Expert in Regulatory Affairs and CMC joins Arex Advisor

Josefin Jönsson joins Arex Advisor from a position as Consultant Drug Development, CMC at ProPharma Group.

Josefin holds a MSc Pharmacy from Uppsala University and has been working in the pharmaceutical sector for almost 20 years. Josefin started her career at the Swedish Medical Products Agency where she was an assessor of product information. After that she moved to the pharmaceutical industry to a position as regulatory affairs manager at Fresenius Kabi, before she became a consultant in global projects and drug development at Sofus Regulatory Affairs, nowadays ProPharma Group.

Josefin is specialized in regulatory CMC and global regulatory affairs. She is especially skilled in the role as project leader, and has extensive experience from working with pharmaceutical companies, focusing on the coordination of CMC documentation for EU and US submissions from Phase I to new drug applications.

Linda Thunell, CEO at Arex Advisor, says: “We are so happy that Josefin has joined the Arex team. She is a true team player, highly appreciated by clients as well as colleagues. While being an expert in CMC and Regulatory affairs, she is also a dedicated and experienced project leader of the type that can be the deciding factor for a project's success. Her professional work and mindset will for sure contribute to the success of our clients’ projects but also to our company.”

About Arex Advisor:

Arex offers strategic advice and operational support in pharmaceutical research and business development. Our mission is to help leaders and companies bring their projects to commercialization. We are passionate about the development process, leading to submission and a successful commercialization of important medicines – which ultimately helps to improve the lives of patients.

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