



# SOT FDA Colloquia on Emerging Toxicological Science: Challenges in Food and Ingredient Safety

May 20, 2016

## Safety Assessment Approaches in Young Children

**Allen Rudman, Chair, US FDA CFSAN, College Park, MD**  
**Elaine Faustman, Co-chair, University of Washington, Seattle, WA**

### Food Safety Colloquia Series

The Society of Toxicology (SOT) and US 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 *(All times are Eastern US, GMT-5)*

8:00 AM–8:30 AM	<b>Badge Pick Up</b> Meeting Desk
8:30 AM–8:40 AM	<b>US FDA Welcome and Overview</b> <b>Suzanne Compton Fitzpatrick</b> , CFSAN, US FDA, College Park, MD
8:40 AM–8:50 AM	<b>Welcome from SOT and Introductions</b> <b>Peter Goering</b> , SOT Past President, US FDA, Silver Spring, MD <b>Speaker Introductions, Allen Rudman</b> , US FDA, College Park, MD
8:50 AM–9:30 AM	<b>Children Matter: Using a Lifecourse Approach to Understanding Safety Assessment Needs for Children</b> <b>Elaine Faustman</b> , University of Washington, Seattle, WA
9:30 AM–10:10 AM	<b>Early Life Development of Pharmacokinetic Pathways: Framework and Case Examples with Implications for Safety Assessment</b> <b>Gary Ginsberg</b> , Connecticut Department of Public Health, Hartford, CT
10:10 AM–10:30 AM	<b>Break</b>
10:30 AM–11:10 AM	<b>Ensuring Safety for Early Life Exposures: Adequacy of Current Methods and Opportunities to Advance the Science</b> <b>Susan Felter</b> , Procter & Gamble Company, Mason, OH
11:10 AM–11:50 AM	<b>Toxicology Challenges in Lifestage-Specific Safety Assessments</b> <b>April P. Neal-Kluever</b> , US FDA, College Park, MD
11:50 AM–12:50 PM	<b>Roundtable Discussion</b> <b>Peter Goering</b> , Moderator All Speakers
1:00 PM–2:00 PM	<b>Informal Lunch for Speakers and US FDA Employees</b> <b>Room 1A-001</b> (Bring your own lunch) US FDA employees are welcome up to the capacity of the room.

**US FDA, College Park, Maryland • Live Webcast**

## Organizing Committee

**Ivan Rusyn**, MD, PhD, Colloquia Series Chair, Texas A&M University, College Station, TX

**Jason L. Aungst**, PhD, US FDA, CFSAN, College Park, MD

**Ronald Chanderbhan**, PhD, US FDA, CFSAN, College Park, MD

**Bryan Delaney**, PhD, DABT, ATS, DuPont Pioneer, Johnston, IA

**Suzanne Compton Fitzpatrick**, PhD, DABT, US FDA, CFSAN, College Park, MD

**Kristi Muldoon Jacobs**, PhD, US FDA, CFSAN, College Park, MD

**Norbert E. Kaminski**, PhD, Michigan State University, East Lansing, MI

**Jieun Lee**, PhD, DABT, Kellogg, Battle Creek, MI

**James J. Pestka**, PhD, Michigan State University, East Lansing, MI

**Allen Rudman**, PhD, US FDA, CFSAN, College Park, MD, Chair of this colloquium

**Catherine Whiteside**, PhD, US FDA, CFSAN, College Park, MD

**Peter Goering**, PhD, SOT Council Contact, US FDA, CDRH, Silver Spring, MD

**April P. Neal-Kluever**, US FDA, CFSAN, College Park, MD, for this colloquium

## Other SOT FDA Colloquia

### Upcoming Colloquia

Watch for the Announcement of the 2016–2017 Schedule

### Previous Colloquia

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

**State of the Art in the Cramer Classification Scheme and  
Threshold of Toxicological Concern, March 29, 2016**

**Role of Mode of Action in Dose-Response Assessment for Carcinogens, January 25, 2016**

**A Path Forward for Using Computational and *In Vitro* Methods for  
Food Ingredient Assessments, October 13, 2015**

**Contemporary Issues in Risk Assessment, June 17, 2015**

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

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

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

Visit [www.toxicology.org](http://www.toxicology.org) to explore other learning opportunities.