CLINICAL OPERATIONS SOLUTIONS

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Clinical Research Associate

KLIFO is looking for a Clinical Research Associate (CRA) for a one year maternity cover for our office in Glostrup

KLIFO would like to engage a CRA in a one year maternity cover position. 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 and trustful working climate with exiting projects among competent colleagues where your contribution is valuable and makes a difference.

The position as CRA:
The 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 CRA 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 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 organisational 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.

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

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 has offices in both Glostrup, Denmark and Munich, Germany. This position is located at our office in Glostrup.

Contact:
For more information, please contact Tina Hjorth, Clinical Research Director of COS at + 45 44 222 934.

Applications should be sent to:
job@klifo.com marked 'Clinical Research Associate'.

Deadline: 01 September 2019

KLIFO is an established and **integrated drug development consultancy** with offices in Denmark and Germany. 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**

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KLIFO
An Integrated Drug Development Consultancy