Hogan Lovells

Blake E. Wilson

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Biography

When clients need help with premarket clearance and approval of new medical devices, Blake Wilson leverages his background in the life sciences industry to smooth regulatory hurdles.

Blake has advised on 510(k) premarket notifications, de novo submissions, premarket approval applications (PMAs), and investigational device exemption applications, among other regulatory filings. He also has experience in drafting clinical trial agreements and contracts for device development, manufacturing, and associated Quality System Regulation (QSR) responsibilities.

Prior to joining Hogan Lovells, Blake worked as an associate at another international law firm and was a lead research assistant at Brown University, where he managed phase 1 and 2 pharmaceutical clinical trials. Blake takes advantage of his clinical research background to help companies navigate the FDA's clinical data requirements.

While attending law school, Blake was executive editor for the University of Pennsylvania *Journal of International Law* and published a student comment regarding the use and regulation of foreign clinical trials in the FDA's drug marketing approval process. He also served as a judicial intern to the Honorable Leonard P.

Practices

Medical Device and Technology Regulatory

Pharmaceuticals and Biotechnology Regulatory

Industries

Life Sciences and Health Care

Areas of focus

Postmarket Compliance and Enforcement Actions

Advertising and Promotion Compliance

Advisory Panel Preparation

Stark of the U.S. District Court for the District of Delaware.

Awards and rankings

- Healthcare: Life Sciences, Rising Star, *Legal 500 US*, 2020

Latest thinking and events

- News
  - FDA issues new policy for evaluating impact of viral mutations on COVID-19 tests
- Hogan Lovells Publications
  - Life sciences and health care horizons 2021
- Press Releases
  - Hogan Lovells welcomes the New Year and 25 new partner and 60 new counsel promotions
- Sponsorships and Speaking Engagements
  - In-House Impact: Life Sciences 2020
- News
  - A step in the right direction: Encouraging diversity in clinical trial populations
- Published Works
  - Pandemic accelerates expanding role of real-world evidence in FDA medical device submissions *Med Device Online*

Combination Products

Medical Devices

Pharmaceuticals and Biotechnology

Clinical Trials

Cell, Tissue, and Gene Therapies

Education and admissions

Education

M.S. Biostatistics, Columbia University, Mailman School of Public Health, 2021

J.D., University of Pennsylvania Law School, 2012

Certificate in Business Economics and Public Policy, University of Pennsylvania, Wharton School of Business, 2012

B.S., Northeastern University, summa cum laude, 2007

Memberships

Pennsylvania Bar Association, 2011 - Present

Bar admissions and qualifications

Pennsylvania

New Jersey