

Jane Summerfield

Partner
London

Biography

Jane Summerfield is Co-Head of our Life Sciences & Health Care industry sector and leads the Hogan Lovells life sciences regulatory and commercial practice in London. She advises life sciences companies on a wide range of regulatory matters, including clinical trial requirements, early access schemes, marketing authorisations, manufacturing and distribution licences, CE marking, product labelling, advertising and marketing activities, non-promotional activities, and pricing and reimbursement.

Jane also advises on commercial contractual arrangements, including consultancy, sponsorship, co-promotion, collaboration, manufacturing, distribution, services, quality and pharmacovigilance agreements.

Jane works with clients to resolve issues with UK enforcement authorities and regulatory bodies, including the Medicines and Healthcare products Regulatory Agency (MHRA), Prescription Medicines Code of Practice Authority (PMCPA), Advertising Standards Authority (ASA), and Trading Standards.

Jane combines a scientific background with deep regulatory knowledge and a commercial approach, providing "clear responses with business impact" Chambers UK.

Representative experience

Advising various multinational pharmaceutical companies on the regulatory requirements relating to



Phone

+44 20 7296 2000

Fax

+44 20 7296 2001

Email

jane.summerfield@hoganlovells.com

Languages

English

Practices

Commercial

Marketing and Advertising

Medical Device and Technology
Regulatory

Pharmaceuticals and Biotechnology
Regulatory

Health

Industries

Life Sciences and Health Care

Areas of focus

marketing authorisations and manufacturing, import and wholesale dealer licences.

Advising an international healthcare services provider on the regulatory requirements governing the provision of remote medical services.

Advising a UK pharmaceutical manufacturer on the regulation of the authorisation, pricing and reimbursement, and sale and advertising of medicinal products in the UK.

Advising a U.S. pharmaceutical manufacture on EU and UK Standard Operating Procedures covering approval of marketing materials and activities, hospitality, promotional gifts, samples and medical advisory boards.

Advising an internet-based consumer goods retailer on competitor challenges to price claims and a Trading Standards challenge to its on-line ordering process for selling age restricted products.

Advising a leading soft drinks manufacturer on labelling and compositional legislation, advertising copy and defending challenges to products and advertising materials by regulatory authorities.

Advising a UK pharmaceutical company on a co-promotion agreement involving the joint promotion of a new treatment with another major pharmaceutical manufacturer.

Advising a medical device manufacturer on an innovative risk share agreement with a prime contractor supplying services to a Clinical Commissioning Group.

Latest thinking and events

- Hogan Lovells Podcasts
 - Podcast: Talking the cure
- News
 - COVID-19 Report for Life Sciences and Health Care Companies
- News

Advertising and Copy Clearance

Agency and Distribution

Digital Health

False Advertising and Unfair Competition

Food and Beverages

Food Advertising and Regulation

Global Regulatory in the UK

Medical Devices

Pharmaceuticals and Biotechnology

Product Compliance

Product Development and Approval

Sales Promotions

Cell, Tissue, and Gene Therapies

Education and admissions

Education

MA Biological Sciences, University of Oxford, New College

PgDL and LPC, BPP University Law School

Bar admissions and qualifications

Solicitor, England and Wales

- Video - The latest in UK medical device regulation
- News
 - COVID-19 Report for Life Sciences and Health Care Companies (Jan - Feb 2022)
- News
 - COVID-19 Report for Life Sciences and Health Care Companies (December 2021)
- News
 - UK ABPI transfer of value disclosures: transitioning from consent to legitimate interests