KLIFO
A step in the right direction

CLINICAL TRIAL Services

Clinical Research Associates

KLIFO A/S is looking for talented and committed persons to strengthen our CTS department.

KLIFO is expanding and wants to engage Clinical Research Associates 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. The areas covered by KLIFO include Clinical Trial Supply, Clinical Trial Services, Pharmacovigilance, Regulatory Affairs and Drug Development Counselling.

The position:
The Clinical Research Associate (CRA) is responsible for the pro-active site management including monitoring activity of trials, i.e.:

- Participation in the preparation of trial documents for submissions to Competent Authorities and Ethics Committees/Institutional Review Boards
- Visiting investigator and investigational site before a specific trial: pre-trial/site assessment visits
- Performing initiation, monitoring and close-out visits
- Elaboration of trial specific procedures
- Support to data management activities
- Continuous relationship with the Principal Investigators and trial staff to assure the success of the trial in terms of enrolment and quality
- Assist in ensuring site compliance with protocol and trial objectives
- Work in the clinical trial team, reporting to a project manager for trial related deliverables

The ideal candidate is a dedicated and collaborative team player, possesses excellent planning skills and is fluent, spoken and written, in English and Danish.

Education, experiences, knowledge and skill:
The Clinical Research Associate should possess the following qualifications:

- B.Sc. in the life sciences field or CRA specific diploma and a minimum of 2 years in a similar position in the pharmaceutical industry/BioTech/CRO
- Knowledge of GCP guidelines, applicable regulatory requirements, medical terminology and clinical trial processes
- Willingness to travel
- Excellent communication skills
- Excellent computer skills, ability to develop and maintain excel spreadsheets and to prepare PowerPoint presentations

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 and positive working environment

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

Contact:
For more information, please contact Annette Stender, Clinical Research Director of CTS at +45 44 222 928.

Applications should be sent to:
Mette.Widen@klifo.com, marked Clinical Research Associate.

Deadline:
01 August, 2017

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