Senior Pharmacovigilance Specialist

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

At KLIFO we meet the increasingly complex regulatory requirements for drug safety to ensure optimal compliance with international and national regulations and guidelines. Our pharmacovigilance experts offers our clients flexible and effective solutions from early clinical development to marketed product support.

As our new Senior Pharmacovigilance Specialist you will be part of our Clinical Operations Solutions (COS) team ensuring operational and strategic advice and support to our clients in the Life Science Industry.

Our strong team of experienced consultants support our clients from our offices in Denmark, Sweden and Germany or from a client site if needed.

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

- Support clients with strategic advice and liaison with competent authorities
- Prepare or review pharmacovigilance or safety documents (For example safety management plans and periodic reports)
- Perform monitoring of safety profile and risk-benefit balance of products
- Leading or participating in safety oversight committees

Furthermore you are responsible for all aspects of pharmacovigilance in your own projects including management, planning, PV set-up, execution, monitoring and documentation.

Location
Glostrup, Denmark

Employment
Full time

Deadline for application
29 August 2024

Contact
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Senior Clinical Research Director
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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

Besides a positive, proactive and self-driven mindset, you are keen to work in an advisory role translating client's needs into solid deliveries. You thrive in a workplace with a great deal of flexibility, experienced colleagues and exiting projects to help our clients advance their unique projects.

Furthermore, you have:

- MSc/BSc in the life science field and a minimum of 8-10 years of experience in a similar position in the pharmaceutical industry/biotech/CRO
- Experience within Pharmacovigilance from both clinical and post marketing phases
- In-depth knowledge of international PV guidelines and regulatory requirements
- Experience with safety databases
- Excellent communication skills in English both verbally and in writing.

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 Pharmacovigilance Specialist no later than 29 August 2024. 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 Randi Rahbæk, Senior Clinical Research Director, at Randi.rahbaek@klifo.com or +45 44 778 704 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).

For information on KLIFO's processing of personal information see https://klifo.com/disclaimer-privacy-policy/.