

KLIFO is looking for a Senior QA Specialist for our Glostrup office

Would you like to use your solid QA experience as **Senior QA Specialist** and support our highly skilled manufacturing department as well as our increasing number of international biotech and pharma clients with a variety of quality support tasks? And would you like to work for a small and very client-focused company?

At KLIFO, we're looking for a new colleague to become part of our highly experienced Quality Assurance Team reporting to Anne Ploug Jørgensen, Senior Director Quality Assurance.

The position:

As Senior QA Specialist you will be working with a diversity of jobs from traditional QA tasks for our local production and writing of SOPs to QA counselling of small biotech companies. You must expect to be working sometimes from a client site for a period either full time or some days a week. Audits will also be a part of the jobs and therefore some traveling must be expected.

Qualifications:

Preferable you are a pharmacist or have gained knowledge to become a QP/QP Delegate. You are experienced in aseptic preparation and able to release biological products. A certain knowledge of Advanced Therapy Medicinal Products is an asset.

You must have solid experience with QA work and QMS maintenance and not be afraid to write SOPs, review and release batch documentation, support and approve in validation, perform audits and handle quality incidents and CAPAs.

We offer:

As our new senior QA Specialist you will:

- Work with a diverse client pool (pharmaceutical companies, established biotech, inexperienced, virtual biotech, investigators/ academia)
- Get the opportunity to bring all your pharmaceutical competences into play in a position with many different QA tasks
- Join an open, flexible and positive working environment with a high degree of influence and transparency
- Collaborate with a team of dedicated and experienced colleagues

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 Anne Ploug Jørgensen, Senior Director QA, +45 44 222 982.

Applications should be sent to:

job@klifo.com marked **Senior QA Specialist**. 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