Cooley

Jessica Koffel

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Brussels

Life Sciences and Healthcare Regulatory
Life Sciences
Healthcare
Medtech
Digital Health
Biotechnology

Jessica focuses on European law and the regulation of medicinal products and medical devices, including in vitro diagnostic medical devices. She helps guide life sciences companies through the regulatory processes and technical requirements governing the approval and marketing of their products – from classification, clinical trials and audits to promotion and marketing. Jessica provides strategic advice from the early development stages to placing on the market and post-marketing activities. She also assists clients in their interactions with regulatory authorities, notified bodies and business partners.

Prior to joining Cooley, Jessica worked as an associate in leading international law firms, where she focused on the EU regulation of life sciences products and chemicals.

Jessica is a native English and French speaker.

Education

Université Catholique de Louvain
Master in European Law, 2017

Admissions & Credentials

Brussels (A List)