway^{1, 3}, L. J. Everett¹, G. Patlewicz¹, I. Shah¹, D. Chang¹, K. Paul Friedman¹, J. Wambaugh¹, J. A. Harrill¹

¹US Environmental Protection Agency, Center for Computational Toxicology and Exposure, Office of Research and Development, Durham, North Carolina, United States of America

²Oak Ridge Institute for Science and Education (ORISE) Postdoctoral Fellow, Oak Ridge, Tennessee, US

³Oak Ridge Associated Universities (ORAU) National Student Services Contractor, Oak Ridge, Tennessee, US

SOC02-04

Mutations in the filaggrin gene determine immune response after dermal chemical exposure

E. Rietz Liljedahl², H. K. De Paoula², M. Engfeldt², A. Julander¹, C. Lidén¹, C. Lindh², **K. Broberg**^{1, 2}

¹Karolinska Institutet, Institute of Environmental Medicine, Stockholm, Sweden

²Lund University, Department of Laboratory Medicine, Lund, Sweden

SOC02-05

The impact of pooling animal histopathology control data on the statistical detection of treatment-related findings

P. S. R. Wright¹, K. A. Briggs², R. Thomas², G. F. Smith³, G. Maglennon⁴, P. Mikulskis⁵, M. Chapman⁶, N. Greene⁷, A. Bender¹ University of Cambridge, Chemistry, Cambridge, UK

²Lhasa Limited, Leeds, UK

³AstraZeneca, Data Science and AI, Clinical Pharmacology and Safety Sciences, R&D, Cambridge, UK

⁴AstraZeneca, Oncology Pathology, Clinical Pharmacology and Safety Sciences, R&D, Melbourn, UK

⁵AstraZeneca, Imaging and Data Analytics, Clinical Pharmacology & Safety Sciences, R&D, Gothenburg, Sweden

⁶AstraZeneca, Toxicology, Clinical Pharmacology and Safety Sciences, R&D, Melbourn, UK

⁷AstraZeneca, Imaging and Data Analytics, Clinical Pharmacology & Safety Sciences, R&D, Waltham, Massachusetts, US

SOC02-06

Novel biomimetic membranes for nanoparticle transport studies at biological barriers

<u>L. A. Furer</u>¹, A. Díaz Abad¹, G. Fortunato², S. Schürle-Finke³, T. Buerki-Thurnherr¹

¹ Empa, Laboratory for Particles-Biology Interactions, St. Gallen, Switzerland

² Empa, Laboratory for Biomimetic Membranes and Textiles, St.Gallen, Switzerland

³ ETH Zürich, Responsive Biomedical Systems Lab, Zürich, Switzerland

SOC02-07

Validation of the ToxProfiler reporter assay for toxicological profiling and determination of the underlying mode of action **B. ter Braak**¹, L. Wolters¹, T. Osterlund¹, B. Van de Water², G. Hendriks¹



¹Toxys b.v., Leiden, Netherlands ²Leiden University, Leiden Academic Center for Drug Research, Leiden, Netherlands



Wednesday, 2	Wednesday, 29 September 2021	
09h00-18h30	Exhibition & Poster Viewing	
10h00-11h00	Specialty Sections meetings Individual Members meeting	
12h00-13h00	Virtual Coffee Break, Exhibition & Poster Viewing/Discussion	
13h00-14h00	SOT Merit Award Lecture 1 Chairs: George Daston, SOT Past President and Heather Wallace, EUROTOX President	
	Unraveling the molecular mechanisms of cannabinoid-mediated immune modulation and cannabinoid receptor 2 as a putative therapeutic target Norbert E. Kaminski, Michigan State University, East Lansing, MI, US	
14h00-16h00	Session 13 – Workshop Modes of action in non-genotoxic carcinogenesis Chairs: Jan Vondracek, Czech Republic and William H Bisson, US	
	Developing an integrated approach for the testing and assessment of chemical non-genotoxic carcinogens for global regulatory purposes Miriam Jacobs, Public Health England, Chilton, UK	
	Cellular and newly proposed models to study the transforming ability of pollutants for translational toxicology William H Bisson & Annamaria Colacci, OHSU Knight Cancer Institute & ARPAE, Portland & Bologna, US & Italy	
	Non-coding RNAs mechanisms enforcing oncogenic programs and allowing establishment of metastatic inches Martin Bushell, The Beatson Institute, Glasgow, UK	
14h00-16h00	Session 14 – Symposium New approaches using in vitro assays and 3D models can improve prediction of immune reactions to xenobiotics Chairs: Marc Pallardy, France and Saadia Kerdine-Römer, France	
	Immune response to chemicals and drugs: understanding is key for prediction Marc Pallardy, University Paris-Sud, Châtenay-Malabry, France	



