

## 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 proactive 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:

17 September 2018

