



Detailed Programme

Status: 10 September, 2015

Sunday, 13 September, 2015		
08h00 - 18h00	Congress Registration	16h00 – 21h00 Exhibition
09h00 - 16h00	Continuing Education Courses (CEC) <i>including individual coffee & lunch breaks</i>	
09h00 - 15h30 D. Luis Hall	CEC 1: Removing Obstacles on the Way to Implement 3R Methods in Toxicology Chairs: Herman Koëter, Belgium and Marc Teunis, The Netherlands Intro: General overview of the variables that determine the process of the regulatory acceptance and use of 3R models (Historical view & new vision) Herman Koëter, Belgium and Marc Teunis, The Netherlands 09:00 – 09:30 CEC1-1 Removing Obstacles on the Way to implement 3R Methods in Hazard and Risk Assessment Herman Koëter <i>Orange House Partnership, S.Lorenzo di Moriano, Lucca, Italy</i> 09:30 – 10:00 CEC1-2 Lessons learned from the 'SLIM' project; regulatory acceptance and use of 3R methods *Cyrille Krul ^{1,2} , Marie-Jeanne Schiffelers ³ , Marc Teunis ² , Raymond Pieters ^{2,3,4} ¹ TNO, Healthy Living, Zeist, Netherlands ² University of Applied Sciences Utrecht, Life Sciences & Chemistry, Utrecht, Netherlands ³ Utrecht University, School of Governance, Utrecht, Netherlands ⁴ Utrecht University, IRAS, Utrecht, Netherlands 10:00 – 10:15 Coffee Break 10:15 – 10:45 Industrial point of view CEC1-3 The importance of the undescribed for industrial and regulatory application of animal-free methods for safety assessment Erwin L. Roggen <i>3Rs Management and Consulting ApS, Lyngby, Denmark</i>	



	<p>10:45 – 11:15</p> <p>Regulators point of view</p> <p>CEC1-4 Removing obstacles on the way to implement 3R methods in toxicology: Regulators point of view Sonja Beken <i>Federal Agency for Medicines and Health Products (FAMHP), DG PRE, Dept. Evaluators, Brussels, Belgium</i></p> <p>11:15 – 11:45</p> <p>Academia's point of view</p> <p>CEC1-5 Implementing 3Rs methods in toxicology: a view from academia Ian Kimber <i>University of Manchester, Faculty of Life Sciences, Manchester, United Kingdom</i></p> <p>11:45 – 12:00 Wrap-up before lunch Herman Koëter, Belgium and Marc Teunis, The Netherlands</p> <p>12:00 – 13:00 Lunch Break</p> <p>13:00 – 15:00</p> <p>CEC1-6 Validation; truth or dare! Round 1: Prioritizing factors that can be influenced by stakeholders Round 2: Actions to be taken to improve implementation and acceptance *Marc Teunis¹, Raymond Pieters^{1,2}, Cyrille Krul^{1,3} ¹<i>University of Applied Sciences, Innovative Testing in Life Sciences & Chemistry, Utrecht, Netherlands</i> ²<i>Utrecht University, Institute for Risk Assessment Sciences, Utrecht, Netherlands</i> ³<i>TNO, Healthy Living, Zeist, Netherlands</i></p> <p>15:00 – 15:30 Wrap-up and conclusions Herman Koëter, Belgium and Marc Teunis, The Netherlands</p>
<p>09h30 – 15:30</p> <p>S. João Hall</p>	<p>CEC 2: Revisiting the Challenges Posed by New Recreational Drugs Chairs: <i>Simon Gibbons, United Kingdom and Bruno Mégarbane, France</i></p> <p>09:30 – 10:00</p> <p>CEC2-1 Novel psychoactive substances: a chemical overview for the toxicologist Simon Gibbons <i>UCL School of Pharmacy, London, United Kingdom</i></p> <p>10:00 – 10:15 General Discussion</p>



10:15 – 10:45

CEC2-2

Pharmacological characterization of designer cathinones in vitro

Matthias Liechti

University Hospital Basel, Basel, Switzerland

10:45 – 11:00

General Discussion

11:00 – 11:15

Coffee Break

11:15 – 11:45

CEC2-3

Understanding mechanisms of toxicity of designer cathinones: contribution of animal models

Bruno Mégarbane

Paris-Diderot University, Lariboisière Hospital, Paris, France

11:45 - 12:00

General Discussion

12:00 – 12:30

CEC2-4:

Toxicovigilance of new psychoactive substances – Perspectives from the EU Early Warning System

Michael Evans-Brown

EMCDDA, Lisbon, Portugal

12:30 – 12:45

General Discussion

12:45 – 13:45

Lunch Break

13:45 – 14:15

CEC2-5

New psychoactive substances: data from the STRIDA project

*Matilda Bäckberg¹, Olof Beck^{2,3}, Anders Helander^{2,3}

¹*The Swedish Poisons Information Centre, Stockholm, Sweden*

²*Karolinska Institutet, Department of Laboratory Medicine, Stockholm, Sweden*

³*Karolinska University Laboratory, Stockholm, Sweden*

14:15 – 14:30

General Discussion

14:30 – 15:00

CEC2-6

The new recreational drugs in the emergency department

David M. Wood

Guy's and St Thomas' NHS Foundation Trust, London, United Kingdom

15:00 – 15:15 General Discussion

15:15 – 15:30 Conclusions



09h30 - 16h00

Arrabida Hall

CEC 3: Thresholds of Toxicological Concern – Basics and Latest Developments *Organised and supported by the International Life Sciences Institute (ILSI Europe)*

Chairs: Susan Barlow, United Kingdom and Kirstin Kosemund, Germany

09:30 – 09:35

Introduction to the CEC on TTC

Susan Barlow, United Kingdom

09:35 - 10:05

CEC3-1

Introduction into the Tiered TTC Concept and regulatory status globally

*Kirstin Kosemund¹, Susan Felter²

¹Procter & Gamble, Global Product Stewardship, Schwalbach am Taunus, Germany

²Procter & Gamble, Central Product Safety, Mason, OH, United States

10:05 – 10:20

Questions and answers

10:20 – 10:50

CEC3-2

Cancer thresholds, Cohort of Concern and other excluded substance groups

Alan Boobis

Imperial College, London, United Kingdom

10:50 – 11:05

Questions and answers

11:05 – 11:30

Coffee Break

11:30 – 12:00

CEC3-3

Non-cancer oral toxicity databases for TTC

Susan Barlow

Consultant, Brighton, United Kingdom

12:00 – 12:15

Questions and answers

12:15 – 12:45

CEC3-5

Estimation of Toxic Hazard - A Revised Cramer-Ford-Hall Decision Tree

*Jürgen Schnabel¹, Sean Taylor²

¹Givaudan International AG, Product Safety and Regulatory Affairs, Kempththal, Switzerland

²International Organisation of the Flavor Industry, Brussels, United States

12:45 – 13:00

Questions and answers

13:00 – 14:00

Lunch Break



	<p>14:00 – 14:30 CEC3-4 Computational modelling for TTC assessment *Andrew Worth¹, Chihae Yang² ¹European Commission, Joint Research Centre, Ispra, Italy ²Altamira LLC, Ohio, United States</p> <p>14:30 – 14:45 Questions and answers</p> <p>14:45 – 15:15 CEC3-6 TTC concept: non oral routes and influence of local effects Sylvia Escher Fraunhofer ITEM, Chemical risk assessment, Hannover, Germany</p> <p>15:15 – 15:30 Questions and answers</p> <p>15:30 – 16:00 Conclusions</p>
09h30 - 16h00 Porto Hall	<p>CEC 4: Evaluating and Expressing Uncertainty in Hazard Characterization: New Guidance from the World Health Organization <i>Organised and supported by the World Health Organization (WHO)</i></p> <p>Chairs: Carolyn Vickers, Switzerland and Matthias Herzler, Germany</p> <p>09:30 – 09:45 Welcome and introductions Carolyn Vickers, World Health Organization, Switzerland</p> <p>09:45 – 10:30 CEC4-1 Probabilistic Hazard Characterization: The basic principles, and the general approach Wout Slob RIVM, Bilthoven, Netherlands</p> <p>10:30 – 11:15 CEC4-2 Deriving generic distributions from historical data for interspecies, intraspecies, and subchronic-chronic extrapolation, and how to deal with other uncertainties Weihsueh Chiu Texas A&M University, Veterinary Integrative Biosciences, College Station, United States</p> <p>11:15 – 11:30 Coffee Break</p> <p>11:30 – 12:00 CEC4-3 Illustration of the software for probabilistic analysis (APROBA) using deoxynivalenol (DON) as a case study Matthias Herzler Federal Institute for Risk Assessment (BfR), Berlin, Germany</p>