	Development of new approaches to predict drug-induced hypersensitivity with an increased understanding of reaction mechanisms Dean Naisbitt, University of Liverpool, Liverpool, UK The challenges of predicting biological products immunogenicity using T-cell assays Bernard Maillère, University of Paris Saclay, Paris, France Challenges and opportunities of 3D-skin models: the way forward for assessing chemical sensitizers? Sue Gibbs, Amsterdam University Medical Center, Amsterdam, The Netherlands
14h00-16h00	Session 15 – Symposium Impact of climate change on food safety Chairs: Angela Mally, Germany and George Kass, Italy The impact of climate change on mycotoxin and related fungi risks in Europe: current scenario and future perspectives Antonio Moretti, Institute of Sciences of Food Production, National Research Council, Bari, Italy Climate change impacts on harmful algal bloom toxicity Dedmer van de Waal, Netherlands Institute of Ecology (NIOO-KNAW), Wageningen, Netherlands Ocean warming and Ciguatera poisoning Elisa Berdalet, Institute of Marine Sciences (ICM-CSIC), Barcelona, Spain Tetrodotoxins in seafood from European waters Ron Hoogenboom, RIKILT Wageningen University & Research, Wageningen, Netherlands
14h00-16h00	Session 29 – Symposium Is there a human risk to PFAS exposure? Chairs: Philippe Grandjean, Denmark and Martin Wilks, Switzerland Investigating developmental effects of PFAS using a 3D human induced pluripotent stem cell differentiation model Anne Marie Vinggaard, Technical University of Denmark, Kgs. Lyngby, Denmark Epidemiological approaches to PFAS toxicity



	Philippe Grandjean, University of Southern Denmark & Boston University, Odense & Boston, Denmark & US
	Exposure to real-life PFAS mixtures present in several human matrices, assessed with ex vivo effect biomarkers Eva Cecilie Bonefeld-Jørgensen ^{1,2} , Vicente Mustieles ^{3,4,5,6} , Andrea Rodríguez ^{3,4,5,6} , Maria Wielsøe ¹ , Mariana F. Fernandez ^{3,4,5,6} ¹ Centre for Arctic Health & Molecular Epidemiology, Department of Public Health Aarhus University, Denmark ² Greenland Centre for Health Research, University of Greenland, Nuuk, Greenland ³ University of Granada, Center for Biomedical Research (CIBM), Spain ⁴ Department of Radiology and Physical Medicine, School of Medicine, University of Granada, Granada, Spain ⁵ Instituto de Investigación Biosanitaria Ibs Granada, Spain; ⁶ Consortium for Biomedical Research in Epidemiology & Public Health (CIBERESP), Spain Wide-spread PFAS contamination of drinking water in Sweden – exposure and health risk assessment Anders Glynn, SLU, Uppsala, Sweden
16h00-16h30	Virtual Coffee Break, Exhibition & Poster Viewing/Discussion
16h30–18h30	Session 16 – Symposium Human microengineered organs-on-chips: advancing regulatory science through innovation Chairs: Suzanne Fitzpatrick, US and Adrian Roth, Switzerland Organs-on-chips for safety testing and disease 18odelling Geraldine A. Hamilton, Emulate Inc., Boston, US
	Human on a chip – are we there yet? Uwe Marx, TissUse GmbH, Berlin, Germany
	Incorporating organ-on-chip into integrated approaches to testing and assessment Sofia Batista Leite, European Commission's Joint Research Centre, Ispra, Italy
	An industry perspective: importance of organs-on-chips for advancing drug discovery and development Adrian Roth, Roche Pharma, Basel, Switzerland
16h30–18h30	Session 17 – Symposium Designing toxicology studies to support development of cell-based therapies Chair: Niklas Öhrner, Denmark, Ridhirama Bhuwania, Germany



