



Christine E. Kirk

Partner

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OVERVIEW

Christine Kirk is a partner and a member of the Healthcare and FDA practice group. Prior to joining the firm, Christine was a counsel at a Washington, DC-based boutique life sciences law firm where she focused her practice on providing clients strategic advice in the biopharma space. She worked with a multidisciplinary team of legal, scientific, and communications colleagues to assist biopharma clients from early drug development through post-approval compliance, including REMS requirements. She also assisted clients with drafting and pre-submission review of new drug application (NDA) and supplemental NDA (sNDA) documents, agency meeting requests and responses, citizen petitions, etc.

This is a temporary biography. A complete biography will be available soon.

PROFESSIONAL BACKGROUND

Prior to her work as counsel Christine was an associate at the Washington, DC, office of an international law firm. During her time as an associate Christine was a member of the food and drug law group where she assisted clients with FDA-related legal and regulatory issues. Christine also worked for the US Food and Drug Administration as a regulatory counsel/attorney advisor. While at the FDA she worked in the Office of Regulatory Policy, Center for Drug Evaluation and Research where she led, co-led, and participated in various rulemaking, guidance, and citizen petition workgroups.

AREAS OF FOCUS

- Healthcare and FDA