After the industry session, the physical Competence Centre TEDD embedded in the Centre for Cell Biology and Tissue Engineering at the ZHAW presented its core competencies, including:
- Cell biology with a focus on stem cell research, cell-based assays, cell differentiation, glycobiology, and cellular engineering
- Applied matrix biology and biophysics (macromolecular crowding, supramolecular aggregates) to develop metabolically active human models such as fatty tissue and skeletal muscle
- Establishment and analysis of tissue equivalents on scaffolds used in the fields of regenerative medicine and substance testing
- 3D cell culture model development that is suitable for the pharmaceutical industry and personalized medicine, including technologies such as bioprinting

Dr Markus Rimann concluded meeting with an outlook for the TEDD network. The goal of the network is to grow further and to closely collaborate with other networks and associations to integrate new stakeholders from different industry sectors.

Katarzyna S. Kopanska and Markus Rimann
Competence Centre TEDD, Institute of Chemistry and Biotechnology (ICBT), ZHAW Zurich University of Applied Sciences, Wädenswil, Switzerland; Centre for Cell Biology & Tissue Engineering, Institute of Chemistry and Biotechnology (ICBT), ZHAW Zurich University of Applied Sciences, Wädenswil, Switzerland (katarzyna.kopanska@zhaw.ch)

The afternoon session started with the keynote of Prof Zhongze Gu, in which he presented the activities of Southeast University (SEU) in China for the TEDD community similarly as the day before at the workshop. Then, Dr Wing Chang, director of research and development at STEMCELL Technologies, Cambridge, UK talked about robust and efficient tools for pluripotent stem cell and organoid research.

Dr Parto Toofan from REPROCELL Europe Limited, presented the company’s efforts to produce GMP-grade iPSCs for drug screening and translational medicine using a safe mRNA strategy. iPSCs produced in GMP-grade for drug screening and translational medicine have the potential to provide patient-specific scalable biologic material of various tissue types, useful for investigating the pathophysiology of rare disorders. iPSCs not only serve as excellent stem cell models, but they also can differentiate into a wide variety of cell types for preclinical studies, providing an unlimited source for the development of healthy or diseased cell models in which to study the effectiveness and toxicity of pharmaceuticals.

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man and animal health, together with the application of the 3Rs principle. Alternative in vitro models, either as stand-alone methods or as parts of an integrated testing strategy, as simple barrier models or more complex 3D cultures, were described in the context of their role in risk assessment evaluation.

**Helena Kandárová**, CEM & FCHT Bratislava, gave a lecture on “Skin irritation testing in vitro – 3D reconstructed human skin models”. Human skin models (RHSM) and respective in vitro protocols have been validated and adopted by regulatory agencies as full or partial replacements of animal experiments, i.e., for skin corrosion and skin irritation as OECD Test Guidelines (TG 431 and 439) and for chemical testing (EU REACH regulation). Recently, the ICH guideline S10 implemented RHSM for phototoxicity assessment of topically applied substances as part of a tiered testing strategy. RHSM are predominantly used for the evaluation of hazard, but they can be applied also in the risk assessment process if appropriate protocols and exposures are used. The cosmetic industry is using RHSM not only to assess the skin tolerance of their products but also to test the efficacy of novel ingredients and formulations. Most recently, RHSM were recognized as useful tools in the assessment of human skin irritation effects of medical devices (De Jong et al., 2018; Kandarova et al., 2018). Wound healing and efficacy studies conducted with RHSM were also discussed.

**Marisa Meloni**, CEO VitroScreen, Milan, Italy, presented a talk entitled: “Eye irritation: an endless challenge for in vitro science”. The process leading to replacement of the Draize rabbit test with alternative non-animal approaches, such as the Integrated Approach on Testing and Assessment (IATA) for Serious Eye Damage and Eye Irritation (OECD, 2017), has taken almost 20 years. The industrial need to evaluate the tolerance of ophthalmic formulations in a more relevant, sensitive and reproducible way compared with the in vivo Draize rabbit test has driven the development of alternative approaches that better assess these products under realistic exposure conditions. A multiparametric approach (MEA) with defined acute and repeated protocols on reconstructed human corneal epithelium (HCE) has been used to assess delayed cytotoxicity effects of eye drops and exclude any potential chronic damage to the ocular surface structure. Information on the mechanism of action and reversibility of the damage can be obtained by investigating the expression level of occludin, a functional protein of the tight junctions (Meloni et al., 2010). The results confirmed the reliability, sensitivity and predictivity of the MEA approach on HCE in detecting subclinical signs of cellular toxicity also for so-called “soft” preservatives, suggesting the need to include delayed toxicity evaluation in the biocompatibility and risk assessment of ophthalmological formulations intended for long-term use (Meloni et al., 2019).

**Arno C. Gutleb**, Environmental Research and Innovation (ER-IN) Department, Luxembourg Institute of Science and Technology, gave a lecture entitled: “Current status of in vitro models to evaluate pulmonary toxicity”. Inhalation of chemicals can have a wide range of effects from irritation in the airways, development of respiratory sensitization, to systemic effects. In vitro air-liquid interface (ALI) cell culture models have been developed to assess inhalation toxicology relevant endpoints. Such models range in complexity from single cell lines to complex 3D models consisting of several cell types. Adequate exposure techniques are essential when it comes to exposing cells grown at the ALI, and an overview of available exposure devices was presented. ALI cultures can serve as an excellent test case to show the impact of replacing animal-derived products, such as foetal calf serum, which are known to be a source of high experimental variability. The status of development, validation, and needs for further improvements of in vitro models and their potential to replace animal experiments in a regulatory context was also discussed.

