QUALITY AND REGULATORY AFFAIRS SPECIALIST

Amsterdam, The Netherlands

Contact: monia@elemed.eu

THE COMPANY

This company has been voted over and over again to be one of the best employers in Medtech! With one of the highest employee satisfaction and retention rates in the Netherlands, this company offers security, stability and long term career opportunities.

Come and join this leading medtech company. With a brand known globally, and innovative devices from Class I-III!

RESPONSIBILITIES

- CAPA process handling, tracking progress and status.
- MDR QMS processes reporting to stakeholders
- Independently working in teams regarding MDR Process and Organization Change Management
- Support the Organization regarding Quality Policy, QMS, Regulatory and Quality in general

ESSENTIAL REQUIREMENT

- A minimum of 2 years' experience in similar role in the Medical Device Industry
- In depth knowledge of EU Medical device Regulation (MDR) and experience implementing change regarding MDR
- Fluent in English

Interested to explore this further?

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

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