	Regulatory considerations for cell based therapies David Jones, MHRA, UK Non-clinical study design considerations in the development of cellular therapeutics Mark Johnson, Northern Biomedical Research, Norton Shores, MI, US
	Preclinical assessment of a pluripotent cell therapy for Parkinson's Disease Agnete Kirkeby, Lund University, Lund, Sweden
	How cell therapies derived from pluripotent stem cells are evaluated for tumorigenicity in vivo and in vitro Dorthe Bach Toft, NovoNordisk, Måløv, Denmark
16h30–18h30	Session 18 – Symposium Mechanistic toxicology as the basis for modelling and prediction of organ-specific toxicity Chairs: Anna Bal-Price, Italy and Ulla Vogel, Denmark
	Applying the adverse outcome pathways network for understanding and predicting neurotoxicity Anna Bal-Price, European Commission Joint Research Centre, Ispra, Italy
	Application of the adverse outcome pathway conceptual framework for translation of mechanistic data into regulatory decisions: adverse outcome pathways for kidney injury as case study Angela Mally, University of Würzburg, Würzburg, Germany
	Novel means of enabling high-throughput toxicogenomics and adverse outcome pathways for prediction of lung toxicity Penny Nymark, Karolinska Institute, Stockholm, Sweden
	Development and application of an adverse outcome pathway of cholestatic liver injury Mathieu Vinken, Vrije Universiteit Brussels, Brussels, Belgium
16h30–17h30	Industry Symposium V
17h30-18h30	Industry Symposium VI
18h30-20h00	Short Oral Communications III Chairs: tba.



SOC03-01

Rat biodistribution of cerium dioxide and titanium dioxide nanomaterials after single and repeated inhalation exposure

<u>I. Gosens</u>¹, E. Duistermaat¹, J. Boere¹, P. Fokkens¹, J. Vidmar², K. Löschner², C. Delmaar¹, A. L. Costa³, R. Peters⁴, F. Cassee^{1, 5}

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²Technical University of Denmark, National Food Institute, Kongens Lyngby, Denmark

³Institute of Science and Technology for Ceramics, National Research Council, Faenza, Italy

⁴Wageningen Food Safety Research, Wageningen, Netherlands

⁵Institute for Risk Assessment Studies,, Utrecht, Netherlands

SOC03-02

Estimation of uncertainty in multi-level in silico models predicting biomarkers of drug-induced proarrhythmic risk

K. Kopanska¹, J. C. Gómez-Tamayo^{1, 2}, J. Llopis-Lorente³, B. A. Trenor-Gomis³, J. Sáiz³, M. Pastor¹

¹ Universitat Pompeu Fabra, Research Programme on Biomedical Informatics (GRIB), Department of Experimental and Health Sciences, Barcelona, Spain

² Janssen, Research & Development, Beerse, Belgium

³ Universitat Politècnica de València, Centro de Investigación e Innovación en Bioingeniería, Valencia, Spain

SOC03-03

Next generation risk assessment for skin sensitization combining non-animal data and read- across: a case study with Resorcinol

<u>F. Gautier</u>¹, F. Tourneix², H. Assaf Vandecasteele¹, D. Bury¹, N. Alépée²

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SOC03-04

Expanding the H295R steroidogenic assay using LC-MS/MS and an ER-alpha reporter gene assay as read-outs using azole fungicides as test compounds

<u>P. Vazakidou</u>, C. Koopmans, S. Grimberg, S. Evangelista, J. Koekkoek, M. Lamoree, P. Leonards, M. van Duursen Vrije University of Amsterdam, Department Environment and Health, Amsterdam, Netherlands

