

# FACULTY

**Randall L. Christian**, *Partner, Bowman and Brooke LLP*, is an accomplished trial attorney with significant experience in pharmaceutical mass tort product liability. He has served as trial counsel in numerous pharmaceutical product liability cases across the country including Arkansas, California, New Jersey, Pennsylvania, Nevada, and Mississippi. Most recently, he has focused his practice on the recruitment and development of testifying expert witnesses in national pharmaceutical product liability litigation. Mr. Christian has recent and substantial experience in conducting litigation risk assessments for companies preparing to market new products and marketed products where litigation is anticipated. He earned his Juris Doctor from Baylor Law School and received his Bachelor of Arts with honors from Texas Tech University.

**Mark DuVal**, *President, DuVal & Associates, P.A.* counsels pharmaceutical, medical device, biotech, food and nutritional supplement companies. Having worked extensively with drug delivery systems businesses, Mr. DuVal's practice also includes combination products, specifically drugs and devices as well as biologics and devices. He regularly advises clients in all areas of FDA laws and regulations, from pre-IDE meetings and pre-market submissions (510(k)s and PMAs, Requests for Designation) to post-marketing responsibilities (inspections, recalls, warning letters, etc.). Mr. DuVal counsels a wide range of medical device and pharmaceutical companies on sales and marketing programs that are appropriately aggressive, yet compliant, counseling on FDA's advertising and promotion regulations, the Anti-kickback Statute and False Claims Act.

**Michelle Rognien Gilboe**, *Partner, Bowman and Brooke LLP*, serves as national and regional counsel, successfully defending products including heart monitors, pumps, respiratory care devices, intravenous access devices, fluid warmers, breast implants, implantable cardiac care devices, laser devices and many more. She also has significant experience in complex class action litigation and MDL litigation including representation of multiple pharmaceutical manufacturers in various states. She has defended manufacturers of branded and generic pharmaceuticals as well as manufacturers of OTC medications.

**Cindy Khin**, *CPCU, AIC, ARM, Assistant Vice President of Claims, Medmarc Insurance Group*, is a veteran claims expert with more than 15 years of experience managing life sciences technology product liability litigation. Ms. Khin has great insights into third party liability risk management and defense techniques that help companies maintain their marketplace reputations. She has negotiated many multi-million dollar settlements, directed legal defense strategies that resulted in favorable outcomes for numerous healthcare product manufacturers and distributors, and helped healthcare technology executives preserve their companies from potentially lethal litigation.

**Kim M. Schmid**, *Managing Partner, Bowman and Brooke LLP*, defends manufacturers of medical devices and pharmaceuticals in courtrooms nationwide, including individual lawsuits, class actions, and cases involved in mass tort and multi-district litigation (MDL). As National Coordinating Counsel in medical device mass tort litigation, she has defended IDE products utilized in clinical trials as well as Class I, II and III FDA cleared devices. Kim has defended pharmaceutical cases involving generics, branded, OTC and Rx medications. She has written articles and presented at national seminars on issues specific to drug and device litigation such as preemption, successful *Daubert* challenges and other evidence based motions. She is an active member of the DRI and ABA's Drug and Medical Device Subcommittees and LifeScience Alley.

**George W. Soule**, *Partner, Bowman and Brooke LLP*, is one of seven founding partners who in 1985 formed the law firm of Bowman and Brooke. George has defended commercial, personal injury and product liability claims for nearly three decades, and has brought dozens of cases to verdict in 17 states. He coordinates product liability litigation nationwide and frequently counsels manufacturers on product safety issues. His clients include manufacturers of medical devices, farm and industrial equipment, vehicles, and consumer products. In addition to his trial experience, George served as the Chair and Vice-Chair of the Minnesota Commission on Judicial Selection for more than 11 years. Outside of the courtroom, George spends time working on political campaigns. He graduated *magna cum laude* from Harvard Law School.