	<p>12:00 – 13:00 Lunch Break</p> <p>13:00 – 14:30 Case Studies</p> <p><i>Students are encouraged to bring their notebook computers (fully charged) to work in pairs on a practical training exercise using the software APROBA.</i></p> <p><i>Link to the WHO Guidance on Evaluating and Expressing Uncertainty in Hazard Characterization:</i> http://www.who.int/ipcs/methods/harmonization/areas/hazard_assessment/en/</p> <p>14:30 – 14:45 Short break and preparation of final discussion</p> <p>14:45 – 16:00 Guided group discussion on applications of the probabilistic approach Matthias Herzler <i>Federal Institute for Risk Assessment (BfR), Berlin, Germany</i></p>
09h30 - 15h00 Miragaia Hall	<p>CEC 5: Co-Exposure Risk Assessment: Approaches and Options for Prioritisation and Refinement <i>Supported by CEFIC LRI</i></p> <p>Chairs: <i>Heli M Hollnagel, Switzerland and Paul Price, United States of America</i></p> <p>09:30 – 10:15 CEC5-1 Refresher: Terminology, Models and tiered Schemes in Co-Exposure Risk Assessment <i>*Heli M Hollnagel, Nathalie Vallotton</i> <i>Dow Europe GmbH, TERC, Horgen, Switzerland</i></p> <p>10:15 – 11:00 Types and sources of exposure data in co-exposure risk assessment</p> <p>CEC5-2 Data and models needed to calculate the risk of co-exposure in Europe <i>*Jacob van Klaveren¹, Hilko van der Voet²</i> ¹<i>RIVM National Institute Public Health and the Environment, Bilthoven, Netherlands</i> ² <i>Wageningen UR, Biometris, Wageningen, Netherlands</i></p> <p>11:00 – 11:30 Coffee Break</p> <p>11:30 – 12:15 CEC5-3 Value and challenges of screening level assessments of co-exposures Paul Price <i>US EPA, Computational Exposure Scientist National Exposure Research Laboratory, United States</i></p> <p>12:15 – 13:15 Lunch Break</p>



	<p>13:15 – 14:00 Higher Tier Co-Exposure Assessments – Options for Refinements</p> <p>CEC5-4 Regulatory challenges and methodological aspect for cumulative risk assessment Roland A. Solecki <i>Federal Institute for Risk Assessment, Pesticide Safety, Berlin, Germany</i></p> <p>14:00 – 15:00 Practical Exercise based on Case Studies All speakers</p>
<p>09h30 - 15h00</p> <p>D. Maria Hall</p>	<p>CEC 6: Modern Risk Assessment in Food Safety Chairs: A. Wallace Hayes, <i>United States of America</i> and Dieter Schrenk, <i>Germany</i></p> <p>9.30 - 10.05 CEC6-1 What is GRAS? *Wally Hayes¹, Claire Kruger² ¹<i>Harvard University, Andover, United States</i> ²<i>Spherix, Rockville, United States</i></p> <p>10.05 - 10.40 CEC6-2 Functional feed for farm animals from oil mill waste waters Demetrios Kouretas <i>University of Thessaly, Biochemistry-Biotechnology, Larisa, Greece</i></p> <p>10.40 - 11.15 CEC6-3 The Margin-of-exposure approach in food safety risk assessment: Acrylamide as an example Dieter Schrenk <i>University of Kaiserslautern, Food Chemistry and Toxicology, Kaiserslautern, Germany</i></p> <p>11.15 - 11.40 Coffee Break</p> <p>11.40-12.15 CEC6-4 The cadmium case: data use and what to make out of it Eugenia Dogliotti <i>Istituto Superiore di Sanità, Rome, Italy</i></p> <p>12.15 - 13.15 Lunch Break</p> <p>13.15 - 13.50 CEC6-5 Assessment of the known and the unknown: Brominated flame retardants *Marco Binaglia, Luisa Ramos Bordajandi, Ake Bergman, Alan Boobis, Sandra Ceccatelli, Jean-Pierre Cravedi, Metka Filipic, Peter Fuerst, Nicklas Johansson, Hel- le Knutsen, Miroslav Machala, Franco Merletti, Olaf Papke, Dieter Schrenk, Rolaf Van Leeuwen, Stefan Van Leeuwen <i>European Food Safety Authority, Unit on Biological Hazards and Contaminants, Parma, Italy</i></p>



	13.50 - 14.25 CEC6-6 New approaches to uncertainty in chemical risk assessment - the example of bi-sphenol A *Trine Husøy, Andy Hart, Ralph Pirow, Wim C. Mennes, Detlef Wölflé, Paul A. Fowler, Ursula Gundert-Remy, Natalie von Goetz, Rudolf A. Woutersen, Davide Arcella, Anne Theobald, Cristina Croera, Anna F. Castoldi <i>European Food Safety Authority, Parma, Italy</i> 14.25 - 15.00 Final Discussion	
16h00	Opening of the Exhibition (Exhibition Area West Ground Floor)	
17h00 - 19h00 Archive Hall	Opening Ceremony Chair: Aristidis Tsatsakis, <i>President of EUROTOX, Greece</i> 17.00-17.45 Welcome Address by Félix Carvalho President of the EUROTOX 2015 congress, Porto, Portugal Welcome Address by Aristidis Tsatsakis President of EUROTOX, Greece 17:45 – 18:00 EUROTOX Merit Award Ceremony 18:15 – 18:45 Keynote Lecture: Public Understanding of Risk Alexandre Tiedtke Quintanilha, Porto, Portugal	
19h00 - 21h00	Welcome Reception (Exhibition Area West Ground Floor)	
Monday, 14 September, 2015		
08h00 - 18h00	Congress Registration	09h00 – 18h00 Exhibition
09h00 - 09h45 Archive Hall	Keynote Lecture: <i>Chair: David Bell, Finland</i> K-1 Systems Medicine, Microbiomes and Personalised Healthcare Jeremy Nicholson <i>Imperial College London, London, United Kingdom</i>	
09h45 - 10h00	Coffee Break, Exhibition and Poster Viewing	
10h00 - 12h00 Archive Hall	Symposium S01: Experiences of Substance Evaluation under REACH - Perspectives from ECHA, Member States and Industry Chairs: Ingo Bichlmaier, <i>Finland</i> and Eva Bonefeld-Jørgensen, <i>Denmark</i> S01-1 Substance Evaluation under REACH - EU-cooperation to increase knowledge on safety of chemicals Pia Korjus <i>European Chemicals Agency, Directorate Evaluation, Helsinki, Finland</i> S01-2	



