Hogan Lovells

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Biography

Bonella Ramsay is a dedicated life science regulatory law, IP and technology lawyer. She advised life sciences (biopharma and med tech) companies on a wide range of regulatory matters including clinical trial requirements, early access schemes, marketing authorisations, CE marking, product labelling, promotional and non-promotional activities, Brexit issues and regulatory compliance generally.

Bonella also advises on international commercial transactions in the life sciences/med-tech sectors, including complex, strategic partnering, clinical/commercial agreements spanning the life cycle of medicinal products/medical devices; from tech, transfer and licensing, research and clinical trial agreements, outsourcing of regulated functions/services, supply chain arrangements, market access, quality and pharmacovigilance agreements and public procurement/ tenders.

Prior to joining Hogan Lovells, Bonella was a partner at another major international law firm where she co-chaired the firm's Global Life Sciences Sector for 10+ years.

Representative experience

Advising various multinational pharmaceutical companies on the regulatory requirements relating to marketing authorisations and manufacturing, import and wholesale dealer licences.

Languages

English

Practices

Commercial

Pharmaceuticals and Biotechnology Regulatory

Health

Industries

Life Sciences and Health Care

Areas of focus

Agency and Distribution

Clinical Trials

Advising widely on the impact of Brexit on pharma and medical device companies and transitional arrangements.

Advising pharma companies on regulatory strategy for orphan drug applications, orphan drug regulations and intervening in the General Court where application is refused.

Advising on the regulatory aspects of the acquisition by AmerisourceBergen of the Alliance Health wholesale distribution and logistics group in Europe and the Middle East.

Leading pharma regulatory and code compliance reviews pharma companies and their European affiliates.

Advising a pharmaceutical company on the use of AI in product development.

Advising on the introduction of genomic services as a standard care within the NHS for rare diseases and cancer and the provision of genomic testing laboratory services.

Advising on multi-jurisdictional clinical trials and investigator led studies, ICFs, clinical data transparency, GCP and data privacy breaches, and secondary use of clinical data.

Providing guidance on use of new technologies and security tools to modernize clinical trial methodologies and clinical development services agreements.

Advising a global biopharma company on R&D spin-outs, collaborations, outsourcings and services arrangements, including IMI funded stem cell bank project and co-promotion partnering deals.

Advising on the regulatory aspects of the acquisition by AmerisourceBergen of the Alliance Health wholesale distribution and logistics group in Europe and the Middle East.

*Matter handled prior to joining Hogan Lovells.

Latest thinking and events

Digital Health

Medical Device Artificial Intelligence

Medical Devices

Pharmaceuticals and Biotechnology

International Regulatory Compliance

Licensing and Commercial Transactions

Education and admissions

Education

Diploma in Law, City, University of London, Diploma, 1988

B.A. Philosophy, University of Warwick, 2:1, 1986

Bar admissions and qualifications

Solicitor, England and Wales

- Webinar
  - Select IP with Hogan Lovells | Webinar no. 18: IP Transactions: Comparative discussion of royalty provisions and other key economic terms in license agreements
- News
  - To opt out or not to opt out, that is the question……
- News
  - Safety first: MHRA response to UK medical device regulation consultation
- News
  - Emerging AI issues affecting EU, UK life sciences firms
- Hogan Lovells Events
  - Health Care AI Law and Policy Summit
- Press Releases
  - Hogan Lovells advises Fressnapf on the purchase of the Italian Arcaplanet Group from Permira