**Hassan Rashidi**, NIHR Great Ormond Street Hospital Biomedical Research Centre, UCL Great Ormond Street Institute of Child Health, University College London, provided a short history of 2D and 3D cell culture techniques and described the development of a 3D platform based on hepatocytes generated from human pluripotent stem cells (hPSCs) as an in vitro tool to evaluate liver toxicity of new lead compounds. 2D-derived HLCs exhibit foetal features and transient phenotype in vitro, limiting their clinical application (Cameron et al., 2015). However, novel 3D liver organoids are not suitable for clinical application due to their reliance on animal-derived and undefined biological components. A novel platform was developed to generate hPSC-derived 3D hepatospheres (3D Heps) under xeno-free and GMP-ready conditions (Rashidi et al., 2018). The 3D Heps downregulated the expression of alpha-fetoprotein, a foetal marker, and remained metabolically active and drug-inducible for over a year in culture, providing an in vitro platform to evaluate long-term hepatotoxicity. Notably, generated 3D tissues provided critical liver support in tyrosinemia type-I animal models, indicating that these tissues may be able to treat certain liver diseases in the future.

**Yula Sambuy**, CREA-Research Centre for Food & Nutrition in Rome, Italy, introduced the most common in vitro intestinal barrier models and some promising advancements in this field. The human intestinal Caco-2 cell line has been used for thirty years to study intestinal toxicology and physiology. Derived from a colon adenocarcinoma, it can be made to differentiate on permeable substrates into a monolayer of polarized cells, coupled by functional tight and adherence junctions and expressing several transport and metabolic features of the absorptive enterocytes of the small intestine. However, cell- and culture-dependent factors strongly affect the expression of fully differentiated features (Sambuy et al., 2005). Newer approaches involve co-culture models of Caco-2 with other intestinal cell types, and 3D-intestinal preparations from normal human tissue have become available. Human induced pluripotent stem cells and organoid technologies (Nakamura and Sato, 2018) promise to be the next step towards obtaining human tissue-relevant models of the intestinal mucosa.

**Teresa CocciNi**, Laboratory of Clinical and Experimental Toxicology, ICS Maugeri, IRCCS Pavia, Italy, presented a talk entitled “Human cell-based models for neurotoxicity”. In vitro cell-based models for neurotoxicology research are moving on from simple in vitro systems with single cell types to more advanced models including 3D models (spheroids), cell co-cultures and primary human cells. Human umbilical cord lining (CL) can be used to isolate pluripotent mesenchymal stem cells (hCL-MSCs), which have strong, long-term proliferative ability (self-renewal)
and show stable amplification in vitro, are widely available without ethical restrictions, and can be easily differentiated into specific cells. In view of the limited access to human tissue, stem cells are likely to gain further importance as an alternative to primary human cells.

Doris Wilflingseder, Institute of Hygiene and Medical Microbiology, Medical University of Innsbruck, gave a talk on “Studying infectious diseases in animal free human 3D models”. 3D animal-free barrier/immune cell systems were developed to study interactions of respiratory barriers with pathogenic fungi, SARS-CoV-2 or influenza or to test novel approaches against HIV-1 infection of a mucosal model. Confocal microscopy and high content screening of 3D respiratory/mucosal samples provide high-resolution pictures and enable quantitative analyses of a high number of cells (Zaderer et al., 2019). Culture under perfused conditions results in accelerated differentiation of barrier models (Chandorkar et al., 2019). Primary respiratory cells of the bronchial and small airway epithelial tract can be cultured over a period of more than two years without losing epithelial integrity, impacting mitochondrial fitness or affecting mucus production by goblet cells. Interactions with fluorescent beads, fungi and viruses can be studied in such cultures.

An interactive debate with the participants on the perspectives and future role of alternative in vitro models on replacement concluded the event.

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Francesca Caloni1, Alessandra Cazzaniga2, Teresa Coccini3, Arno C. Gutleb4, Helena Kandárová5,6, Marisa Meloni7, Hassan Rashidi8, Yula Sambuy8, Doris Wilflingseder9,10 and Giulio Casati11

1Università degli Studi di Milano, Department of Environmental Science and Policy (ESP), Milan, Italy; 2Fondazione Alessandro Volta, Como, Italy; 3Laboratory of Clinical and Experimental Toxicology – Toxicology Unit, Istituti Clinici Scientifici Maugeri S.p.A. - SB, IRCCS Pavia, Italy; 4Luxembourg Institute of Science and Technology, Environmental Research and Innovation (ERIN) Department, Luxembourg, Luxembourg; 5Centre of Experimental Medicine (CEM), Institute of Experimental Pharmacology and Toxicology, Slovak Academy of Sciences, Bratislava, Slovakia; 6Faculty of Chemical and Food Technology (FChFT), Institute of Biochemistry and Microbiology, Slovak University of Technology (STU), Bratislava, Slovakia; 7ViroScreen, Milan, Italy; 8NIHR Great Ormond Street Hospital Biomedical Research Centre, UCL Great Ormond Street Institute of Child Health, University College London, London, UK; 9Council for Research in Agriculture and Economics (CREA) – Research Centre for Food & Nutrition, Rome, Italy; 10Institute of Hygiene and Medical Microbiology, Medical University of Innsbruck, Innsbruck, Austria; 11Lake Como School of Advanced Studies, Como, Italy