SOC03-05

A novel prediction model to evaluate genotoxicity based on a gene signature in metabolically competent human HepaRGTM cells

A. Thienpont¹, S. Verhulst², L. van Grunsven², V. Rogiers¹, T. Vanhaecke*¹, B. Mertens*^{3, 4}

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SOC03-06

N. N.

SOC03-07

Air pollutants induced immunotoxicity linked to COVID-19 complications: toxicogenomic approach

<u>D. Jorgovanovic</u>, K. Živančević, K. Baralić, A. Buha Djordjevic, E. Antonijević Miljaković, B. Antonijević, D. Đukić-Ćosić University of Belgrade - Faculty of Pharmacy, Department of Toxicology, Belgrade, Serbia

18h30-20h00

Short Oral Communications IV

Chairs: tba.

SOC04-01

Estimating the kinetics of titanium dioxide nanoparticles in rats after inhalation using physiologically based kinetic modelling

J. Minnema¹, C. Delmaar¹, I. Gosens¹, F. R. Cassee^{1, 2}, L. Tran³, B. Bokkers¹

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³Institute of Occupational Medicine, Edinburgh, UK

SOC04-02

Glyphosate and T cells: an immunotoxicity in vitro evaluation

A. Maddalon¹, V. Galbiati¹, M. Iulini², M. Marinovich¹, E. Corsini²

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SOC04-03

Analysis of Murine Liver mRNA Expression, DNA Methylation, And Histone After Repeated Exposure To Chemicals

J. Kanno, K.-I. Aisaki, R. Ono, S. Kitajima

National Institute of Health Sciences, Division of Cellular & Molecular Toxicology/Center for Biological Safety & Research, Kawasaki, Japan

SOC04-04

N.N.



SOC04-05

An *in vitro* harmonized strategy to assess the toxicity of chemicals using multiple human induced pluripotent stem cell (hiPSC)-derived models

<u>C. Nunes</u>^{1, 2}, P. Singh^{3, 4}, Z. Mazidi^{5, 6}, C. Murphy⁷, A. Bourguignon^{8, 9}, S. Wellens¹⁰, S. Ghosh¹¹, M. Zana⁸, C. Verfaillie¹¹, M. Culot¹⁰, A. Dinnyés^{8, 13}, P. Jennings⁷, J. Grillari^{5, 6, 12}, T. Exner¹⁴, M.-G. Zurich^{1, 2}

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Pharmaceutical Sciences, Amsterdam, Netherlands

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10 University of Artois, Laboratoire de la Barrière Hémato-Encéphalique (LBHE), Faculté des sciences Jean Perrin, Lens, France

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SOC04-06

Brain organoids to study SARS-Cov-2 infection of developing CNS

L. Smirnova¹, C. K. Bullen², H. T. Hogberg¹, A. Pekosz³, W. Bishai², T. Hartung¹

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SOC04-07

In silico approaches to link adverse outcomes to molecular initiating events through AOPs

<u>A. A. Oliveira</u>, A. Caley, S. A. Stalford, S. Kane, R. Foster, E. Hill, G. Kocks, A. Fowkes, A. Myden, D. Newman, J. D. Vessey Lhasa Limited, Leeds, UK



