KLIFO
A step in the right direction

CLINICAL TRIAL Services

Clinical Research Associate

KLIFO A/S is looking for a talented and committed Clinical Research Associate.

KLIFO is expanding and wants to engage a Clinical Research Associate 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 collaborative 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 as Clinical Research Associate

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 qualifications of the CRA

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
- Extensive knowledge of GCP guidelines, applicable regulatory requirements, medical terminology and clinical trial processes
- Willingness to travel
- Excellent verbal and communication skills
- Excellent computer skills, ability to develop and maintain excel spreadsheets and to elaborate PowerPoint presentations
- Strong organizational skills with attention to details

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

The offer

KLIFO offers a job where you can:

- 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

As KLIFO is a smaller service provider, you will experience a high 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 Tina Hjorth, Clinical Research Director of CTS at +45 44 222 934.

Applications should be sent to:

job@klifo.com, marked Clinical Research Associate

Deadline:

21 October 2018