Job opportunity

Pharmacovigilance (PV) Consultant

ZWIERS Regulatory Consultancy

Job description

We offer a job as a **Pharmacovigilance (PV) Consultant**, in which you will:

- work in an international environment of pharmaceutical and medical device industry
- ensure compliance with safety-related marketing authorisation commitments and obligations
- take on the role of QPPV
- write and review pharmacovigilance-related documents (eg. PSUR, RMP, SOPs, PSMF)
- monitor and interpret safety findings from literature
- manage and maintain pharmacovigilance databases
- provide input on safety for clinical development

Profile

You have:

- preferably, a PhD in Pharmacy, Medical Biology, Life Sciences
- minimal 5 years of experience in the pharmaceutical industry and PV
- knowledge of pharmacovigilance, pharmacovigilance systems and GVP modules
- the ability to handle deadlines and changing priorities in a dynamic environment
- the ability to communicate effectively based on scientific knowledge
- The will to live up to the Zwiers “professional approach with a personal touch” spirit

Offer

Zwiers Regulatory Consultancy offers a fulltime job, but is willing to discuss a part time contract
Salary will depend on education, knowledge and experience

Contact

For further information you can contact Alex Zwiers, CEO, at tel. +31 (0) 6 344 848 51.
We look forward to your application. You can apply via email: info@az-regulatory.com.