	<p>Looking beyond REACH standard information requirements: Testing requested under Substance Evaluation Gabriele Schöning <i>ECHA, Evaluation, Helsinki, Finland</i></p> <p>S01-3 Industries' perspective: How an 'identified concern' can drive 'better science?' Violaine Verougstraete <i>Eurometaux, EHS, Brussels, Belgium</i></p> <p>S01-4 Member state priorities and role in Substance Evaluation Magnus Løfstedt <i>Danish Environmental Protection Agency, Chemicals, Copenhagen, Denmark</i></p> <p>S01-5 Conclusions of substance evaluation and possible following regulatory actions Evelin Fabjan <i>European Chemicals Agency, Helsinki, Finland</i></p>
<p>10h00 - 12h00</p> <p>Infante Hall</p>	<p>Workshop W01: Natural and Process-Related Carcinogens in Food: How Should the Risk be Assessed? Chairs: Dieter Schrenk, Germany and Ans Punt, The Netherlands</p> <p>W01-1 Risk assessment of plant genotoxins *Ans Punt, Ivonne M.C.M. Rietjens <i>Wageningen University, Wageningen, Netherlands</i></p> <p>W01-2 Current risk assessment of pyrrolizidine alkaloids in food Diane Benford <i>Food Standards Agency, London, United Kingdom</i></p> <p>W01-3 Aristolochia species: A metabolomic and ethnopharmacological risk assessment focusing on local uses in Bangladesh *Michael Heinrich, J. Michl <i>UCL School of Pharmacy, Centre for Pharmacognosy and Phytotherapy, London, United Kingdom</i></p> <p>W01-4 Natural and Process-Related Carcinogens in Food: Macromolecular Adducts in Animal Models and Human Blood and Tissue Samples *Hansruedi Glatt^{1,2}, Walter Meinl², Wolfram Engst², Fabian Schumacher^{2,3}, Benjamin Sachse², Kristin Herrmann², Gitte Barknowitz², Mareike Bernau², Carolin Bendadani², Melanie Wiesner^{2,4}, Monika Schreiner⁴, Roman Tremmel⁵, Achim Bub⁶, Ulrich Zanger⁵, Bernhard Monien^{1,2} ¹Federal Institute for Risk Assessment, Berlin, Germany ²German Institute of Human Nutrition, Potsdam-Rehbrücke, Germany ³University of Potsdam, Nuthetal, Germany ⁴Leibniz-Institute of Vegetable and Ornamental Crops, Grossbeeren, Germany ⁵Dr. Margarete Fischer-Bosch-Institute of Clinical Pharmacology, Stuttgart, Germany ⁶Max Rubner Federal Research Institute of Nutrition and Food, Karlsruhe, Ger-</p>



	<p>many</p> <p>W01-5 Genotoxic and carcinogenic constituents in food and medicinal drugs: Margin of exposure, Virtually Safe Dose, TTC – which way to go? Dieter Schrenk <i>University of Kaiserslautern, Food Chemistry and Toxicology, Kaiserslautern, Germany</i></p>
<p>10h00 - 12h00</p> <p>Despachantes Hall</p>	<p>Workshop W02: Nanotoxicology Modelling for Risk Assessment: The Regulators' Dilemma Chairs: Karin Aschberger, Italy and João Paulo Teixeira, Portugal</p> <p>W02-1 Feasibility and Challenges of Health Risk Assessment of Nanomaterials *Zuzana Klöšlová¹, Karin Aschberger², Jos Bessems², Kirsten Gerloff² ¹ECHA, Evaluation, Helsinki, Finland ²European Commission, JRC-IHCP, Systems Toxicology Unit, Ispra, Italy</p> <p>W02-2 Evaluating and Modeling Oxidative Stress Responses of Immune Cells to Nanoparticles: Usefulness in Risk Assessments Anna Shvedova^{1,2} ¹WVU, Physiology and Pharmacology, Morgantown, United States ²West Virginia University, Physiology and Pharmacology, Morgantown, WV, United States</p> <p>W02-3 Achievements and perspectives of computational nanotoxicology Tomasz Puzyn <i>University of Gdansk, Faculty of Chemistry, Laboratory of Environmental Chemometrics, Gdansk, Poland</i></p> <p>W02-4 Proteomics approaches for hazard assessment of nanomaterials and for supporting NM classification *Andrea Haase¹, Marc Driessen¹, Rainer Ossig², Bryan Hellack³, Antje Vennemann⁴, Jürgen Schnekenburger², Martin Wiemann⁴, Thomas Kuhlbusch³, Wendel Wohleben⁵, Andreas Luch¹ ¹German Federal Institute for Risk Assessment, Safety of Chemicals and Consumer Products, Berlin, Germany ²Westfälische Wilhelms-Universität, Biomedical Technology Center, Münster, Germany ³Institute of Energy and Environmental Technology (IUTA) e.V., Duisburg, Germany ⁴IBE R&D gGmbH, Münster, Germany ⁵BASF SE, Ludwigshafen, Germany</p> <p>W02-5 Moving towards a safe by design approach for ENM: linking ENM relevant properties to toxicological concerns *Teresa Borges¹, Maria João Silva², Henriqueta Louro² ¹General-Directorate of Health, Occupational and Environmental Health Division, Lisbon, Portugal ²Instituto de Saúde Dr. Ricardo Jorge, Genética Humana, Lisboa, Portugal</p>
<p>10h00 - 12h00</p> <p>Noble Hall</p>	<p>Workshop W03: Opportunities to Enhance Quality and Impact of Omics Sciences Chairs: Bennard van Ravenzwaay, Germany and Thomas Hartung, United States of America</p>



	<p>W03-1 RNA-sequencing in toxicogenomics Jos Kleijnans <i>Maastricht University, Toxicogenomics, Maastricht, Netherlands</i></p> <p>W03-2 The importance of data quality to enhance the impact of omics sciences Timothy Gant <i>Public Health England, Fermie Avenue, Oxford, United Kingdom</i></p> <p>W03-3 Identification of Potential Endocrine Disrupting Chemicals Using Gene Expression Biomarkers Chris Corton <i>US-EPA, ISTD, NHEERL, Durham, NC, United States</i></p> <p>W03-4 10 years of metabolomics research: the importance of quality control *Hennicke Kamp¹, Eric Fabian¹, Markus Frericks¹, Michael Herold², Gerhard Krennrich¹, Ralf Looser², Werner Mellert¹, Gina Montoya¹, Erik Peter², Tzutzy Ramirez¹, Michael Spitzer², Volker Strauss¹, Alexander Strigun², Tilmann Walk², Bennard van Ravenzwaay¹ ¹BASF SE, Ludwigshafen, Germany ²metanomics GmbH, Berlin, Germany</p> <p>W03-5 Metabolomics: an opportunity for systemic toxicity assessment in vitro Thomas Hartung <i>The Johns Hopkins University, Center for Alternatives to Animal Testing, Baltimore, United States</i></p>
12h00 - 14h00	Lunch Break, Exhibition & Poster Viewing
12h00 - 13h00 Despachantes Hall	<p>HESI CITE Lecture Chair: Syril D Pettit, United States</p> <p>K-2 Phylotoxicology: breaking the artificial divide between human- and ecotoxicology *John Colbourne¹, Mark Viant¹, Joseph Shaw^{1,2} ¹University of Birmingham, Birmingham, United Kingdom ²Indiana University, School of Public and Environmental affairs, Bloomington, United States</p>
13h30 - 15h30 Infante Hall	<p>SpS Sponsored Symposium of the European research initiative SEURAT: Predicting Long Term Toxic Effects Using Computer Models Based on Systems Characterization of Organotypic Cultures – The NOTOX Project Chairs: Elmar Heinzle, Germany and Fozia Noor, Germany</p> <p>Welcome and introduction Elmar Heinzle, Germany</p> <p>SpS-1 Improved in vitro systems for prediction of hepatotoxicity *Magnus Ingelman-Sundberg¹, Lisa Fredriksson Puigvert¹, Sebastian Klein², Peter</p>



