



Experienced Data Manager/Programmer

KLIFO is expanding and developing the eClinical department to provide eClinical solutions as an integrated part of Clinical Trial Services.

KLIFO wants to engage an experienced Data Manager/Programmer to actively engage in building the department and to support the project team to get oversight over data, including metrics for risk-based approaches.

The person would like to work in a consulting environment and has 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 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 Data Manager/Programmer:

The Data Manager/Programmer is overall responsible for the data during clinical trials, i.e.:

- EDC set-up and management
- General data management tasks e.g. input to protocol, validation plan, DM plan, data cleaning and coding
- Validation of computerized systems including writing test protocols, perform tests and prepare documentation
- Visualization of data and metrics for study oversight
- Explain and demonstrate DM related tasks during client meetings
- Attending cross-functional project team in relation to trial(s)
- Write/review SOPs to support eClinical activities

Education, experience, knowledge and skills:

The Data Manager/Programmer should possess the following qualifications:

- Bachelor or master degree in computer science and a minimum of 3-5 years of overall experience in the pharmaceutical industry/Biotech/CRO
- General knowledge of GCP guidelines, applicable regulatory requirements, medical terminology and data management
- Documented data management skills incl. the ability to facilitate a fast data base lock by continuous data cleaning
- Excellent verbal and written communication skills
- Expertise within key metrics and acceptance criteria for use with risk-based approaches
- Knowledge of CDISC standards and mapping SDTM and ADaM is beneficial

In addition to the above-mentioned qualifications, the candidate is a dedicated and collaborative team player that possesses excellent planning skills and has professional proficiency in English and understands 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
- The opportunity for you to shape the eClinical department for future success

As KLIFO is a smaller provider in drug development 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 Christoffer von Sehested, Director eClinical at telephone +45 44 222 916

Applications should be sent to:

job@klifo.com, marked Data Manager

Deadline:

17th September, 2018

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