KLIFO is looking for a Biostatistician or Sr. Biostatistician for our Munich office

For our German office (KLIFO GmbH) we now want to appoint a skilled and motivated (Senior) Biostatistician. The people we want to engage like to work in a consulting environment and have a positive, proactive, self-driven personality. We can offer a highly flexible, dynamic, free and trustful working environment. With exciting customers and projects in cooperation with competent colleagues. Where your knowledge, experience and your contribution are appreciated and highly valued.

The position as Biostatistician:

The Biostatistician is responsible for biostatistical deliverables on projects conducted by KLIFO (and can function as a mentor for other employees):

- Provide statistical consultancy on general or trial related aspects, internally and externally
- Cooperate closely with study team members such as data managers, programmers and project managers
- Statistical planning of clinical trials: provide statistical input to trial designs and perform sample size calculations
- Writing or peer review of the statistical sections of clinical trial protocols Ensure appropriate randomisation procedures are applied
- Writing or peer review of statistical analysis plans Coordinate small teams of programmers, perform quality control of programs
- Perform statistical analyses using SAS® (or other appropriate software)
- Provide statistical contributions to clinical trial reports Provide statistical support for publications
- Interpretation and visualization of trial data
- Provide input to Risk Based Quality Management (RBQM) including risk assessment and risk controls
- Support maintenance and further development of internal standards and of Standard Operating Procedures
- Ad-hoc tasks within the department
As a Senior Biostatistician you would in addition:

- Act as the primary contact person for clients concerning statistical aspects
- Act as mentor for other biostatisticians
- Work with no or only minimal supervision

**Qualifications of the Biostatistician**

- Diploma or master in (bio-)statistics, mathematics or comparable degree
- At least 1-4 years professional experience with planning, analysis and reporting of clinical trials in the pharmaceutical or biotech industry, in a CRO or in an academic environment.
- Knowledge of relevant regulatory guidelines
- Profound knowledge of SAS®
- Good knowledge of Microsoft® Office
- Good verbal and written communication skills
- Fluent in English (spoken and written)
- Proven knowledge of tasks and deliverables within biostatistics

The ideal candidate is a dedicated and collaborative team player.

**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)
- Use – and elaborate – your competences and experience
- A team of experienced colleagues
- Work in an interactive, flexible and positive working environment

**Location:**

KLIFO has offices in Glostrup, Denmark, Munich, Germany, Lund, Sweden and Eindhoven, The Netherlands. This position is located at our office in Munich or remote in Germany.
Contact:

For more information, please contact Simone Ahrens-Mende, Director Data Management & Biostatistics, KLIFO GmbH at +49 1522 6355866

Applications should be sent to:

job@klifo.com, marked ‘Biostatistician Munich Office’.

KLIFO processes your application and all related personal data exclusively for the specific hiring process. Your data is processed as confidential information, cf. the current data protection law (GDPR).

KLIFO is an established and integrated drug development consultancy with offices in Denmark, Germany, Sweden and The Netherlands. We provide end-to-end expert capabilities, enabling our partners to maximise opportunity, mitigate risks, drive innovation and achieve efficient project advancement.

Further information about KLIFO can be found at www.klifo.com