Marie Ramm LL.M.

Marie Ramm, LL.M., born in 1993, studied law at the University of Augsburg from 2013 to 2018 with a focus on bio -, health - and medical law and so deepened her knowledge in pharmaceutical and medical device law. During her studies, Ms. Ramm worked continuously as a research assistant in a well-known Munich law firm in hospital and social law. Ms. Ramm completed her legal clerkship at the Higher Regional Court of Munich from 2018 to 2020, including stations at the Government of Upper Bavaria in the area of environment, health and consumer protection as well as at a renowned international pharmaceutical law firm in Frankfurt am Main.

Ms. Ramm was admitted as a attorney in 2021 and joined the pharmaceutical and healthcare team at Meisterernst Rechtsanwälte in 2022. She previously worked in the legal department of a university hospital in Munich. In addition to general medical law, in particular medical malpractice law, the focus of her work included providing legal advice to employees on the professional law of the medical professions as well as to clinics and the board of directors on all aspects of research and inventions in the context of clinical trials, e.g. drafting and negotiating contracts.

Ms. Ramm's practice focuses on all matters relating to pharmaceutical and medical device law as well as related fields, e.g. intellectual property law (competition law) and compliance (including anti-corruption, professional law for doctors, medical and medical criminal law and industry codes).

Her advice covers the entire life cycle of a product:

- Negotiation and drafting of research and development contracts, clinical trial agreements and cooperation agreements
- Securing and distributing rights to research results (in particular intellectual property "IP" and know-how) and compliance with legal requirements
- Regulatory issues, including the distinction between medicinal products, medical devices, foodstuffs and biocides
- Questions regarding the authorization, certification, approval, notification or registration of products
- Manufacturing and distribution of the product (including license, manufacturing and distribution agreements, including pharmacovigilance), including checking packaging for legal compliance
- Reviewing the possibilities of communication, information and advertising in connection with medicinal products and medical devices (e.g. marketing materials, campaigns, corporate/image advertising, scientific presentations and articles) in terms of the law on medicinal products advertising
- Advice and support in dealing with authorities and competitors in competition disputes or proceedings

In addition to successfully completing the specialist lawyer course in medical law, Ms. Ramm completed the part-time Master's degree course in medical law in 2022/2023 with the additional qualification "Master of Laws" (LL.M.). The subject of her thesis was the gender distribution in commercial pharmaceutical clinical trials in Europe.

Ms. Ramm advises in German and English.

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