Current Trends in the Use of Non-Animal Data

For Risk Assessment

National Capital Area Chapter - Society of Toxicology

2011 Spring Symposium – April 19, 2011

Lister Hill Auditorium - National Institutes of Health - Bethesda, Maryland

7:45 AM Registration and Continental Breakfast

8:30 AM Welcome and Opening Comments (Program Chair: Pam Chamberlain)

8:45 – 9:20 LECTURE 1 – History and Overview of FDA Use of Non-Animal Data (Abigail Jacobs, US FDA, CDER)


9:55 – 10:30 BREAK

10:30 – 11:05 LECTURE 3 – 10 Years of ICCVAM: Challenges, Successes (Thomas Hartung, JHSPH)


11:40 – 1:00 LUNCH will be provided

Student Poster Presentations

1:00 – 1:35 LECTURE 5 – Current In Silico Methods and Their Implications in Risk Assessment (Richard Judson, US EPA NCCT)
1:35-2:10  LECTURE 6 – An NGO Perspective on Regulatory Acceptance of Non-animal Data and Related Issues (Martin Stephens, HSUS)

2:10–2:45  PANEL DISCUSSION – Global Public and Regulatory Pressures to Reduce, Refine, or Replace the Use of Animals in Biomedical Research. How Far Have We Come? How Far Can We Go?

2:45 – 3:15  Questions, discussion, and concluding remarks