Preliminary Agenda

Advancing Alternatives to Animal Testing

SCIENTIFIC ADVISORY COMMITTEE ON ALTERNATIVE

TOXICOLOGICAL METHODS

Wednesday, September 2, 2020

10:00 AM – approximately 3:10 PM (meeting may end earlier or later). Breaks will be taken at the discretion of the chair.

Time	Agenda Item	Presenter
10:00 AM	Introductions	Nadira DeAbrew, The Proctor and Gamble Co., Chair
	Welcome and Opening Remarks	 Dr. Rick Woychik, National Institute of Environmental Health Sciences (NIEHS)/ National Toxicology Program (NTP) Dr. Anna Lowit, ICCVAM Co-chair, Environmental Protections Agency (EPA) Dr. Emily Reinke, Interagency Coordinating Committee on the Validation of Alternative Methods (ICCVAM) Co-chair, Department of Defense (DoD) Dr. Nicole Kleinstreuer, NIEHS/NTF Interagency Center for the Evaluation of Alternative Toxicological Methods (NICEATM)
	COI and Housekeeping	Dr. Elizabeth Maull, NIEHS/DNTP, Designated Federal Official
	ICCVAM – Past, Present, Future	
	20 Years of Scientific Accomplishments	Dr. Reinke
	Implementing the Strategic Roadmap	Dr. Nicole Kleinstreuer, NIEHS/NICEATM
	NTP Approaches to Assessment of Dermal Hypersensitivity: Using Alternative Methods to Predict Skin Sensitization	Dr. Dori Germolec, NIEHS/DNTP
Measuring Success of 3R Initiatives The Strategic Roadmap: What Lies on the Horizo	Dr. Suzanne Fitzpatrick, Food and Drug Administration (FDA)/Center for Food Safety and Applied Nutrition	
	• The Strategic Roadmap: What Lies on the Horizon?	Dr. Warren Casey, NIEHS/DNTP

Webinar



Interagency Coordinating Committee on the Validation of Alternative Methods

Time	Agenda Item	Presenter
	Public CommentSACATM Discussion	Dr. DeAbrew
12:30 PM	Lunch	
1:15 PM	Fostering International Partnerships	
	International Partner Updates	Dr. Lowit
	• FDA, ICH, and the 3Rs	Dr. Paul Brown, FDA/Center for Drug Evaluation and Research
	 Non-animal Test Methods for Hazard Classification – Update on UN GHS Activities 	Dr. Janet Carter, Occupational Safety and Health Administration
	Moving Away from Animal-based Antibodies	Dr. Casey
	COVID-19 Therapeutic Development with Synthetic Antibody Technology	Dr. Sachdev Sidhu, University of Toronto
	Public CommentSACATM Discussion	Dr. DeAbrew
3:25 pm	ADJOURN Day 1	Dr. DeAbrew

Thursday, September 3, 2020

10:00 AM – approximately 3:00 PM (meeting may end earlier or later). Breaks will be taken at the discretion of the chair.

Time	Agenda Item	Presenter
10:00 AM	Introductions Day 2	Nadira DeAbrew, The Proctor and Gamble Co., Chair
	COI and Housekeeping	Dr. Elizabeth Maull, NIEHS/DNTP, Designated Federal Official
	Curating and Characterizing Data for Alternative Methods Use	
	 Incorporating Variability in Animal Studies into Regulatory Frameworks and NAM Assessment 	Dr. Kleinstreuer
	 Quantitative Variability in Repeat Dose Toxicity Studies: Implications for Scientific Confidence in New Approach Methodologies 	Dr. Katie Paul-Friedman, EPA/ Office of Research and Development
	 Machine Learning in Toxicology: Towards Intelligent Access to the Content of Research 	Dr. Robert Patton, Oak Ridge National Laboratory



Interagency Coordinating Committee on the Validation of Alternative Methods

Time	Agenda Item	Presenter
	Stem Cells and Genomics for Precision Cardiovascular Medicine	Dr. Joseph Wu, Stanford University School of Medicine
	Public Comment	Dr. DeAbrew
	SACATM Discussion	
12:10 PM	Lunch Break	
12:55 PM	Computational Resources	
	Introduction	Dr. Kleinstreuer
	 Integrated Chemical Environment (ICE) & In Vitro to In Vivo Extrapolation (IVIVE) 	Dr. Shannon Bell, ILS
	 Collaborative Modeling Project for Predicting Acute Oral Toxicity (CATMoS) 	Dr. Kamel Mansouri, ILS
	Toxicokinetic and Toxicological Based Geospatial Risk Mapping	Dr. Kyle Messier, NIEHS/DNTP
	Public Comment	Dr. DeAbrew
	SACATM Discussion	
	Concluding Remarks - SACATM	Dr. DeAbrew
	Concluding Remarks	Dr. Berridge
3:05 PM	ADJOURN Day 2	Dr. DeAbrew