CLINICAL PROJECT MANAGER

→ MONTÉLIMAR (HYBRID UP TO 4 DAYS REMOTE WORK)

→ Contact: clarisse@elemed.eu

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THE COMPANY

Are you looking to strengthen your experience in Clinical and take the next step in your career?

We have a fantastic opportunity to share with you to join a leader of medical devices in their field. This is a company that has a real international presence, with a number of subsidiaries in different countries as well as a strong global exportation. As Clinical Project Manager, you will be based in their main site, which means being at the very heart of the decision-making and the company operations, so you will be in the ideal place to influence the business.

Working at this company, you'll enjoy working across a wide variety of products: the portfolio ranges from class I to class III. What's more? They are continually expanding their portfolio with a strong investment in the development of new products. So this role is key to help them with the new regulation deadline approaching.

This well-established company is looking for a clinical project manager to help them in this transition. You will report to the Clinical Evaluations Manager, and your main mission will be to write the clinical files to obtain the CE marking under the MDR. Working at this company, you'll enjoy working across a wide variety of products and learn from the best of the best professionals in the industry. This is not an opportunity to miss out on!

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→ YOUR RESPONSIBILITIES:

As Project Clinical Manager you will:

- Coordinate the writing of clinical evaluations of medical devices
- Manage the clinical bibliographic review applicable
- Participate in the development of clinical evaluation strategies
- Autonomously write clinical evaluation plans and reports
- Take charge of the statistical analysis
- Participate in the answers to questions asked by the notified bodies
- Participate in the risk management
- Autonomously write pre-clinical investigation plans and reports and/or post CE marking/marketing authorization
- Participate in the development of the methods and tools used in the department: instruction sheets, model documents, etc.

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YOUR QUALIFICATIONS

- Experience in participating in a CER
- Knowledge of the MDR
- English writing ability
- Able to go to the site for the first month

GET IN TOUCH

If you are interested in this exciting role, please send your application directly to **clarisse@elemed.eu** – this could be a life-changing opportunity for you!

Would you like to find out more about our open opportunities? Visit https://www.elemed.eu/vacancies/

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