GREATER LYON, FRANCE
(REMOTE WORK POSSIBLE)

Clinical Project Manager

MONIA@ELEMED.EU

The company

Are you looking for your next challenge?

Are you looking for the next step in your Clinical career?

If yes, this opportunity is for you! Join a well-established, growing company designing, developing, and producing orthopaedic devices that change patients' lives.

To further support the growth of its clinical department, this company has released an exciting position for a Clinical Project Manager to work alongside their Director Clinical Affairs.

This is a great opportunity to join a close-knit team with one vision, surrounded by other senior experts in their field. You can be sure that no two days will be the same! You'll enjoy a wide variety of responsibilities covering various elements and work on the international market.

The opportunity

You will have the opportunity to join the team and shape your role as a Clinical Project Manager where you will have autonomy and directly impact the products from a clinical perspective.

As well as writing CERs, writing clinical protocols, and handling data, you will also have the opportunity to expand your knowledge into other clinical activities such as study management and study design.

With the Medical Device Regulation in full force, you will ensure that all reports and data (existing & new) are kept compliant with the relevant regulations and highlight important information in cross-functional meetings. This is also an opportunity to experience new and exciting markets as the company continues to expand globally.

The responsibilities

- Develop and review documents and reports for the clinical parts of regulatory submissions

- Coordinate cross-functional teams and liaise with CROs throughout clinical trials in order to ensure alignment and compliance with the company's goals and necessary regulations

- Prepare, organise, and interpret clinical data

- Support the development of the clinical strategy for the company for various projects as they arise

- Manage relevant timelines required by the internal teams and external bodies from a clinical perspective

- Write CERs, clinical protocols, clinical data reports, and more

Qualifications

- Scientific background
- Minimum 2 years experience in a similar position within the Medical Device industry
- In-depth knowledge of regulations and standards/guides associated with clinical investigations/assessments
- Experience in writing Clinical Evaluation Reports for Medical Devices

INTERESTED IN FURTHER CONVERSATION?

IF YOU ARE INTERESTED IN THIS EXCITING ROLE, PLEASE SEND YOUR APPLICATION DIRECTLY TO:

MONIA@ELEMED.EU

elemed