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History and Mission

KLIFO, established in 1994, is an integrated drug development consultancy with offices in Denmark and Germany. From our offices in Munich, we offer consulting and operational services for strategic drug development, CMC and regulatory affairs and clinical trial management. **Our ambition is to support our clients to accelerate the progress of their drug development project effectively.**

Highlights

At KLIFO, we strive to build flexible and trust-based partnerships in which we work as an integrated part of your team.

Over the past two decades, KLIFO has grown to bring all necessary drug development expertise and competences under one roof. In 2019, KLIFO acquired medicomp GmbH, expanding our skills and expertise.

We support our clients in navigating the drug development process from end to end to help them bring their life-changing products closer to market.

KLIFO is a member of PSN Research (www.psnresearch.com), a worldwide operating CRO, and is able to offer services throughout Europe, the US and Canada.

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BVMA member since 1996
Audits passed in 2000, 2003, 2006, 2009, 2012, 2015 and 2018

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Services

Integrated drug development consultancy from KLIFO

KLIFO Drug Development Counselling increases the value of development projects and operational strategies by bringing together a wide range of expertise across all key R&D functions to build robust drug development strategies.

KLIFO Clinical Operations Solutions navigates the unique complexities of individual clinical studies to ensure the shortest possible development times, reduce risks and constrain costs. Our experienced staff leverage their in-depth expertise to create scalable and flexible solutions to meet your specific needs.

KLIFO CMC Development Solutions applies scientific excellence to ensure the optimal development of your drug candidate. Our experts ensure that your drug presentation is suitable for the relevant preclinical and clinical stages.

KLIFO Regulatory Affairs Solutions merges competence and experience to assist you in navigating all regulatory requirements. With our extensive experience in interacting with national and regional agencies, we can support all your regulatory affairs needs.

KLIFO Clinical Trial Supply Solutions optimises clinical trial supply by offering proactive management throughout the supply and logistics process. We have international reach, highly flexible solutions and are compliant with EMA, FDA and national requirements.

KLIFO Pharmacovigilance Solutions meets the complex regulatory requirements of drug safety to ensure optimal compliance with regulations and guidelines.

KLIFO Quality Assurance Solutions helps manage and ensure the correct level of quality and integrity across every stage of your drug development project.