	<p>Peters³, Sabrina Moro¹, Catherine Bell¹, Delilah Hendriks¹, Daniel Müller², Viola Schweitzer², Fozia Noor², Elmar Heinzle²</p> <p>¹Department of Physiology and Pharmacology, Karolinska Institutet, Stockholm, Sweden</p> <p>²Saarland University, Biochemical Engineering, Saarbruecken, Germany</p> <p>³Institute of Nanoscopy, University of Maastricht, Maastricht, Netherlands</p> <p>SpS-2 Toxicoproteomics applied to in vitro investigation of liver toxicity using HepaRG cells</p> <p>*Fabrice Bertile¹, Georg Tascher¹, Daniel Müller², Sebastian Klein², Lisa Fredricks-son³, Inger Johansson³, Valery Shevchenko⁴, Christophe Chesne⁴, Magnus Ingelmann-Sundberg³, Elmar Heinzle², Fozia Noor², Alain Van Dorsselaer¹</p> <p>¹CNRS-IPHC, Université de Strasbourg, Strasbourg, France ²Saarland University, Biochemical Engineering, Saarbruecken, Germany ³Karolinska Institutet, Physiology & Pharmacology, Stockholm, Sweden ⁴Biopredic International, St Gregoire, France</p> <p>SpS-3 Model and in vitro based prediction of human hepatotoxicity</p> <p>*Jan Georg Hengstler³, Dirk Drasdo^{1,2}, Geraldine Cellière¹, Raymond Reif³, Marcel Leist⁴, Jörg Rahnenführer⁵, Regina Stöber³,</p> <p>¹INRIA, Institute for Research in Computer Science, Le Chesnay Cedex, France</p> <p>²Universität Leipzig, Interdisciplinary Center for Bioinformatics, Leipzig, Germany</p> <p>³Leibniz Research Centre for Working Environment and Human Factors, Toxicology, Dortmund, Germany ⁴University of Konstanz, TheDorenkamp-Zbinden Chair, Konstanz, Germany ⁵University of Dortmund, Department of Statistics, Dortmund, Germany</p> <p>SpS-4 Prediction of long term toxic effects by genome based network models</p> <p>*Lothar Terfloth¹, Joachim Bucher¹, Sebastian Klein², Georg Tascher³, Inger Johansson⁴, Silvia Magioni⁵, Fabrice Bertile³, Magnus Ingelman-Sundberg⁴, Alain van Dorsselaer³, Emilio Benfenati⁵, Fozia Noor², Elmar Heinzle², Klaus Mauch¹</p> <p>¹Insilico Biotechnology AG, Stuttgart, Germany ²Saarland University, Saarbrücken, Germany ³CNRS Strasbourg, Strasbourg, France ⁴Karolinska Institute, Stockholm, Sweden ⁵Instituto di Ricerche Farmacologiche Mario Negri, Milan, Italy</p> <p>Final remarks and closure Fozia Noor, Germany</p>
14h00 - 16h00	Poster Session 1 - in the Poster Areas on the Ground Floor (Exhibition Area, Foyers & Miniaturas Hall) and 1st Floor (Republic Room and Foyers)
15h30 - 16h00	Coffee Break, Exhibition & Poster Viewing
16h00 - 18h00	<p>Workshop W04: New approaches to repeated dose toxicity assessment - are we ready to replace animal testing?</p> <p>Chairs: Amaia Irizar, United Kingdom and David Bell, Finland</p> <p>W04-1 Safety Evaluation Ultimately Replacing Animal Testing: the SEURAT-1 approach? Maurice Whelan European Commission Joint Research Centre, Institute for Health and Consumer Protection, Ispra, Italy</p> <p>W04-2</p>

	<p>A High Throughput Microscopy Toxicity Pathway Reporter Platform for Chemical Safety Assessment <i>*Steven Wink¹, Bob van de Water²</i> ¹<i>Leiden University, Division of Toxicology & Leiden Cell Observatory High Content Imaging Screening Facility Leiden Academic Centre for Drug Research (LACDR), Leiden, Netherlands</i> ²<i>Leiden University, Leiden Academic Centre for Drug Research, Leiden, Netherlands</i></p> <p>W04-3 Functional intravital imaging of hepatotoxicity: comparing intact livers to 3D in vitro systems <i>*Jan Georg Hengstler¹, Dirk Drasdo², Geraldine Celliere², Seddik Hammad¹, Ahmed Ghallab¹, Raymond Reif¹, Rosemarie Marchan¹, David A. Fluri³, Jens M. Kelm³, Patricio Godoy¹</i> ¹<i>Leibniz Research Centre for Working Environment and Human Factors, Toxicology, Dortmund, Germany</i> ²<i>Institute National de Recherche en Informatique et en Automatique (INRIA), Le Chesnay Cedex, France</i> ³<i>InSphero AG, Schlieren, Switzerland</i></p> <p>W04-4 A 3D Liver Co-Culture System for Evaluating Drug-Induced Adverse Outcome Pathways Leading to Fibrosis <i>Leo van Grunsven</i> <i>Vrije Universiteit Brussel, Liver Cell Biology Laboratory, Brussels, Belgium</i></p> <p>W04-5 Exploiting data derived from non-standard methods for chemical risk assessment <i>George Daston</i> <i>Procter & Gamble, Victor Mills Society Research Fellow, Cincinnati, United States</i></p>
<p>16h00 - 18h00</p> <p>Despachantes Hall</p>	<p>Symposium S02: New Advances in In Vivo Mutagenicity Testing and Application in the Regulatory Setting Chairs: <i>Frank Le Curieux, Finland and David Kirkland, United Kingdom</i></p> <p>S02-1 The Transgenic Rodent Gene Mutation Test <i>Carol Beevers</i> <i>Covance Laboratories Ltd, Harrogate, United Kingdom</i></p> <p>S02-2 The comet assay – Peculiarities, pitfalls and interpretation <i>Brian Burlinson</i> <i>Huntingdon Life Sciences / Harlan Laboratories, Huntingdon, United Kingdom</i></p> <p>S02-3 The Pig-A Assay – Its potential applications in regulatory mutagenicity testing <i>Roland Froetschl</i> <i>BfArM Federal Institute for Drugs and Medical Devices, Genetic and Reproductive Toxicology, Bonn, Germany</i></p> <p>S02-4 Application of in vivo assays in a regulatory setting <i>David Kirkland</i></p>



	<p><i>Kirkland Consulting, Tadcaster, United Kingdom</i></p> <p>S02-5 New advances in vivo mutagenicity tests: application of the guidance of the European Food Safety Authority <i>*Riccardo Crebelli¹, Maria Carff², Juan Manuel Parra Morte², Anna Maria Rossi², Maria Vittoria Vettori², Daniela Maurici²</i> ¹<i>Istituto Superiore di Sanità, Rome, Italy</i> ²<i>European Food Safety Authority, Parma, Italy</i></p>
<p>16h00 - 18h00</p> <p>Archive Hall</p>	<p>Workshop W05: Drug Induced Liver Injury: Prediction Based on Preclinical Approaches Chairs: <i>Bob van de Water, The Netherlands and Richard Weaver, France</i></p> <p>W05-1 Predicting risk of human drug-induced liver injuries from non-clinical studies in R&D <i>Richard Weaver</i> <i>Institut de recherches internationales servier, Scientific Dept, Paris, France</i></p> <p>W05-2 DILI: from chemical and metabolism to man <i>Brian K. Park</i> <i>University of Liverpool, Translational Medicine, Liverpool, United Kingdom</i></p> <p>W05-3 Mechanistic Insights in TNF signaling and Drug-induced Liver Injury: Towards a Predictive Preclinical Toolbox <i>Bob van de Water</i> <i>Leiden University, Leiden Academic Centre for Drug Research, Leiden, Netherlands</i></p> <p>W05-4 Role of the immune system in DILI; lessons learned from animal studies <i>Raymond Pieters</i> <i>Utrecht University-IRAS, Utrecht, Netherlands</i></p> <p>W05-5 Unravelling the impact of hepatotoxic drugs by dynamic pathway modelling <i>Ursula Klingmueller</i> <i>DKFZ, B200, Heidelberg, Germany</i></p>
<p>16h00 - 18h00</p> <p>Infante Hall</p>	<p>Workshop W06: Needs and Challenges in Developing a Weight of Evidence Approach for Endocrine Disruptors Chairs: <i>Matthias Öberg, Sweden and Anna Beronius, Sweden</i></p> <p>W06-1 Endocrine disruptors and PCOS: Pathophysiological aspects <i>Evanthia Diamanti-Kandarakis</i> <i>University of Athens, Medicine, Athens, Greece</i></p> <p>W06-2 Information/testing strategies for identification of substances with endocrine disrupting properties</p>



