

KLIFO is looking for a talented and committed Project Manager (PM) based in Denmark

KLIFO is expanding our office in Denmark and wants to engage Project Managers into a dynamic and experienced team within Clinical Operations Solutions. The people we want to engage like to work in a consulting environment and have a positive, proactive, flexible, self-driven and self-confident personality. We can offer a highly flexible, free and trustful working climate with exciting customers and projects among competent colleagues where your knowledge, experience and contribution are valuable and highly appreciated.

The position as Project Manager:

The Project Manager (PM) is responsible for the delivery of an agreed trial as defined in the trial outline/protocol, to the agreed quality, budget and timelines:

- Lead the clinical trial team and coordinate and manage all activities related to clinical trials
- Planning, delivery and management of clinical trial activities (start-up, operational conduct and closure) according to SOPs and a responsibility split with Sponsor
- Liaison with clinical trial team members regarding trial design and protocol preparation and ensure the production of the final trial protocol
- Lead the Risk Based Quality Management (RBQM) process including risk assessment and risk controls
- Development of project plans for the conduct of the clinical trial.
- Communicate timelines, identify milestones, handle trial budget and report progress for the clinical trials to sponsor and to COS management team

The qualifications of the Project Manager:

- BSc/MSc in the life sciences field and a minimum of 3-4 years in a similar position in the pharmaceutical industry/BioTech/CRO
- Experience within clinical trial management, preferably within phase 1
- Knowledge of GCP guidelines, applicable regulatory requirements, medical terminology and clinical trial processes
- Ability to translate client's needs into a solid project plan

- Excellent communication skills (written and verbal) in English as well as in native language (if other than English)
- Microsoft Office Skills

We offer:

- Work within different therapeutic areas and with tasks of varying complexity
- Work with a heterogeneous client pool (pharmaceutical companies, established biotech, inexperienced biotech, investigators/academia)
- Build international client relations
- Use and elaborate your competences and experience
- Work in an interactive, flexible and positive working environment

Location:

KLIFO has offices in both Denmark, Germany, Sweden and The Netherlands This position will be located at our office in Glostrup, Denmark.

Contact:

For more information, please contact Jennie Wilborgsson, Clinical Research Director, at +45 93 63 88 20

Applications should be sent to:

job@klifo.com marked Project Manager. 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).

Deadline: 23 November 2020 (although applications are reviewed on an ongoing basis)

KLIFO is an established and **integrated drug development consultancy** with offices in Denmark, Germany and Sweden. We provide end-to-end expert capabilities, enabling our partners to maximise opportunity, mitigate risks, drive innovation and achieve efficient project advancement. **For more information, visit KLIFO.com**