



# Draft Agenda

## Workshop: BioMed21 – A Human Pathways Approach to Disease Research

**Co-organized by the National Toxicology Program and the Human Toxicology Project Consortium**

Fishers Lane Conference Center  
National Institutes of Health  
Bethesda, MD  
June 26-27, 2017

Monday, June 26

Time	Agenda Item	Presenter
8:00 a.m.	<b>Registration and continental breakfast outside meeting room</b> <i>Refreshments sponsored by the Human Toxicology Project Consortium</i>	
8:30 a.m.	<b>Welcome, summary of workshop, goals, work products</b>	Kate Willett Humane Society of the United States (HSUS)/Humane Society International (HSI)
8:45 a.m.	<b>Keynote Lecture – A Call to Action</b>	Chris Austin National Center for Advancing Translational Sciences (NCATS)
9:30 a.m.	<b>Session 1 – Setting the Stage: What is Needed and Why?</b>	Chair: Warren Casey National Institute of Environmental Health Sciences (NIEHS)
9:35 a.m.	<b>• Challenges and needs to use existing data in drug development</b>	François Pognan Novartis
9:55 a.m.	<b>• NIEHS-led U.S. roadmap to implementation of alternatives</b>	Warren Casey NIEHS
10:20 a.m.	<b>• Clinical point of view: current practices, challenges, and needs</b>	Bruce Cuthbert National Institute of Mental Health
10:45 a.m.	<b>Break</b> <i>Refreshments sponsored by the Human Toxicology Project Consortium</i>	
11:00 a.m.	<b>• FDA collaborations for addressing practical applications</b>	Suzanne Fitzpatrick Food and Drug Administration (FDA)
11:20 a.m.	<b>• Case studies in 21<sup>st</sup> century disease models</b> <ul style="list-style-type: none"> <li>◦ A tiered approach to in vitro-based compound testing: Using computational modeling, big data approaches and fit-for-purpose in vitro assays to streamline compound development</li> <li>◦ Using organoids to define key pathways in chronic obstructive pulmonary disease (COPD) pathogenesis</li> <li>◦ Organs-on-chips: A platform for advancing drug development and disease modeling</li> </ul>	Rebecca Clewell ScitoVation  Ian Adcock Imperial College London  Daniel Levner Emulate, Inc.
12:30 p.m.	<b>Lunch (on your own)</b>	

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Monday, June 26 *(continued)*

Time	Agenda Item	Presenter
1:45 p.m.	<b>Breakout Group Discussion Session 1</b>	
2:45 p.m.	<b>Break</b> <i>Refreshments sponsored by the Human Toxicology Project Consortium</i>	
3:00 p.m.	<b>Session 2 – Big Data; Information Into Knowledge Into Action</b>	Chair: Ajay Pilli Library of Integrated Network-based Cellular Signatures (LINCS) Program, National Institutes of Health (NIH)
3:10 p.m.	• <b>Overview of NIH big data projects, LINCS in particular</b>	Ajay Pilli NIH LINCS
3:30 p.m.	• <b>Biomedical Data Translator – What’s it going to take?</b>	Christine Colvis NIH Translator
3:50 p.m.	• <b>Translator and Fanconi anemia</b>	Christopher Chute NIH Translator
4:10 p.m.	<b>Break</b> <i>Refreshments sponsored by the Human Toxicology Project Consortium</i>	
4:30 p.m.	<b>Breakout Group Discussion Session 2</b>	
5:30 p.m.	<b>Break</b> <i>Refreshments sponsored by the Human Toxicology Project Consortium</i>	
5:40 p.m.	<b>Recap Day 1</b>	
6:00 p.m.	<b>Adjourn</b>	

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Tuesday, June 27

Time	Agenda Item	Presenter
8:00 a.m.	<b>Registration and continental breakfast outside meeting room</b> <i>Refreshments sponsored by the Human Toxicology Project Consortium</i>	
8:30 a.m.	<b>Session 3 – Current Tools to Support Pathway-based Decisions</b>	Chair: Suzanne Fitzpatrick FDA
8:30 a.m.	• <b>Tox21 and beyond for pharma and biomed</b>	Anton Simeonov NCATS
8:50 a.m.	• <b>Organs-on-a-chip: Applications for testing and research</b>	Lucie Low NCATS
9:10 a.m.	• <b>Primary cell-based phenotypic profiling for building human outcome pathways</b>	Ellen Berg DiscoverX Corporation
9:30 a.m.	• <b>Systems biology approach to cancer</b>	Shannon Hughes National Cancer Institute
9:50 a.m.	<b>Break</b> <i>Refreshments sponsored by the Human Toxicology Project Consortium</i>	
10:00 a.m.	• <b>How might a “pathway-based” approach help (e.g., AOPs)</b>	Kate Willett HSUS/HSI
10:20 a.m.	• <b>A network-based approach to understanding drug toxicity and its application to human liver disease</b>	Jeff Sutherland Consultant, Indiana Biosciences Research Institute
10:40 a.m.	• <b>Systems pharmacology (PredicTox)</b>	Darrell Abernethy FDA
11:00 a.m.	<b>Break</b> <i>Refreshments sponsored by the Human Toxicology Project Consortium</i>	
11:15 a.m.	<b>Breakout Group Discussion Session 3</b>	
12:15 p.m.	<b>Lunch (on your own)</b>	
1:15 p.m.	<b>Summary of Breakout Group Discussions 1-3</b>	

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Tuesday, June 27 *(continued)*

Time	Agenda Item	Presenter
2:45 p.m.	<b>Session 4 – Coordination and Support: How to Make it Work</b>	Chair: Troy Seidle HSI
	<ul style="list-style-type: none"><li>• <b>Panel discussion</b><ul style="list-style-type: none"><li>◦ Role of funding agencies</li><li>◦ Role of pharma</li><li>◦ Role of academia/SMEs</li><li>◦ Role of regulatory agencies</li></ul></li></ul>	Chris Austin NCATS François Pognan Novartis Daniel Levner Emulate Frank Weichold FDA
3:45 p.m.	<b>Break</b> <i>Refreshments sponsored by the Human Toxicology Project Consortium</i>	
4:00 p.m.	<b>Group Discussion Session 4</b>	
4:40 p.m.	<b>Break</b> <i>Refreshments sponsored by the Human Toxicology Project Consortium</i>	
4:50 p.m.	<b>Wrap-up: Summary of Discussion Questions</b>	Chairs: Warren Casey NIEHS Kate Willett HSUS/HSI
5:00 p.m.	<b>Adjourn</b>	