



Interagency Coordinating Committee on the Validation of Alternative Methods

Agenda

SCIENTIFIC ADVISORY COMMITTEE ON ALTERNATIVE TOXICOLOGICAL METHODS

Remote Meeting

Wednesday, September 21, 2022

10:00 AM – approximately 3:30 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	Dr. Nadira DeAbrew, The Procter & Gamble Co., Chair
	Welcome and Opening Remarks	Dr. Rick Woychik, National Institute of Environmental Health Sciences (NIEHS)/ National Toxicology Program (NTP) Dr. John Gordon, ICCVAM Co-chair, U.S. Consumer Product Safety Commission (CPSC) Dr. Nicole Kleinstreuer, NTP Interagency Center for the Evaluation of Alternative Toxicological Methods (NICEATM), NIEHS/DTT Dr. Warren Casey, Executive Director, ICCVAM, NIEHS/DTT
	COI and Housekeeping	Dr. Milene Brownlow, NIEHS/Division of the National Toxicology Program (DTT), Designated Federal Official
	Session I: Major ICCVAM Accomplishments in 2022	Dr. John Gordon, CPSC
	<ul style="list-style-type: none"> Public Comments 	
	<ul style="list-style-type: none"> SACATM Discussion 	
	Session II: Implementing the Strategic Roadmap: Incorporation of Alternatives and Associated Metrics	Dr. Warren Casey, NIEHS
	<ul style="list-style-type: none"> Metrics Case Studies from ICCVAM Agencies 	
	<ul style="list-style-type: none"> Communicating Progress in Advancing Alternative Methods for Regulatory Use at the FDA 	Dr. Paul Brown, U.S. Food and Drug Administration
	<ul style="list-style-type: none"> U.S. Consumer Product Safety Commission Metrics on New Approach Methods Synopsis 	Dr. John Gordon, CPSC



Interagency Coordinating Committee on the Validation of Alternative Methods

Time	Agenda Item	Presenter
	<ul style="list-style-type: none"> Animal Reduction Metrics Used by U.S. EPA Office of Pesticide Programs 	Dr. Monique Perron, U.S. Environmental Protection Agency
	<ul style="list-style-type: none"> Department of the Interior: Potential Metrics to Track and Encourage Use of Alternative Methods in Ecotoxicological Research and Testing 	Dr. Barnett Rattner, U.S. Department of the Interior
12:15 PM	Lunch Break	
12:45 PM	<ul style="list-style-type: none"> Metrics Case Studies from Industry 	
	<ul style="list-style-type: none"> Animal Metrics: Tracking New Approach Methods (NAMs) Impacts on Animal Use 	Dr. Sue Marty, Dow Chemical
	<ul style="list-style-type: none"> A Data-Driven Decision Making Framework for the Selection, Application, and Development of Advanced In-Vitro Models for Preclinical Drug Development 	Dr. Daniela Ortiz Franyuti, Roche
	<ul style="list-style-type: none"> Consideration of Alternative Methods Workgroup 	Ms. Jessie Carder, U.S. Department of Agriculture
	<ul style="list-style-type: none"> Public Comments 	Dr. DeAbrew
	<ul style="list-style-type: none"> SACATM Discussion 	
3:30 PM	ADJOURN Day 1	Dr. DeAbrew

Thursday, September 22, 2022

10:00 AM – approximately 3:30 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	Dr. Nadira DeAbrew, The Procter & Gamble Co., Chair
	COI and Housekeeping	Dr. Milene Brownlow, NIEHS/DTT, Designated Federal Official
	Session III. Validation and Establishing Scientific Confidence in New Approach Methodologies (NAMs)	
	<ul style="list-style-type: none"> ICCVAM Validation Workgroup: Updating the ICCVAM Guidance on Validation – Progress Report 	Dr. Suzanne Fitzpatrick, FDA
	<ul style="list-style-type: none"> Technical Framework for Enabling High Quality Measurements in NAMs 	Dr. Elijah Petersen, National Institute of Standards and Technology



Interagency Coordinating Committee on the Validation of Alternative Methods

Time	Agenda Item	Presenter
	<ul style="list-style-type: none"> Scientific Confidence Framework 	
	<ul style="list-style-type: none"> Biological Relevance: A Better Benchmark 	Dr. Nicole Kleinstreuer, NIEHS/DTT
	<ul style="list-style-type: none"> Variability of Reference Data 	Dr. Agnes Karmaus, Inotiv, Inc.
	<ul style="list-style-type: none"> Transparency, Data Integrity, and External Review 	Dr. João Barroso, European Union Reference Laboratory for Alternatives to Animal Testing
	<ul style="list-style-type: none"> Public Comments 	
	<ul style="list-style-type: none"> SACATM Discussion 	
12:20 PM	Lunch Break	
1:00 PM	<ul style="list-style-type: none"> Understanding Context of Use: Case Studies 	
	<ul style="list-style-type: none"> Understanding Context of Use for Medical Devices: Case Study - U.S. Food and Drug Administration 	Dr. Shelby Skoog, FDA
	<ul style="list-style-type: none"> Context of Use of Mammalian Median Lethal Dose (LD₅₀) in Ecological Assessment at U.S. Environmental Protection Agency 	Dr. William Eckel, EPA
	<ul style="list-style-type: none"> Development of a Rapid Risk Assessment Process and Software Tools to Support Air Force Operational Decision-making and Technology Acquisition 	Dr. Rebecca Clewell, U.S. Air Force
	Public Comment	Dr. DeAbrew
	SACATM Discussion	
	Session IV: Computational Resources	Dr. Helena Hogberg-Durdock, NIEHS/DTT
	Public Comment	Dr. DeAbrew
	SACATM Discussion	
3:30 PM	ADJOURN Day 2	Dr. DeAbrew

Next SACATM Meeting: September 21 – 22, 2023