	<p>*Sofie Christiansen¹, Henrik Holbech², Poul Bjerregaard², Ulla Hass¹ ¹Technical University of Denmark, National Food institute, Søborg, Denmark ²University of Southern Denmark, Department of Biology, Odense M, Denmark</p> <p>W06-3 Practical example of a WoE-approach with BPA and other endocrine disruptor compounds Claire Beausoleil ANSES, Risk Assessment Department, Maisons-Alfort Cedex, France</p> <p>W06-4 A proposal for systematic review and assessment of endocrine disruption *Anna Beronius¹, Marlene Ågerstrand¹, Christina Rudén¹, Åke Bergman² ¹Stockholm University, Department of Environmental Science and Analytical Chemistry, Stockholm, Sweden ²Swetox, Södertälje, Sweden</p>	
18h30 - 20h00	AstraZeneca Reception (by invitation only) , Venue: Hotel Vincci, Porto	
20h00	Speaker's Dinner (by invitation only) , Venue: Taylor's, Porto	
Tuesday, 15 September, 2015		
08h00 - 18h00	Congress Registration	09h00 – 17h00 Exhibition
08h30 - 09h30 Archive Hall	<p>Keynote Lecture Bo Holmstedt Memorial Fund Chair: Herman Autrup, Denmark</p> <p>K-3 Exposome science for public health protection and innovation Dimosthenis Sarigiannis Aristotle University of Thessaloniki, Chemical Engineering, Thessaloniki, Greece</p>	
09h30 - 10h00	Coffee Break, Exhibition and Poster Viewing	
10h00 - 12h00 Infante Hall	<p>Workshop W07: Molecular Stress at Mucosal Surfaces: Contribution to Adverse Immune Responses Chairs: Emanuela Corsini, Italy and Mojmir Mach, Slovakia</p> <p>W07-1 Skin perturbations, stress surveillance and atopy Jessica Strid Imperial College London, Medicine, London, United Kingdom</p> <p>W07-2 Epithelial stress and contact dermatitis *Stefan F. Martin, Philipp R. Esser University of Freiburg - Medical Center, Dermatology, Allergy Research Group, Freiburg, Germany</p> <p>W07-3 Immunomodulation of chemicals at the lung epithelium Jitka Stilund Hansen The National Research Centre for the Working Environment, Copenhagen, Denmark</p> <p>W07-4</p>	



	<p>Interaction between intestinal epithelial cells and intraepithelial lymphocytes in food allergy *Marianne Bol-Schoenmakers¹, Saskia Braber^{1,2}, Joost J. Smit¹, Raymond H. H. Pieters¹ ¹<i>Utrecht University, Institute for Risk Assessment Sciences (IRAS), Utrecht, Netherlands</i> ²<i>Utrecht University, Utrecht Institute for Pharmaceutical Sciences, Utrecht, Netherlands</i></p>
<p>10h00 - 12h00</p> <p>Noble Hall</p>	<p>Workshop W08: Doping in Sports: A Toxicological Perspective Chairs: Christina Tsitsimpikou, Greece and Luis Horta, Brazil</p> <p>W08-1 Reported Target Organ Toxicity of Doping Substances: Animal Studies and Human Case Reports Arif Ahmet Başaran^{1,2} ¹<i>Hacettepe University Faculty of Pharmacy, Pharmacognosy, Ankara, Turkey</i> ²<i>Hacettepe University, Turkish Doping Control Center, Ankara, Turkey</i></p> <p>W08-2 Detection of doping substances residues in biological material: a comparative approach *Andreas Tsakalof¹, Manolis Tzatzarakis², Christina Tsitsimpikou³ ¹<i>University of Thessaly, Faculty of Medicine, Larisa, Greece</i>, ²<i>University of Crete, Medical School, Heraklion, Greece</i>, ³<i>General Chemical State Laboratory of Greece, Athens, Greece</i></p> <p>W08-3 Doping substances in nutritional supplements: a possible risk for public health Christina Tsitsimpikou <i>General Chemical State Laboratory of Greece, Directorate of Energy, Industrial and Chemical Products, Athens, Greece</i></p> <p>W08-4 Medical conditions or pathologies that could require treatment with prohibited substances: the Therapeutic Use Exemption approach Luis Horta <i>Autoridade Brasileira de Controle de Dopagem, SAN Edifício DNIT, Brasília, Brazil</i></p> <p>W08-5 Fighting doping during Olympic Games: the experience gained and a glance in the future Sasho Popovski <i>Macedonian Olympic Committee, Skopje, Former Yugoslav Republic of Macedonia (FYROM)</i></p>
<p>10h00 - 12h00</p> <p>Archive Hall</p>	<p>Symposium S03: Reactive Metabolites and Molecular Mechanisms of Adverse Drug Reactions Chairs: Hilmi Orhan, Turkey and Heather M Wallace, United Kingdom</p> <p>S03-1 Reactive metabolites: In vitro screening techniques and potential extrapolation to in vivo Olavi Pelkonen <i>University of Oulu, Pharmacology and Toxicology, Oulu, Finland</i></p>



	<p>S03-2 Reactive electrophiles: Toxicity to target Michael Cameron <i>Scripps Research Institute, Molecular Therapeutics, Jupiter, United States</i></p> <p>S03-3 Role of genetic polymorphism of protective enzymes in the inactivation of reactive drug metabolites Jan Commandeur <i>VU University Amsterdam, Molecular Toxicology, Amsterdam, Netherlands</i></p> <p>S03-4 Pharmacogenomic factors affecting adverse drug reactions involving reactive metabolites Ann Daly <i>Newcastle University, Institute of Cellular Medicine, Newcastle upon Tyne, United Kingdom</i></p>
<p>10h00 - 12h00</p> <p>Despachantes Hall</p>	<p>Symposium S04: Challenges for Combined Effects of Chemical Mixtures in Risk Assessment Chairs: <i>Claudio Colosio, Italy and Antonio Hernández-Jerez, Spain</i></p> <p>S04-1 Modelling risk for chemical mixtures * Athanasios Alegakis¹, Vasilis Androutsopoulos¹, Spyros Karakitsios², Dimosthenis Sarigiannis² ¹<i>University of Crete, Laboratory of Toxicology, Heraklion, Greece</i> ²<i>Aristotle University of Thessaloniki, Department of Chemical Engineering, Thessaloniki, Greece</i></p> <p>S04-2 A prototype algorithm to calculate health-based exposure limits for a safe use of pesticides * Claudio Colosio, Stefan Mandic-Rajcevic, Federico Maria Rubino <i>Department of Health Sciences of the University of Milan, International Centre for Rural Health of the University Hospital San Paolo, and Laboratory for Analytical Toxicology and Metabonomics, Milan, Italy</i></p> <p>S04-3 Interpretation of biological monitoring data in exposure to complex mixtures * Lode Godderis^{1,2}, Radu-Corneliu Duca¹, Nathalie Grova³, Katrien Poels⁴, Jeroen Vanoirbeek¹, Brice MR Appenzeller³ ¹<i>KULeuven, Centre for Environment and Health, Leuven, Belgium</i> ²<i>DEWE, Heverlee, Belgium</i> ³<i>Luxembourg Institute of Health, LABH, Esch sur Alzette, Luxembourg</i> ⁴<i>KULeuven-DEWE, Centre for Environment and Health, Leuven, Belgium</i></p> <p>S04-4 Assessment of chemical mixtures toxicity by novel target organ-specific biomarkers * Antonio Hernández-Jerez¹, David Lozano¹, Fernando Gil¹, Tesifón Parrón^{1,2}, Raquel Alarcón², Mar Requena², Marina Lacasaña³ ¹<i>University of Granada School of Medicine, Legal Medicine and Toxicology, Granada, Spain</i> ²<i>University of Almería School of Health Sciences, Neurosciences and Health Sciences, Almería, Spain</i></p>



