Senior QA Specialists (GMP or GCP)

Join an international drug development consultancy and contribute with strategic advice and operational support to clients based in Europe, the US and Asia.

At KLIFO, you'll be part of a highly specialised team of experts who value workplace flexibility and collaborate to help our life-science clients across different therapeutic areas realise their unique projects.

Become part of an experienced and dedicated team

KLIFO Quality Assurance Solutions (QAS) helps manage and ensure the correct level of quality and integrity across every stage of drug development projects and clinical research. KLIFO provides you with a unique combination of interesting and challenging tasks related to our own KLIFO GCP/GMP site in Glostrup and QA specialist tasks for pharmaceutical clients.

In QAS, we are a team of 9 highly specialised experts. We take pride in tailoring our solutions to suit each individual client. We provide practical consulting and operational services related to GMP, GDP, GCP, GLP and GVP to ensure client projects can fulfil the complex regulatory requirements.

In the role as Senior QA Specialist, you'll contribute by:

- Supporting small pharmaceutical companies/clients with QA support and counselling in developing exciting new products. This may involve working from the client's site at intervals
- Developing and designing Quality Management Systems for small upcoming pharmaceutical companies
- Acting as QA specialist for our local KLIFO GCP/GMP areas
- Auditing of a wide range of companies, including KLIFO and clients' suppliers/manufacturers/vendors. Some travelling must be expected
- In periods, working from a client site some days a week

Location
Glostrup, Denmark

Employment
Full time

Deadline for application
9 November 2023

Contact
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About KLIFO
- 150+ employees
- Offices in Denmark, Sweden, Germany and The Netherlands
- Headquarter in Glostrup, Denmark

KLIFO is a leading North-European drug development consultancy with significant experience in partnering with biotech and pharmaceutical companies.

We provide end to end solutions and support our clients with strategic advice and operational support across therapeutic areas and disciplines.

Your background and qualifications

You possess a collaborative mindset and work in a structured way with an eye for detail. You enjoy working independently and conscientiously, and you are not afraid to take responsibility. You are eager to share your knowledge and thrive in a client focused and supportive role. You are diplomatic and accommodating, and you can motivate and communicate your points of view.

If you apply as a GMP specialist experience in aseptic preparation and ability to release biological products is an advantage.

As a GMP specialist you also have:

- the education qualifying you to become QP/QP Delegate
- experience with pharmaceutical manufacturing

If you apply as GCP specialist solid experience with auditing of clinical trials is required.

Furthermore, all applicants must:

- Possess solid experience with QA work and QMS maintenance within a pharmaceutical company
- Be fluent in English and Danish, both written and spoken

Why join KLIFO?

- Join an organisation where we value people and their expertise as the greatest asset
- Enter a flexible workplace with a culture based on trust, transparency and respect
- Work with some of the most experienced and dedicated colleagues in the life-science industry
- Contribute with your expertise across different therapeutic areas
- Develop tailor-made solutions based on cross-disciplinary collaboration
- Cultivate successful relationships with our clients
- Be part of an organisation that sees knowledge-sharing as the road to success

Share your application

Share your application with us at job@klifo.com marked Senior QA Specialist no later than 9 November 2023. Kindly state how you heard about this position. Interviews will be carried out continuously.

If you have questions about the position, please reach out to Anne Ploug Jørgensen, Senior Director QA at anne.ploug@klifo.com or +45 44 222 982 for more information.

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).