Job opportunity

Medical writer, consultant

ZWIERS
Regulatory Consultancy
A ProductLifeGroup Company

Job description

We offer a job as a **Medical Writer**, in which you will:

- optimally combine your passion for writing with your Life Sciences background
- work in an international environment of pharmaceutical industry
- develop and manage the drafting of medical documentation, such as clinical study protocols (CSP) and reports (CSR), clinical trial and investigational new applications (CTA/IND)
- write Common Technical Document (CTD) modules for Marketing authorization applications (MAAs) and New Drug Applications (NDA)
- write manuscripts for publication
- Initiate/manage development of templates, guidelines and SOPs for medical documentation and workflow procedures
- perform literature searches/reviews
- be coached by Zwiers and benefit from being part of ProductLife Group

Profile

You have:

- an MSc/PhD in Pharmacy, Biomedical Sciences, Life Sciences
- scientific and technical writing and editing skills (English)
- an eye for detail
- the ability to handle stringent deadlines
- the eagerness to learn, develop and excel
- the ability to communicate in a convincing way
- at least 3 years of experience as a medical writer or comparable writing position.

Offer

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.