PHARMACO-VIGILANCE SOLUTIONS

PHARMACOVIGILANCE SOLUTIONS Pharmacovigilance Associate

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

KLIFO is expanding our office in Denmark and wants to engage an additional Pharmacovigilance Associate into a dynamic and experienced team within Pharmacovigilance Solutions. The person we want to engage like to work in an a consulting environment and has a positive, proactive, flexible, self-driven and self-confident personality. We can offer a highly flexible, free and trustful working climate with exciting customers and projects among competent colleagues where your knowledge, experience and contribution are valuable and highly appreciated.

The position as PV Associate:
The Pharmacovigilance Associate is responsible for collaboration with clients and execution of projects according to the clients expectations:

- Single Adverse Event case handling including data entry, evaluation, assessment and writing of safety narratives
- Monitoring of the pharmacovigilance mailbox
- Handling of safety databases
- Preparation of pharmacovigilance and safety documents; for example Safety Management Plans, PSURs, DSURs, and safety signal reports
- Drive medical monitoring activities
- Maintain pharmacovigilance system master files
- Author and review SOPs

The qualifications of the PV Associate:

- MSc/BSc in the life sciences field, nurse or holding a diploma within Pharmacovigilance
- Preferably > 3years of experience from a similar position in the pharmaceutical industry/Biotech/CRO
- Knowledge of drug safety guidelines, terminology and processes
- Experience with safety databases
- Excellent communication skills in English both written and verbal
- Experienced user of MS Office and good understanding of databases
- Able to translate client's needs into solid deliveries

We offer:

- Work within different therapeutic areas and with tasks of varying complexity
- Work with a heterogeneous client pool (pharmaceutical companies, established biotech, newly started biotech and academia)
- Build international client relations
- Use and elaborate your competences and experience
- Work in an interactive, flexible and positive working environment

Location:
KLIFO has offices in 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 Pharmacovigilance Associate. 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: 27 October 2021 (although applications are 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

KLIFO A/S
Smedeland 36
DK-2600 Glostrup
Denmark
+45 44 222 900

KLIFO GmbH
Heimeranstr. 35
D-80339 München
Deutschland
+49 (0) 898 952 860

KLIFO AB
Medicon Village
Scheeletorget 1
223 81 Lund, Sweden
+46 709 135 125

KLIFO
An Integrated Drug Development Consultancy