Medical Writer

Remote with one day per month on site in Lyon

monia@elemed.eu

The Company

Tired of working for the same big slow-moving corporation where the key decisions are made outside of Europe? Are you looking for a role where you will never get bored?

Located near the beautiful cultural city of Lyon, is Elemed's newest client; a small medical device manufacturer. Small in terms of people, but not small in terms of presence. This is not an opportunity in a mono-product, mono-franco environment. This company has a significant product portfolio and designs, develops, manufactures and sells their products for the global market: EMEA, N.AMERICA, LATAM & APAC.

Your role:

Attached to the Quality, Regulatory and Clinical Department, you will:

- Participate in the clinical evaluation strategy
- 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 development of new products as a clinical referent
- Participate in the risk management
- Autonomously write pre-clinical investigation plans and reports and/or post CE marking / marketing authorization

Your qualifications:

- Scientific Background
- Minimum 2 years experience in a similar position within Medical Devices
- In-depth knowledge of regulations and standards/guides associated with clinical investigations/assessments
- Fluent English and French

Interested in this role?

If you are interested in this exciting role, please send your application directly to monia@elemed.eu

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