

## Speakers' Biographies

**Gregory H. Levine** is a Partner in the Washington, D.C. office of the law firm of Ropes & Gray LLP. Greg focuses his practice on FDA regulation of pharmaceuticals, biotechnology, medical devices, and cosmetics. He also counsels pharmaceutical and medical device clients on compliance with health care fraud and abuse laws. A former legislative staff member in the U.S. House of Representatives and former member of a White House healthcare reform task force, Greg has extensive FDA experience. He regularly represents clients before state and federal regulators on all phases of product life cycle and assists clients with both internal and government compliance investigations and enforcement actions. Greg is a frequent author and lecturer on subjects relating to pharmaceuticals, biologics and medical devices.

**Kirsten Mayer** is counsel in the Boston office of Ropes & Gray. Since joining the firm in 1999, she has represented clients in criminal and civil matters involving federal and state law enforcement agencies as well as in complex class action and multi-district litigation. Ms. Mayer has significant experience defending corporations and individuals in the health care industry in grand jury, civil, and administrative investigations by United States Attorneys Offices, Main Justice, the Office of the Inspector General of the U.S. Department of Health and Human Services, and various state agencies. She has represented clients in the pharmaceutical and medical device industries in off-label, anti-kickback, AWP, and Medicaid Rebate Program fraud matters.