	³ Andalusian School of Public Health, Granada, Spain S04-5 Assessing the feasibility of mixture risk assessment – case studies with pesticides and environmental pollutants <i>*Andreas Kortenkamp, Richard Evans, Olwenn V. Martin</i> <i>Brunel University London, Institute of Environment, Health and Societies, Uxbridge, United Kingdom</i>	
10h00 - 10h30 D. Luis Hall	Exhibitor Hosted Session by QIAGEN	
12h00 - 13h00 Archive Hall	EUROTOX-SOT Debate: D In Vitro Alternatives are Ready to be Implemented and Relied Upon for Human Safety Testing <i>Chairs: John Morris, United States of America and Mumtaz Iscan, Turkey</i> From EUROTOX: Maurice Whelan <i>European Commission Joint Research Centre, EURL ECVAM, Ispra, Italy</i> From SOT: George Daston <i>Procter & Gamble, Victor Mills Society Research Fellow, Cincinnati, United States</i> Board of Trustees Distinguished Professor Department of Pharmaceutical Sciences University of Connecticut Storrs, CT 06269-3092	
13h00 - 14h00	Lunch Break, Exhibition & Poster Viewing	
14h00 - 15h00 Archive Hall	Keynote Lecture: Chair: Jyrki Liesivuori, Finland K-4 Challenges for a full replacement of animal models to assess immunotoxicity Emanuela Corsini <i>Università degli Studi di Milano, Dipartimento di Chimica, Milan, Italy</i>	
15h00 - 17h00	Poster Session 2 - in the Poster Areas on the Ground Floor (Exhibition Area, Foyers & Miniaturas Hall) and 1st Floor (Republic Room and Foyers)	
16h30 - 17h00	Coffee Break & Exhibition Viewing	17h00: Closing of the Exhibition
17h00 - 19h00 Infante Hall	Symposium S05: Environment, Epigenetic Mechanisms and Immunotoxicology <i>Chairs: Raymond H. H. Pieters, The Netherlands and Unni C. Nygaard, Norway</i> S05-1 Developmental origin of immune diseases – Environmental influences <i>*Marin Strøm¹, Thorhallur Ingi Halldorsson¹, Susanne Hansen¹, Sjurður Olsen¹, Ronald Dahl², Hans Jurgen Hoffmann³, Dorte Rytter⁴, Bodil Hammer Bech⁴, Allan Linneberg⁵, Panu Rantakokko⁶, Hannu Kiviranta⁶</i> ¹ Statens Serum Institute, Department of Epidemiology Research, Statens Serum Institute, Copenhagen, Iceland ² Odense University Hospital, The Allergy Centre, Odense, Denmark ³ Aarhus University Hospital, Department of Pulmonary Medicine and Allergy, Aarhus, Denmark ⁴ Aarhus University, Department of Public Health, Section for Epidemiology, Aarhus, Denmark ⁵ Department of Clinical Experimental Research, Glostrup University Hospital, Glostrup, Denmark ⁶ National Institute for Health and Welfare, Department of Environmental Health, Kupio, Finland S05-2	



	<p>General introduction to epigenetic mechanisms and methods Mariona Bustamante^{1,2} ¹Center for Research in Environmental Epidemiology, Barcelona, Spain ²Center for Genomic Regulation, Barcelona, Spain</p> <p>S05-3 Epigenetic mechanisms as tool for fetal programming and possible environmental influences Eva Cecilie Bonefeld-Jørgensen Aarhus University, Centre for Arctic Health, Department of Public Health, Aarhus, Denmark</p> <p>S05-4 Environmental exposures, epigenetics, and allergy Wilfried Karmaus University of Memphis, School of Public Health, Epidemiology, Biostatistics, and Environmental Health, Memphis, United States</p> <p>S05-5 Towards Understanding the Immune Mechanism of Air Pollution-Associated Asthma Kari Nadeau Stanford University, Pediatrics, Stanford, United States</p>
17h00 - 19h00 Noble Hall	<p>Symposium S06: Emerging Drugs of Abuse – An Increasing Problem also in Toxicology Chairs: Hans H. Maurer, Germany and Marilyn Huestis, United States of America</p> <p>S06-1 Responding to new psychoactive substances in Europe – the EU Early Warning System and risk assessment *Michael Evans-Brown, Roumen Sedefov EMCDDA, Lisbon, Portugal</p> <p>S06-2 In-vitro toxicokinetics of New Psychotropic Substances (NPS) Hans H. Maurer Saarland University, Dept. of Exper. & Clinical Toxicology, Homburg, Germany</p> <p>S06-3 In vivo Toxicokinetics of Novel Psychoactive Substances (NPS) in Rats *Marilyn A. Huestis¹, Marta Concheiro¹; Karl B. Scheidweiler¹; Sebastien Anizan¹, Kurt R. Lehner², Mohammad O. Bukhari², Masaki Suzuki^{3,4}, Kenner C. Rice³, Michael H. Baumann² ¹Chemistry and Drug Metabolism, Baltimore, MD, United States ²Designer Drug Research Unit, Baltimore, MD, United States ³Drug Design and Synthesis Section, IRP, National Institute on Drug Abuse and National Institute on Alcohol Abuse and Alcoholism, National Institutes of Health, Baltimore, MD, United States, ⁴Medicinal Chemistry Group, Qs' Research Institute, Otsuka Pharmaceutical Co., Ltd., Tokushima, Japan</p> <p>S06-4 Sources of Data on the Acute Toxicity Associated with the use of New Psychoactive Substance (NPS) Paul I. Dargan^{1,2}</p>



	<p>¹Guy's and St Thomas' NHS Foundation Trust and King's Health Partners, London, United Kingdom ²King's College London, London, United Kingdom</p> <p>S06-5 Fatal poisonings caused by NPS, new psychoactive substances Robert Kronstrand National Board of Forensic Medicine, Forensic Toxicology, Linköping, Sweden</p>
17h00 - 19h00 Archive Hall	<p>Workshop W09: Toxicological and Ecotoxicological Aspects of Metal Based Nanomaterials Chairs: Syed Ali, United States of America and Anne Kahru, Estonia</p> <p>W09-1 Intestinal handling of mineral nanoparticles: friends or foes from food? Jonathan Powell MRC Human Nutrition Research, Cambridge, United Kingdom</p> <p>W09-2 Engineered Metallic Nanoparticles: Pro-inflammatory response and effects on integrity of Blood-Brain-Barrier * Syed Ali¹, Susan Lantz-Mc-Peak¹, Bonnie Robinson¹, Hector Rosas_Hernandez¹, Carmen Gonzalez², William Trickler¹, Saber Hussain³ ¹NCTR, Division of Neurotoxicology, Jefferson, United States ²Universidad Autonoma de San Luis Potosi, Coordinacion para la Innovacion y la Aplicacion de la Ciencia y la Tecnologia, San Luis Potosi, Mexico, Mexico ³WP Air Force Research Laboratory, Applied Biotechnology Branch, Human Effectiveness Directorate, AFB, OH, United States</p> <p>W09-3 Novel developments in ecosafety of metal-containing nanomaterials *Anne Kahru, Irina Blinova, Angela Ivask, Kaja Kasemets, Olesja Bondarenko, Monika Mortimer, Villem Aruoja National Institute of Chemical Physics and Biophysics, Laboratory of Environmental Toxicology, Tallinn, Estonia</p> <p>W09-4 Metallic nanomaterials: From (eco)toxicity to risk assessment *Willie Peijnenburg^{1,2}, Martina Vijver¹ ¹University Leiden, Center for Environmental Sciences, Leiden, Netherlands ²National Institute for Public Health and the Environment (RIVM), Center for Safety of Substances and Products, Bilthoven, Netherlands</p> <p>W09-5 Modeling of Toxicity of Metal Oxide Nanoparticles *Robert Rallo¹, Alberto Fernández², Francesc Giralt² ¹Universitat Rovira i Virgili, Departament d'Enginyeria Informatica i Matematiques, Tarragona, Spain ²Universitat Rovira i Virgili Departament d'Enginyeria Química, Tarragona, Spain</p>
17h00 - 19h00 Despachantes Hall	<p>Workshop W10: Toxicokinetic Modelling as an Integrating Principle in Non-Animal Toxicity Testing Chairs: Ursula Gundert-Remy, Germany and Heidi Foth, Germany</p> <p>W10-1</p>



