Senior Clinical Project Manager

Join an international drug development consultancy and contribute with strategic advice and operational support to clients based in Europe, the US and Asia.

At KLIFO, you'll be part of a highly specialised team of experts who value workplace flexibility and collaborate to help our life-science clients across different therapeutic areas realise their unique projects.

Become part of an experienced and dedicated team

KLIFO Clinical Operations Solutions (COS) ensures operational and strategic advice and support to our clients with tasks of varying complexity. Our team of highly skilled Project Managers work at our offices in Denmark, Sweden and Germany and manages global clinical phase 1-3 trials in multiple indications and therapeutic areas. In addition, we also provide advice and sponsor oversight to a diverse segment of clients (pharma and biotech industries) and we work from the client´s office on occasions if required.

In the role as Senior Clinical Project Manager, you'll contribute by:

- Delivering clinical trial(s) studies as defined in the trial outline/protocol, to the agreed quality, budget, timelines and resources.
- Leading the clinical trial team(s), coordinate and manage all activities related to clinical trials studies.
- Communicating timelines, identify milestones, handle trial budget and report progress for the clinical trials to sponsor and COS management team.
- Possible participation in global development teams on compound level giving input to clinical development plans and operational strategies across a clinical program and taking on a main role in larger programs.
- Being mentor for Project Manager colleagues within COS

Location
This position can be located in our office in Denmark or Sweden

Employment
Full time

Deadline for application
17 July 2024

Contact
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About KLIFO
- 150+ employees
- Offices in Denmark, Sweden, Germany and The Netherlands
- Headquarter in Glostrup, Denmark

KLIFO is a leading North-European drug development consultancy with significant experience inpartnering with biotech and pharmaceutical companies. We provide end to end solutions and support our clients with strategic advice and operational support across therapeutic areas and disciplines.

Your background and qualifications

You have a positive, proactive, flexible and self-driven personality and like to work in a consulting environment with a high variety of tasks and responsibilities. Your communication skills and ability to listen to clients needs will be a substantial asset in the collaborations.

Furthermore, you have:

- M.Sc./B.Sc. in the life science field and a minimum of 7 years in a similar position in the pharmaceutical industry/CRO
- Extensive experience within clinical trial management and understanding the principles of clinical drug development incl. solid knowledge of GCP guidelines, applicable regulatory requirements, medical terminology and clinical trial processes.
- Ability to translate client´s need into a solid project plan.
- Ability to work independently without supervision of a colleague.
- Are skilled in supervising and coaching of non-senior personnel.

Why join KLIFO?

- Join an organisation where we value people and their expertise as the greatest asset.
- Enter a flexible workplace with a culture based on trust, transparency and respect.
- Work with some of the most experienced and dedicated colleagues in the life-science industry
- Contribute with your expertise across different therapeutic areas.
- Develop tailor-made solutions based on cross-disciplinary collaboration.
- Cultivate successful relationships with our clients.
- Be part of an organisation that sees knowledge-sharing as the road to success.

Share your application

Share your application with us at job@klifo.com marked Senior Clinical Project Manager no later than 17th of July 2024.

If you have questions about the position, please reach out to Randi Rahbæk, Senior Clinical Research Director at Randi.rahbaek@klifo.com or +45 44 778 704 for more information.

KLIFO processes your application and all related personal data exclusively for the specific hiring process. Your data is processed as confidential information, cf. the current data protection law (GDPR).