



REGULATORY AFFAIRS

A Conversation with Leading Experts

Yale School of Public Health
60 College St., Winslow Auditorium

10 am - 12 pm (Class Lecture)
12 pm - 1 pm (Special Lecture)

3/31

PETER HUTT, J.D.

Peter Hutt is a senior counsel in the Washington DC law firm of Covington & Burling, specializing in Food and Drug law. Since 1994 he has taught a full course on Food and Drug Law at Harvard Law School. He has represented the national trade associations for the food, prescription drug, nonprescription drug, dietary supplement, and cosmetic industries. He is a member of the Institute of Medicine, and was the past Chief General Counsel to the FDA. He is the co-author the leading book *Food and Drug Law*. Peter will provide a broad overview of the history of drug regulation in the US during the past 220 years.

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NATHAN SCHACTMAN, J.D.

Nathan Schactman's practice in New York has over 30 years of experience in the defense of product liability suits, with an emphasis on the scientific and medico-legal issues that often dominate such cases. His trials, hearings, and appeals have involved prescription and over-the-counter medications, medical devices, and exposure to toxic substances from products and environmental sources.

4/14

LEE SIMON, M.D.

Dr. Lee Simon served as the FDA's Division Director of Analgesic, Anti-inflammatory and Ophthalmologic Drug Products (2001-2003). He served in multiple FDA advisory committees and has extensive experience in drug development.

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JOSEPH SCHEEREN, Pharm. D.

Joseph Scheeren is the Global Regulatory Affairs Head for the Pharmaceuticals and Consumer Care division at Bayer Healthcare. He has more than 25 years of global industry experience, including past Head of Asia Development for Bayer in Beijing. His talk will give a snapshot of the current dynamic regulatory environment in China.

4/28

YU-TE WU, Ph.D.

Yu-te Wu is the FDA's Division Director for Generic Drugs and a YSPH alumni. Dr. Wu will provide a class lecture only.