Medical Writer
Full-time, Remote in either Ireland, UK or Germany
Ref: V197

Summary

Veryan people are the best at what we do. We share a commitment, a passion and a vision for the contribution Veryan makes to innovation in interventional medicine.

Veryan people power Veryan's business. Whether they are design engineers, quality engineers, clinical study managers or sales specialists, our people bring an unmatched expertise in each of their fields that both inspires and challenges.

Veryan's culture of innovation helps each individual develop both professionally and personally, and our culture of respect means that we all work in a collaborative, supportive and exciting environment.

Reporting to the Head of Global Clinical & Regulatory Affairs, the position of Medical Writer will develop and review clinical and regulatory affairs documents, conduct literature search, support publication, presentation and abstract writing, ensuring compliance with international regulatory standards and guidelines.

Primary Job requirements

Principal Responsibilities

- Responsible for providing advice in the development and implementation of best practice in medical writing.
- Author and review clinical and regulatory documents such as Clinical Study Protocol, Clinical Study Report, Investigator Brochure, Clinical Evaluation Plan, Clinical Evaluation Report, Summary of Safety and Clinical Performance, Periodic Safety Update Reports, Clinical Complication Analysis and ensure compliance with regulatory and clinical standards and guidelines.
- Support authoring and review of peer reviewed publications and abstracts for submission to major congresses in close cooperation with Clinical Marketing and Clinical Investigators.
- Support creation of presentation material for scientific congresses in close cooperation with Clinical Marketing and presenters.
- Review and author regular safety update reports as required by competent authorities.
- Ensure that clinical and regulatory documents are accurate, suited for the intended audience and easy to read.
- Conduct regular literature searches.
- Trigger annual reviews and updates of documents, where required.
- Work within the quality system and ensure that the quality policy and company systems and procedures are complied with in line with the Medical Device

Directive 93/42/EEC, Medical Device Directive 2007/47/EC, MDR 2017/745, and FDA Quality System Regulations 21 CFR Part 820 and ISO 14971.

Education & Training

BSc, MSc or University Degree in life science or other relevant scientific discipline required.

Experience

- Previous experience working as a medical writer in the medical device industry; vascular intervention experience is advantageous.
- Excellent spoken and written English.
- Well developed and proven medical writing skills, combined with an ability to interpret and present complex scientific data from different disciplines in a clear, concise format.