PHARMACO-VIGILANCE SOLUTIONS

PHARMACOVIGILANCE SOLUTIONS
Senior PV Specialist

KLIFO is looking for a talented and committed Senior Pharmacovigilance Specialist based in Denmark

We are looking for an additional Pharmacovigilance Specialist to join a dynamic and experienced team within Pharmacovigilance Solutions.

The people we want to engage like to work in an advisory role and have a positive, proactive, self-driven and self-confident personality. We offer a highly flexible, free and trustful working climate with competent colleagues, exciting customers and challenging projects.

As our Senior PV Specialist:
You are responsible for all aspects of pharmacovigilance in your own projects including management, planning, PV set-up, execution, monitoring and documentation.

You will
- Create trial specific Safety Management and SAE flow
- Handle the safety database (not necessarily the technical part)
- Ensure expedited reporting to Competent Authorities and Ethic Committees/Independent Review Board
- Review and present safety data listings (coding, signals)
- Write and submit periodic reports
- Define Reference Safety Information
- Review documents for safety information
- Lead or participate in safety oversight committees

Your qualifications as Senior PV Specialist:
- MSc/BSc in the life sciences 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
- Knowledge with device pharmacovigilance is an advantage
- Experience with safety databases and knowledge of MedDRA and ATC coding
- Excellent communication skills in English both written and verbal
- Master Microsoft Office
- Able to translate client's needs into solid deliveries

We offer:
- Work within different therapeutic areas and with tasks of varying complexity
- A heterogeneous client pool (pharmaceutical companies, established biotech, newly started biotech and academia)
- Build international client relations
- A unique opportunity to elaborate your competences and experience
- An interactive, flexible, trustful and positive working environment

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 Randi Rahbæk, Senior Clinical Research Director, at +45 4477 8704.

Applications should be sent to:
job@klifo.com marked Senior PV 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: 5 August 2022. Applications will be reviewed 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