KLIFO
A step in the right direction

CLINICAL TRIAL Services

Project Managers

KLIFO is expanding and wants to engage Project Managers into a dynamic and experienced team within Clinical Trial Services.
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 projects among competent colleagues where your contribution is valuable and makes a difference.

KLIFO is a service provider of drug development competences and operational services to the pharmaceutical, biotech and med-tech companies in Europe, US and Asia. Our services include Clinical Trial Supply, Clinical Trial Services, Regulatory Affairs, Pharmacovigilance, Chemistry and Manufacturing Control and Drug Development Counselling.

The position as Project Manager:
The Project Manager is overall responsible for managing the clinical trials, i.e.:

- Management of assigned project in all phases, from start-up to the end of the trial
- Leadership of people involved in the project
- Investigational site selection in order to assure the quality and conformity of the sites
- Attending cross-functional project team in relation to trial(s)
- Operational and scientific input to key project documents
- Continuous relationship with the Principal Investigators and Sponsor to assure the success of the trial in terms of enrolment and quality
- Management and resolution of issues occurring during the trial
- Generation and management of Project Plans
- Investigators and Monitors training and organization of meetings
- Elaboration of trial specific procedures
- Participation in data management activities

Education, experiences, knowledge and skill:
The Project Manager should possess the following qualifications:

- MSc in the life sciences field and a minimum of 5-6 years of overall experience out of which 2 years must be in project management in the pharmaceutical industry /Biotech/CRO
- Extensive knowledge of GCP guidelines, applicable regulatory requirements, medical terminology and clinical trial processes
- Willingness to travel
- Demonstrated project management skills including the ability to plan a project and operate within plan and budget
- Excellent verbal and written communication skills
- Computer skills, ability to develop and maintain excel spreadsheets and to generate PowerPoint presentations

In addition to the above-mentioned qualifications, the candidate is a dedicated and collaborative team player that possesses excellent planning skills and is fluent, spoken and written, in English and in a Scandinavian language.

We offer:

- Work within different therapeutic areas and with tasks of varying complexity
- Work with a heterogeneous client pool
- Build international client relations
- Use and elaborate your competences and experience
- Work in an interactive, flexible and positive working environment

As KLIFO is a smaller service provider you will experience a higher level of transparency, influence and a good possibility for individual planning.

Location:
KLIFO is located at Smedeland 36, 2600 Glostrup.

Contact:
For more information, please contact Klas Rådberg, Clinical Research Director of CTS at +45 44 222 935

Applications should be sent to:
Mette.Widen@klifo.com, marked Project Manager

Deadline:
24th August, 2018

For more information on KLIFO, please visit: **www.klifo.com**