Thursday, 30 September 2021	
09h00-18h30	Exhibition & Poster Viewing
09h00-12h00	Business Council Meeting
12h00-13h00	Virtual Coffee Break, Exhibition & Poster Viewing/Discussion
13h00-14h00	SOT Merit Award Lecture 2 Chairs: George Daston, SOT Past President and Heather Wallace, EUROTOX President
	The exciting challenge of working in regulatory toxicology Rogene Henderson, Lovelace Respiratory Research Institute, Albuquerque, NM
14h00-16h00	Session 19 – Workshop Can we panelize seizure? Chairs: Ruth Roberts, UK and Jennifer Pierson, US Seizure liability in drug discovery and development: current state of play Jean-Pierre Valentin, UCB, Braine-I'Alleud, Belgium Exploring the utility of an ion channel panel for detecting seizure liability Mike Morton, ApconiX, Alderley Park, UK Development of seizure prediction methods using in vitro microelectrode array (MEA) Ikuro Suzuki, Tohoku Institute of Technology, Sendai, Japan Panel Discussion focusing on key questions: confidence in the biology, confidence in the robustness of the assay and confidence in translation to the clinic and the patient All speakers
14h00-16h00	Session 20 – Symposium Modernizing cancer risk assessment: beyond the bioassay Chairs: Gina Hilton, UK and Mirjam Luijten, Netherlands
	Current challenges in a paradigm shift for cancer risk assessment



Alan Boobis, Imperial College London, UK Current transcriptional benchmark dose approaches that are available to safely assess carcinogenicity risk Virunya Bhat, Independent Consultant, US Strategies for a weight of evidence-based carcinogenicity assessment of human pharmaceuticals Jan Willem van der Laan, Netherlands Organization for Applied Scientific Research (TNO), Zeist, Netherlands Predicting non-genotoxic carcinogenic potential of agrochemicals: a mechanistic approach Mirjam Luijten, RIVM, Netherlands 14h00-16h00 Session 21 - Symposium Drug - exposome interactions Chairs: Benedikt Warth, Austria and Angela Mally, Germany The exposome: drugs, toxicants, and metabolites Gary W. Miller, Columbia University, New York City, US Impact of dietary xenoestrogens and other food contaminants on drug metabolism and action Benedikt Warth, University of Vienna, Vienna, Austria The Central European Longitudinal Studies of Parents and Children (CELSPAC) from an exposome perspective Jana Klánová, Masaryk University, Brno, Czech Republic Biotransformation-driven interactions and precision responses to chemotherapy Shana Sturla, ETH Zurich, Zurich, Switzerland 14h00-16h00 Session 30 - Symposium Revisiting paracetamol-induced multisystem toxicity: novel mechanistic insights Chairs: Hilmi Orhan, Turkey and Hartmut Jaeschke, US Paracetamol hepatotoxicity: Discovering new drugs based on mechanistic insight from animal studies Hartmut Jaeschke, University of Kansas Medical Center, Kansas City, US Paracetamol-associated adverse reactions in kidney: different mechanistic pathways compared to liver Hilmi Orhan, Ege University, Izmir, Turkey



	Paracetamol and pregnancy: short- and long-term consequences for mother and child N. N. Paracetamol and development – reasons for concern David Kristensen, University of Copenhagen & Inserm, Irset, Copenhagen & Rennes, Denmark & France
16h00–16h30	Virtual Coffee Break, Exhibition & Poster Viewing/Discussion
16h30-18h30	Session 22 – Symposium Computational models to reliably predict chemical mixture toxicity Chairs: Aristidis Tsatsakis, Greece and Michael Aschner, US SMF-1, SMF-2 and SMF-3DMT1 orthologues regulate and are regulated differentially by manganese levels in C. elegans Michael Aschner, Albert Einstein College of Medicine, New York, US Systems toxicology models for the development of AOP networks induced by exposure to complex mixtures Denis Sarigiannis, University School of Advanced Studies IUSS, Pavia, Italy Computational modelling: A new paradigm for chemical mixtures risk assessment? Antonio F. Hernandez, University of Granada School of Medicine, Granada, Spain PBPK modelling: Bridging animal-free toxicology tools and conventional in vivo testing for cumulative risk assessment after long-term-low-dose exposure to chemical mixtures Marina Goumenou, University of Crete Medical School, Heraklion, Greece
16h30-18h30	Session 23 – Symposium The value of micro-physiological systems for drug safety assessment – a series of case studies Chairs: Ekaterina Breous-Nystrom, Switzerland and Thomas Steger-Hartmann, Germany Microphysiological systems evolution – introducing the immune compartment Ekaterina Breous-Nystrom, Roche, Basel, Switzerland A microfluidic two-organ chip to investigate species specific differences of thyroid-liver crosstalk in human and rats Diana Karwelat, Bayer AG, Berlin



