QUALITY ASSURANCE SOLUTIONS

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Senior QA Specialist

KLIFO is looking for a QA Officer for our Glostrup office

Would you like to use your QA experience as **QA Officer** and **QP Delegate** to support our highly skilled Clinical Trial Supply (CTS) department with a variety of quality tasks? And would you like to work in a client-focused company with dedicated colleagues able to support all parts of the drug and device development process?

At KLIFO, we're looking for a new colleague to become part of our highly experienced CTS QA Team reporting to Anders Ravn Sørensen, Team Manager, Senior QA Specialist.

The position:
As **QA Officer** your work will include review and approval of master documents, receipt of goods, batch documentation review, facilitate and approve deviation and complaint handling and SOP review and writing.
As **QP delegate** the tasks will include release of products packed for clinical trials.

Qualifications:
Preferable you are a pharmacist or have a similar education that enables you to become a QP Delegate.

The ideal candidate has minimum 3 years experience in a pharmaceutical QA or at a manufacturing/packaging site.

You must be fluent in Danish as documentation and communication is in Danish.

We offer:
As our new **QA Officer** you will:

- Experience a GXP compliant quality system and production
- Have variation in the job
- Meet a diverse client pool (pharmaceutical companies, established biotech, inexperienced, virtual biotech, investigators/academia)
- Get the opportunity to utilize many pharmaceutical competencies
- Experience an interactive, open and positive working environment
- Join a team of experienced colleagues where you use, and elaborate your skills and competencies
- Work in an interactive, flexible and positive working environment with a high level of trust, transparency and cooperation across.

Location:
KLIFO has offices in both Denmark, Germany, Sweden and the Netherlands. This position is located at our office in Glostrup, Denmark.

Contact:
For more information, please contact Anders Ravn Sørensen, Team Manager, +45 44 778 720 or Anne Ploug Jørgensen, Senior Director QA, +45 44 222 982.

Applications should be sent to:
job@klifo.com marked **QA Officer**.
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).

Deadline:
Please apply as soon as possible. Recruitment interviews will be held on an ongoing basis

KLIFO is a leading integrated drug development consultancy with significant experience in partnering with biotech and pharmaceutical companies to advance their drug development projects. We provide end-to-end expert capabilities, enabling our partners to maximise opportunity, mitigate risks, drive innovation and achieve efficient project advancement. KLIFO has offices in Denmark, Sweden, Germany and the Netherlands and employs more than 150 highly skilled employees.

Join KLIFO and work together with some of the most experienced colleagues in the life-science industry. We have strengthened our competences over many years of dedication to our field of expertise, and we support and learn from each other. This gives us a positive, flexible and informal working environment and allows us to grow as individuals and as a company.
For more information, visit KLIFO.com

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KLIFO
An Integrated Drug Development Consultancy