

Anishiya Abrol

Counsel

Washington, D.C.

Biography

Anishiya Abrol supports companies in their development and commercialization of cutting-edge medical therapies and tools. She advises on and negotiates a variety of strategic transactions from collaborations to in-licensing and out-licensing to maximize intellectual property assets, and then assists the company in implementing and integrating the new assets.

Anishiya has been seconded to multiple life sciences companies. She knows the issues facing in-house counsel and works closely with members of the firm's Regulatory and Privacy Practice Groups to provide integrated and practical advice.

Anishiya's practice encompasses the full life sciences life cycle, from the discovery in the lab to dispensing to a patient. She has an undergraduate science degree, and her curious nature draws her towards early stage research and development transactions and activities, including clinical trial agreements, material transfer agreements, CRO agreements, core lab agreements, sponsored research, grants, and technology transfers, both domestically and internationally. Recognizing though that a therapy's primary value lies in the hands of patients, she assists commercialization efforts with support on distribution agreements, specialty pharmacy agreements, co-promotion agreements,



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Practices

Complex Contracting

Commercial

Intellectual Property

Pharmaceuticals and Biotechnology
Regulatory

Patents

Trade Secrets and Confidential
Know-how

IP and Technology Transactions

Industries

Life Sciences and Health Care

authorized generic agreements, out-licensing arrangements, and other similar commercial activities.

Representative experience

Regularly counsels on and oversees the conduct of global clinical studies.

Advised Novartis AG in connection with its divestment of its animal health business.

Advised Novartis AG in connection with its purchase of oncology assets from GSK.

Awards and rankings

- Healthcare: Life Sciences, Next Generation, *Legal 500 US*, 2017-2020
- Healthcare: Life Sciences, Recommended, *Legal 500 US*, 2020

Latest thinking and events

- Press Releases
 - Hogan Lovells advises Lucira Health in securing Emergency Use Authorization for over the counter at-home COVID-19 test
- Published Works
 - Cell and gene therapies are driving M&A deal activity *Law360*
- Press Releases
 - Hogan Lovells advises Gilead Sciences on its collaboration with Goldfinch Bio to develop novel therapies for kidney disease
- Press Releases
 - Hogan Lovells Represents Gilead Sciences Inc. in Strategic Collaboration with Tango Therapeutics

Areas of focus

IP Rights in Transactions
Manufacturing
Pharmaceuticals and Biotechnology
IP Licensing, Commercialization, and Technology Transfer
Clinical Trials
Technology Contracts
Cell, Tissue, and Gene Therapies

Education and admissions

Education

J.D., George Mason University School of Law, magna cum laude, 2004
B.A. Biology, University of Virginia, 1997
B.A. History, University of Virginia, 1997

Memberships

Chair, Wills for Heroes Program, Virginia Bar Association (2008-2013)
Member, American Health Law Association
Member, Virginia Bar Association

Bar admissions and qualifications

District of Columbia

Virginia