	Adopting organ-chips as internal decision making-tools: a quantitative evaluation of human liver-chip predictive validity Lorna Ewart, Emulate, London, UK Human iPSC-derived retinal organoid model for in vitro toxicity screening Valeria Chichagova, Newcells Biotech, Newcastle, UK
16h30-18h30	Session 24 – Symposium Building confidence in the use of new approach methodologies for safety decision-making Chairs: Alistair Middleton, UK and Ans Punt, Netherlands
	Strategies to evaluate <i>in vitro in silico</i> physiologically based kinetic (PBK) models as essential tool in next generation (animal-free) risk evaluations Ans Punt, RIKILT Wageningen University and Research, Wageningen, Netherlands
	In silico approaches to link adverse outcomes to molecular initiating events through AOPs Oliveira Anax, Lhasa Limited, Leeds, UK
	Strategic use of high-throughput transcriptomics and phenotypic profiling data in support of regulatory decisions Joshua Harrill, US EPA NCCT, Research Triangle Park, US
	An industry perspective on strategies for integrating new approach methodologies for next generation risk assessment Maria Baltazar, Unilever Safety and Environmental Assurance Centre, Bedford, UK
16h30-17h30	Industry Symposium VII
17h30–18h30	Industry Symposium VIII



Friday, 1 October 2021	
09h00-17h00	Exhibition & Poster Viewing
10h00-12h00	EC21-3 Meeting
12h00-13h00	Virtual Coffee Break, Exhibition & Poster Viewing/Discussion
13h00-14h00	HESI CITE Lecture
14h00-16h00	Session 25 – Symposium Safeguarding female reproductive health across disciplines Chairs: Julie Boberg, Denmark and Paul Fowler, UK New insights into how early-life exposure to industrial chemicals can disrupt female reproductive development Hanna KL Johansson, Technical University of Denmark, Kgs. Lyngby, Denmark Reproductive toxicity in wildlife N. N. Influence of EDCs on female puberty – evidence from human epidemiology Anders Juul, Copenhagen University Hospital, Copenhagen, Denmark EDCs and female fertility – what can we learn from human clinical samples? Richelle Duque Björvang, Karolinska University Hospital, Stockholm, Sweden
14h00-16h00	Session 26 – Symposium Computational toxicology – New advances and acceptance in academia, industry and regulation Chairs: Timothy Allen, UK and Ruth Roberts, UK Developing and assessing in silico profilers for organ-level toxicity using non-standard data Mark Cronin, Liverpool John Moores University, Liverpool, UK Artificial Intelligence in drug discovery and computational safety: What is realistic, what are illusions? Andreas Bender, University of Cambridge, Cambridge, UK



	Industrial perspectives on in silico tools – early screening to regulatory applications Catrin Hasselgren, Genentech, San Francisco, US Open source computational toxicology tools in food and feed safety: integrating historical data, meta-analysis and species-specific generic models Jean-Lou Dorne, EFSA, Parma, Italy
14h00–16h00	Session 27 – Symposium Predictive systems to identify etiological factors and pathogenic mechanisms of neurodegeneration Chairs: Jonathan Doorn, US and Jason Cannon, US Translation of mechanistic data into in vivo systems to predict risk for neurodegeneration
	Altered neurotransmitter homeostasis as a mechanistic biomarker of neurotoxicity progressing to neurodegeneration Jonathan Doorn, University of Iowa, Iowa City, US Application of an adverse outcome pathway-based in vitro testing battery for neurotoxicity evaluation Katharina Koch, IUF-Leibniz Research Institute for Environmental Medicine, Düsseldorf, Germany
	In vitro neurotoxicity test methods: from development to degeneration Remco Westerink, Utrecht University, Utrecht, Netherlands
14h00-16h00	Industry Symposium IX
14h00-16h00	Industry Symposium X
16h00-17h00	Closing Ceremony and Awards presentation