Poland Jobs Expertini®

Principal Medical Writer, sponsor-dedicated, EMEA

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Company: Fortrea

Location: Poland

Category: arts-design-entertainment-sports-and-media

As a leading global contract research organization (CRO) with a passion for scientific rigor and decades of clinical development experience, Fortrea provides pharmaceutical, biotechnology, and medical device customers a wide range of clinical development, patient access and technology solutions across more than 20 therapeutic areas. With over 19,000 staff conducting operations in more than 90 countries, Fortrea is transforming drug and device development for partners and patients across the globe.

We are on a mission to deliver solutions that bring life-changing treatments to patients faster. But we can´t do it alone.

We are seeking an experienced Medical Writer to **lead authoring and development of high complexity clinical regulatory documents** that are critical to the project(s) and strategic goals of our partner.

This is why we need you.

Join Fortrea. Your job matters.

WHAT YOU WILL DO

This is a full-time, remote job, EMEA.

You will work for one sponsor, imbedded on their team. Under this framework, you will act as an:

Expert Contributor.

You will serve as an expert contributor on our partner’s project teams. You will use your expertise and experience to write, advise, and coordinate development of complex clinical regulatory documents including Clinical Study Protocols, Clinical Study Reports, Investigator

Brochures, and clinical summary and overview documents in CTD/eCTD format for regulatory submissions world-wide.

Strategist .

This job is tailor-made for Medical Writers passionate about leading development of key documents that inform and align with project strategy.

Project Manager .

You will be the nexus where multiple teams´ expertise converge. You will leverage your experience to manage writing projects, coordinate and collaborate with stakeholders, steer discussion, drive consensus and facilitate decision-making to propel the document development cycle forward.

YOU NEED TO BRING…

Advanced degree (PhD or Masters)

Minimum 6 years medical writing experience, including 3 years as **medical writing project lead** .

Experience and proficiency in **writing and leading development** of a variety of clinical regulatory medical writing deliverables, including extensive experience leading content development of **clinical summaries for eCTDs and drug applications** across different regions.

Proven experience leading stakeholders/project teams through **submission document development** .

To drive development of documents of this scale forward, this role requires visibility, proactivity, collaboration/teamwork, and excellent communication skills:

You must be comfortable leading team discussions, managing complex medical writing tasks and processes, engaging with a variety of stakeholders, and confidently contributing your expertise based on experience.

Integration within the partner´s team is crucial, requiring readiness to assume augmented responsibilities and adaptability across environments and therapeutic areas.

A UNIQUELY DIVERSE CAREER

At Fortrea, your career path is yours to shape. We empower our team to steer their own development.

If you thrive in medical writing and want to remain deeply involved in science with high-profile clients, Fortrea is your destination.

If you are coming to a point where you want to try management, we offer comprehensive training and support to prepare you for leadership roles.

Your aspirations drive your journey with us.
Fortrea is actively seeking motivated problem-solvers and creative thinkers who share our passion for overcoming barriers in clinical trials. Our unwavering commitment is to revolutionize the development process, ensuring the swift delivery of life-changing ideas and therapies to patients in need. Join our exceptional team and embrace a collaborative workspace where personal growth is nurtured, enabling you to make a meaningful global impact.

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