Food Safety Colloquia Series
The Society of Toxicology (SOT) and FDA Center for Food Safety and Applied Nutrition (CFSAN) have partnered to provide a colloquia series that presents scientific information that is high-quality, cutting-edge, future-oriented toxicological science to provide a well-grounded foundation to inform the work of FDA employees. These sessions are open to the public to attend in person or via webcast, but these events are not a public forum for discussion of toxicology regulatory issues.

Schedule
8:00 AM–8:15 AM
Badge Pick Up
Meeting Desk

8:15 AM–8:25 AM
US FDA Welcome and Overview
Suzanne Compton Fitzpatrick, US FDA Center for Food Safety and Applied Nutrition, College Park, MD

8:25 AM–8:30 AM
Welcome from SOT
Peter L. Goering, SOT President, FDA, Silver Spring, MD

8:30 AM–9:15 AM
Problem Formulation and Scoping for Human Health Assessments
Juleen Lam, University of California San Francisco, San Francisco, CA

9:15 AM–10:00 AM
Identification and Selection of the Evidence Base for Human Health Assessments
Kathryn Guyton, International Agency for Research on Cancer Monographs Programme, Lyon, France

10:00 AM–10:15 AM
Break

10:15 AM–11:00 AM
Harmonizing Dose-Response Assessment for Cancer and Non-cancer Endpoints in Human Health Assessments
Weihsueh Chiu, Texas A&M University, College Station, TX

11:00 AM–11:45 AM
The Use of the Mechanistic Evidence in Human Health Assessments
J. Vincent Cogliano (by webinar), US EPA, Crystal City, VA

11:45 AM–12:30 PM
Roundtable Discussion
Ivan Rusyn, Moderator
All Speakers

12:45 PM–1:30 PM
Informal Discussion for FDA Employees and Speakers
Room 1A-001 (Bring your own lunch)
FDA employees are welcome up to the capacity of the room.
Contemporary Issues in Risk Assessment

Organizing Committee

Ivan Rusyn, MD, PhD, Colloquia Series Chair, Texas A&M University, College Station, TX
Suzanne Compton Fitzpatrick, PhD, Colloquium Chair, DABT, US FDA, CFSAN, College Park, MD
Ronald Chanderbhan, PhD, US FDA, CFSAN, College Park, MD
Bryan Delaney, PhD, DABT, ATS, DuPont Pioneer, Johnston, IA
Kristi Muldoon Jacobs, PhD, US FDA, CFSAN, College Park, MD
Ji-Eun Lee, PhD, DABT, Kellogg, Battle Creek, MI
James J. Pestka, PhD, Michigan State University, East Lansing, MI
Allen Rudman, PhD, US FDA, Office of Food Additive Safety, CFSAN, College Park, MD
Catherine Whiteside, PhD, US FDA, CFSAN, College Park, MD
Norbert E. Kaminski, PhD, SOT Council Contact, Michigan State University, East Lansing, MI

Other 2014–2015 SOT FDA Colloquia

Previous Colloquia

Recordings and Materials available at http://www.toxicology.org/fda

Complexities in Evaluating Human Clinical and Observational Data for Ingredient Safety Assessment: Partially Hydrogenated Oils (PHOs) As a Case Study, November 7, 2014

Application of ADME/PK Studies to Improve Safety Assessments for Foods and Cosmetics, February 23, 2015

Immunotoxicology in Food and Ingredient Safety Assessment: Approaches and Case Studies, April 14, 2015

Visit www.toxicology.org to explore other learning opportunities.