

## **Food and Drug Law Journal 2017 Symposium:** FDA and Health Behavior Regulation

October 20, 2017

Jones Day

300 New Jersey Ave, NW | Washington, DC

8:45 AM Registration and Continental Breakfast

9:15 AM Welcome and Announcements

Judy Rein, Director of Publications and Editor in Chief, FDLI

Laurie Lenkel, Director, FDA Office of the Ombudsman and Chair, Food and Drug

Law Journal Editorial Advisory Board

9:25–10:35 AM Medical Products I: Exploring the Public Health Implications of

**Off-Label Communications** 

**Author:** 

Jeffrey Chasnow, SVP & Associate General Counsel, Pfizer

**Discussants:** 

Allison Zieve, Director, Public Citizen Litigation Group, and Immediate Past

Chair, FDLI Board of Directors

Kellie B. Combs, Partner, Ropes & Gray LLP

**Moderator:** 

Amy Comstock Rick, President & CEO, FDLI

10:35–10:45 AM Networking and Coffee Break

10:45 AM-12:00 PM Medical Products II: FDA and Opioid Use

Author:

Patricia Zettler, Associate Professor, Georgia State University College of Law

**Discussants:** 

Frederick (Rick) R. Ball, Partner, Duane Morris LLP, and Member, FDLI Board of

Directors

John A. Gilbert, Jr. Director, Hyman, Phelps & McNamara, PC

**Moderator:** 

Jodi Schipper, Regulatory Counsel, Office of Compliance, Center for Drug

Evaluation and Research (CDER), FDA

12:00–1:30 PM	<b>Luncheon Keynote Peter Barton Hutt</b> , Harvard Law School and Senior Counsel, Covington & Burling LLP
	Introduced by Joseph A. Page, Professor Emeritus, Georgetown University Law Center, and Faculty Advisor, Food and Drug Law Journal
1:30-2:45 PM	Food Labeling and Consumer Perceptions
	Authors: Michael T. Roberts, Executive Director, Resnick Program for Food Law and Policy, UCLA Law Efthimios Parasidis, Associate Professor of Law and Public Health, Moritz College of Law, Ohio State University
	Discussant: Steve Armstrong, EAS Consulting Group, and Member, FDLI Board of Directors  Moderator:
	Deepti Kulkarni, Attorney, Sidley Austin LLP
2:45-3:00 PM	Networking and Coffee Break
3:00-4:30 PM	Tobacco Products: Should FDA Try to Move Smokers to E-Cigarettes or Other Less-Harmful Tobacco-Nicotine Products and, if so, How?
	Moderator/Convener: Eric Lindblom, Director, Tobacco Control and Food & Drug Law, O'Neill Institute for Global and National Health Law, Georgetown University
	Respondents:  Jeffrey Weiss, General Counsel & Senior Vice President of Government Affairs, NJOY LLC Kenneth E. Warner, University Professor of Public Health, Professor, Health Management & Policy, School of Public Health, University of Michigan Matthew L. Myers, President, Campaign for Tobacco-Free Kids Valerie Briggs Solomon, Managing Counsel, R & D Regulatory, RAI Services

Company

Reception

4:30-5:30 PM