	<p>Toxicokinetic modelling: a necessary tool for quantitative risk assessment in animal-free toxicity testing Jos Bessems <i>until 2015-09-01 - EC Joint Research Centre, Systems Toxicology Unit - EURL ECVAM, Ispra, Italy</i></p> <p>W10-2 Building a toxicokinetic model using in vitro/in silico data: what is needed? Olavi Pelkonen <i>University of Oulu, Pharmacology and Toxicology, Oulu, Finland</i></p> <p>W10-3 Building a non-animal toxicokinetic model: what can be done? Case studies and lessons learned Ursula Gundert-Remy <i>Charité, Berlin, Germany</i></p> <p>W10-4 Cosmetics as a test case for non-animal testing? *Vera Rogiers, Tamara Vanhaecke <i>Vrije Universiteit Brussel, In Vitro Toxicology and Dermato-Cosmetology, Brussels, Belgium</i></p>
20h00 - 24h00	Gala Dinner at The Stock Exchange Palace
Wednesday, 16 September, 2015	
08h00 - 13h00	Congress Registration
08h30 - 10h30	<p>Workshop W11: ABC Transporters as Important Key in Xenobiotic (Pharma) Toxicokinetics Chairs: Fernando Remião, Portugal and Xavier Declèves, France</p> <p>W11-1 Induction and activation of P-glycoprotein efflux pump as a therapeutic strategy *Fernando Remião¹, Renata Silva¹, Vânia Vilas-Boas¹, Helena Carmo¹, Ricardo Jorge Dinis-Oliveira^{1,2,3}, Félix Carvalho¹, Maria de Lourdes Bastos¹ ¹UCIBIO-REQUIMTE, Lab Toxicology, Fac. of Pharmacy - University of Porto, Porto, Portugal ²INFACTS - Institute of Research and Advanced Training in Health Sciences and Technologies, Advanced Institute of Health Sciences – North (ISCS-N), Gandra, Portugal ³Department of Legal Medicine and Forensic Sciences, Fac. of Medicine - University of Porto, Porto, Portugal</p> <p>W11-2 ABC Transporters at the Blood-brain Barrier against Neurotoxicity of Xenobiotics Xavier Declèves <i>University Paris Descartes, Department of Pharmacokinetics, Paris, France</i></p> <p>W11-3 Transporters and drug-drug interactions: important determinants of drug disposition and effects Martin F. Fromm <i>Friedrich-Alexander-University Erlangen-Nuremberg, Institute of Experimental and</i></p>



	<p><i>Clinical Pharmacology and Toxicology, Erlangen, Germany</i></p> <p>W11-4 Thioxanthenes derivatives as dual inhibitors of P-glycoprotein and tumor cell growth <i>*Maria Emília Sousa^{1,2}, Andreia Palmeira¹, Ana Oliveira¹, Vanessa Lopes-Rodrigues^{3,4,5}, Marta Correia-da-Silva^{1,2}, Raquel Lima^{3,4}, Maria Helena Vasconcelos^{1,3,4}, Madalena Pinto^{1,2}</i> ¹<i>Faculty of Pharmacy, University of Porto, Porto, Portugal</i> ²<i>CIIMAR – Interdisciplinary Centre of Marine and Environmental Research, Porto, Portugal</i> ³<i>Cancer Drug Resistance Group, IPATIMUP - Institute of Molecular Pathology and Immunology of the University of Porto, Porto, Portugal</i> ⁴<i>Instituto de Investigação e Inovação em Saúde, Universidade do Porto, Porto, Portugal</i> ⁵<i>Institute of Biomedical Sciences Abel Salazar, University of Porto, ICBAS-UP, Porto, Portugal</i></p> <p>W11-5 Pharmacogenomics of ABC drug transporters: Clinical implications <i>Eugenia Yiannakopoulou</i> <i>Technological Educational Institute of Athens, Athens, Greece</i></p>
<p>08h30 - 10h30</p> <p>Infante Hall</p>	<p>Symposium S07: MicroRNAs- Mediators and Markers of Chemically-Induced Toxicity Chairs: <i>David Bell, Finland and Timothy Gant, United Kingdom</i></p> <p>S07-1 miRNAs in toxicology; the link to epigenetic effects <i>*Timothy Gant, Emma Marcxylo</i> <i>Public Health England, Fermie Avenue, Didcot, United Kingdom</i></p> <p>S07-2 MiRNAs in drug induced steatosis and its link to human non-alcoholic fatty liver disease <i>Juergen Borlak</i> <i>Hannover Medical School, Centre for Pharmacology and Toxicology, Hannover, Germany</i></p> <p>S07-3 miRNA as population variability-independent biomarkers of toxicity <i>Ivan Rusyn</i> <i>Texas A&M University, College Station, United States</i></p> <p>S07-4 MicroRNAs in drug-induced liver injury <i>Chris Goldring</i> <i>University of Liverpool, MRC Centre for Drug Safety, Liverpool, United Kingdom</i></p> <p>S07-5 Potential translational safety biomarkers for the small intestine: miRNA vs citrulline <i>Philip Hewitt</i> <i>Merck Serono, Non-Clinical Safety, Darmstadt, Germany</i></p>



<p>08h30 - 10h30</p> <p>Archive Hall</p>	<p>Workshop W12: Physiology of Infant Skin and Considerations for Quantitative Risk Assessment of Dermal Applied Substances Supported by Procter & Gamble as well as Johnson & Johnson</p> <p><i>Chairs: Susan Felter, United States of America and Georgios Stamatas, France</i></p> <p>W12-5 Quantitative Risk Assessment in the EU of Cosmetics for Babies and Children Vera Rogiers <i>Vrije Universiteit Brussel, In Vitro Toxicology and Dermato-Cosmetology, Brussels, Belgium</i></p> <p>W12-1 Infant Skin: Overview of Physiology and Maturation Antonio Torrelo <i>Hospital Niño Jesús, Dermatology, Madrid, Spain</i></p> <p>W12-2 Dermal Penetration: Models and Non-Invasive Measurements for Clinical Studies Georgios Stamatas <i>Johnson & Johnson Sante Beaute France, Issy-les-Moulineaux, France</i></p> <p>W12-3 Quantifying the skin barrier from infant to adult Maeve Kelleher <i>University College Cork, Paediatrics and Child Health, Cork, Ireland</i></p> <p>W12-4 Diapered Skin and Diaper Dermatitis: Implications for Risk Assessment Susan Felter <i>Procter & Gamble, Central Product Safety, Mason, OH, United States</i></p>
<p>08h30 - 10h30</p> <p>Despachantes Hall</p>	<p>Workshop W13: Non-Monotonic Dose-Response Curve in Hormonally Active Substances Supported by CEFIC LRI</p> <p><i>Chairs: Bruno Hubesch, Belgium and Emanuela Testai, Italy</i></p> <p>W13-1 Dose-response relationship: monotone vs non-monotone curve Emanuela Testai <i>Istituto Superiore di Sanità, Environment and Primary Prevention, Rome, Italy</i></p> <p>W13-2 Low Dose/ Dose Response Relationship of Hormonally Active Substances and their Mixture - Testing Endocrine Disruptors in Classical and Molecular Endpoints at Human-Relevant Exposure Levels *Steffen Schneider¹, Karma C. Fussell¹, Stephanie Melching-Kollmuss², Sibylle Gröters¹, Volker Strauss¹, Benazir Siddeek³, Mohamed Benahmed³, Markus Frericks², Bennard van Ravenzwaay¹ ¹BASF SE, Experimental Toxicology and Ecology, Ludwigshafen, Germany ²BASF SE, Product Safety, Ludwigshafen, Germany ³Centre Méditerranéen de Médecine Moléculaire, INSERM U895, Nice, France</p>

