

Philip Katz

Partner

Washington, D.C.

Biography

Philip Katz helps pharma/biotech companies successfully navigate FDA and related regulatory matters. He counsels them on compliance; helps anticipate and address regulatory issues in their day-to-day business operations and strategic planning; and advocates for them before the FDA, other agencies, and in court. His clients develop, manufacture, and distribute drugs and biologics, spanning from large public companies with extensive product portfolios to start-ups trying to bring their first product to market.

Phil is the former longtime head of the firm's Pharma/Biotech practice group, which has 25+ lawyers – many with years of experience at the FDA – who work collegially and in collaboration with our clients to help them achieve their business goals in a highly regulated environment in which good long-term relations with the FDA are essential. The excellence of the group is reflected in its *Chambers* ranking as a Band 1 D.C. Pharmaceutical/Medical Products Regulatory practice.

Phil and his team bring together a detailed and nuanced knowledge of the law and FDA precedents, an informed understanding of the client's business goals, and an experienced appreciation of the public policy implications to craft thoughtful, creative, and practical solutions. He and his team focus particularly on product development, approval, and lifecycle management; responding to agency enforcement activities; and



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Practices

Pharmaceuticals and Biotechnology
Regulatory

Industries

Life Sciences and Health Care

Areas of focus

Product Approvals and Dispute
Resolution

Regulatory Exclusivities, Hatch-
Waxman, and Similar Statutes

Hogan Lovells China Desk

Cell, Tissue, and Gene Therapies

Education and admissions

counseling on business transactions. He has deep knowledge of the Hatch-Waxman Act, Orphan Drug Act, and Biosimilars Price Competition and Innovation Act.

Phil served as Chairman of the board of directors of the Food and Drug Law Institute (FDLI) and as a result of his dedicated work, he received the FDLI Distinguished Service and Leadership Award. Phil's leading role in the food and drug bar is reflected in his many recognitions, including *Legal 500*, *Best Lawyers in America*, *Who's Who Legal*, *LMG Life Sciences* "Regulatory Star," and *Chambers*.

Representative experience

Represented Depomed at FDA and in court to obtain orphan exclusivity for Gralise.

Represented senior executive in negotiating and operating under FDA consent decree and subsequent DOJ investigation.

Authored successful FDA petition ensuring patent certification and opportunity for litigation and 30-month stay with competitor's 505(b)(2) NDA.

Successfully advocated to FDA Exclusivity Board for favorable decision on scope of 3-year exclusivity.

Worked with senior management to help a company develop and implement an NDA resubmission strategy after a Complete Response Letter.

Represented reference listed drug sponsor in negotiating shared REMS with generic competitor and advocating at FDA on related issues.

Represented overseas manufacturer in responding to Form FDA 483 inspectional observations and subsequent Import Alert.

Advised company with FDA strategy for approval of first product, responded to competitor petitions, and interacted with underwriters in IPO.

Awards and rankings

■ Healthcare: Pharmaceutical/Medical Products

Education

J.D., Georgetown University Law Center, magna cum laude, Order of the Coif, 1992

B.A., University of Virginia, with distinction, 1981

Bar admissions and qualifications

District of Columbia

Court admissions

U.S. Supreme Court

U.S. District Court, District of Columbia

Regulatory (District of Columbia), *Chambers USA*, 2016-2020

- Life Sciences: Regulatory/Compliance (Nationwide), *Chamber USA*, 2020
- FDA Law, *Best Lawyers in America*, 2015-2019
- Healthcare: Life Sciences, *Legal 500 US*, 2013-2015, 2017-2020
- Regulatory Star, *LMG Life Sciences*, 2014
- Regulatory, *PLC Life Sciences Cross-border Handbook*, 2011-2012
- USA – Life Sciences: Regulatory, Recommended, *PLC Which Lawyer?*, 2009

Latest thinking and events

- News
 - Restrictions on lawyer ads involving drugs and medical devices are not unconstitutional, says federal court
- Press Releases
 - Hogan Lovells names Lynn Mehler to lead the firm's Pharmaceuticals & Biotechnology Regulatory practice
- Hogan Lovells Podcasts
 - Podcast: Talking the cure
- News
 - COVID-19 Report for Life Sciences and Health Care Companies
- News
 - Eleventh Circuit decision could significantly expand scope of orphan exclusivity
- News
 - COVID-19 Report for Life Sciences and Health Care Companies (Jan - Feb 2022)