	<p>W13-3 Risk assessment of 'endocrine substances': Guidance on identifying endocrine disruptors Richard Green <i>Syngenta, Global Product Safety, Bracknell, United Kingdom</i></p> <p>W13-4 Risk assessment of endocrine disruptors: is there data support to deviate from the traditional approach relying on potency? Helmut Greim <i>Technical University of Munich, Freising-Weihenstephan, Germany</i></p> <p>W13-5 Regulatory Perspective on Non-Monotonic Dose-Response Curves and "Low dose effects" Niklas Andersson <i>European Chemicals Agency (ECHA), Helsinki, Finland</i></p>
10h30 - 11h00	Coffee Break
11h00 - 13h00 Noble Hall	<p>Symposium S08: Long-Term Effects of Pre- and Early Postnatal Exposure to Environmental Contaminants Chairs: Thomas Weiser, Switzerland and Martin Wilks, Switzerland</p> <p>S08-1 Exposure, effects and disease in the real world Jyrki Liesivuori <i>University of Turku, Pharmacology, Drug Development and Therapeutics, Turku, Finland</i></p> <p>S08-2 Use of biomarkers to unravel the risks from prenatal environmental exposures for later health outcomes *Greet Schoeters^{1,2}, Eva Govarts^{1,2}, Sylvie Remy^{1,2} ¹VITO, Environmental Risk and Health, MOL, Belgium ²University of Antwerp, Biomedical Dept, Antwerp, Belgium</p> <p>S08-3 Prenatal long term pesticide exposure and its association with pregnancy problems and birth defects Aristidis Tsatsakis <i>University of Crete, Medical school, Heraklion, Crete, Greece</i></p> <p>S08-4 Neurodevelopmental and neurobehavioural effects of polybrominated and per-fluorinated chemicals Martin Wilks <i>University of Basel, Swiss Centre for Applied Human Toxicology, Basel, Switzerland</i></p>
11h00 - 13h00 Infante Hall	<p>Symposium S09: Pharmacovigilance: Rational and Safe Use of Drugs Chairs: Eren (Civelek) Ozcagli, Turkey and Sini Eskola, Belgium</p> <p>S09-1 Past, present and future of pharmacovigilance: an update for current aspect Eren Ozcagli</p>



	<p><i>Istanbul University, Faculty of Pharmacy, Department of Pharmaceutical Toxicology, Istanbul, Turkey</i></p> <p>S09-2 New features of the pharmacovigilance legislation and their impact on pharmaceutical industry Sini Eskola <i>European Federation of Pharmaceutical Industries and Associations, Regulatory Affairs, Brussels, Belgium</i></p> <p>S09-3 The role of digital and social media on pharmacovigilance and their effect on personalised healthcare *Dionysios Vynias, Aristidis Tsatsakis, Manolis Tzatzarakis <i>University of Crete, Medical School, Laboratory of Toxicology, Heraklion, Greece</i></p> <p>S09-4 Importance of pharmacogenetics in adverse drug reactions Semra Sardas <i>Marmara University, Faculty of Pharmacy, Department of Pharmaceutical Toxicology, Istanbul, Turkey</i></p> <p>S09-5 Drug interactions in pharmacovigilance <i>will be replaced: Alberico L. Catapano</i></p>
<p>11h00 - 13h00</p> <p>Despachantes Hall</p>	<p>Workshop W14: Alternative Toxicity Testing Using Multicellular In Vivo Models Chairs: Marjolein Wildwater, The Netherlands and Herman Spaijk, Netherlands</p> <p>W14-1 C. elegans as robust, high throughput in vivo system for hazard assessment *Marjolein Wildwater¹, Engelen Kerkhof¹, Christien Lokman¹, Raymond Pieters^{2,3} ¹University of Applied Sciences Arnhem and Nijmegen, BioCentre, Nijmegen, Netherlands ²University of Applied Sciences, Utrecht, Innovative testing, Utrecht, Netherlands ³Utrecht University, IRAS, Utrecht, Netherlands</p> <p>W14-2 Automated zebrafish toxicology screening: effect assessment and uptake studies *Herman Spaijk¹, Peter Racz^{1,2}, Anita Ordas¹, Wouter Veneman¹, Martina Vijver³, Marjolein Wildwater⁴, Raymond Pieters⁵, Harshal Zope⁶, Alexander Kros⁶, Vasu Kantae⁷, Elke Krekels⁷, Piet Hein van der Graaf⁷, Thomas Hankemeier⁷ ¹Leiden University, Institute of Biology, Leiden, Netherlands ²ZF-screens, Leiden, Netherlands ³Leiden University, Institute of Environmental Sciences, Leiden, Netherlands ⁴Hogeschool Nijmegen, Nijmegen, Netherlands ⁵Utrecht University, Utrecht, Netherlands ⁶Leiden University, Institute of Chemistry, Leiden, Netherlands ⁷Leiden University, Institute of Pharmacy, Leiden, Netherlands</p> <p>W14-3 What determines chemical uptake by the zebrafish embryo model? *Till Luckenbach, Eberhard Küster, Wibke Busch, Stefan Scholz, Rolf Altenburger UFZ - Helmholtz Centre for Environmental Research, Bioanalytical Ecotoxicology, Leipzig, Germany</p>



	<p>W14-4 Amoeba Tastes Bitter: A Novel non Animal Model for Bitterness Research *Robin S. B. Williams¹, Marco Coccorocchio¹, Paul L. R. Andrews² ¹ School of Biological Sciences, Centre of Biomedical Sciences, Royal Holloway University of London, Egham, United Kingdom, ² Division of Biomedical Science St George's University of London, London, United Kingdom</p> <p>W14-5 Drosophotoxology takes flight: Genomic elucidation of adverse outcome pathways of mercury toxicity in the fruit fly *Matthew Rand¹, Sara Montgomery¹, Daria Vorobjkina¹, Wen Huang², Trudy F MacKay², Robert R. Anholt² ¹University of Rochester, Environmental Medicine, Rochester, United States ²North Carolina State University, Department of Biological Sciences, Genetics Program, Raleigh, NC, United States</p>
11h00 - 13h00 Archive Hall	<p>Hot Topic: HT Identifying and Prioritising Emerging Issues in Human & Environmental Sciences Chair: Ruth Roberts, United Kingdom</p> <p>Identifying and prioritising emerging issues in Human and Environmental Sciences José Manautou University of Connecticut, Pharmaceutical Sciences, Storrs, United States</p> <p>Prioritised topic: How do we ensure Non-animal Methods in Safety Assessment are fit for purpose? Beatriz Silva Lima Universidade de Lisboa, Faculdade de Farmácia, Lisbon, Portugal</p>
13h00 - 13h30 Archive Hall	<p>Closing Ceremony and Awards Presentation</p>

Detailed Online Programme: <http://eurotox2015.abstract-management.de/program/>