AUDITOR ACTIVE MEDICAL DEVICES

Remote from Italy

THE COMPANY

Calling all design, quality, testing and R&D professionals! Do you want to oversee the newest innovations in Software and Artificial Intelligence technologies?

Have you ever thought about joining a Notified Body?

This Notified Body is growing its active device team, and this is your chance to be a part of it! We are looking for professionals with experience in active medical devices (ideally stand-alone software, AI or with software embedded).

You would be joining an International company, liaising with a stimulating network of people and being able to exchange skills and tips with high-level experts in the industry.

GET IN TOUCH WITH KRISTINA AT KRISTINA@ELEMED.EU

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THE OPPORTUNITY

This is a great opportunity to be at the forefront of the newest, cutting-edge technology. If you have previously worked in the industry, want to widen your product scope instead of being limited to one company, and work from home and travel, then this could be the perfect opportunity for you!

We are offering a great chance for professional development in a Notified Body where you will have your voice heard and be more than just a number. You can expect an excellent work-life balance, working from home, with a 40/50% travelling during audits on-site.

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THE OPPORTUNITY

As an Auditor, you will assess some of the most innovative products coming to the market: from start-ups to global corporations. You will add technical competence to this Notified Body in their product review team and ensure their reputation remains unrivalled.

You will analyse and evaluate the manufacturer's technical documentation relating to active medical devices. Your primary responsibility is to aid manufacturers seeking CE marking by performing conformity assessment activities across a range of different devices. You will report directly to the Team Leader of the Active team.

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YOUR RESPONSIBILITIES

- Plan, identify milestones and oversee the entire conformity assessment project plan.

- Assess manufacturer's documentation for CE marking according to MDR 2017/745, carrying out conformity assessment.

- Provide input/recommendations to the audit team on areas of focus for QMS audits, based on your experience of reviewing the technical documentation.

- Manage compliance and regulatory activities related to the Notified Body.

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REQUIREMENTS

- Bachelor's, Master's or PhD in relevant science or engineering subject.
- 4+ years in a design, quality, testing and/or R&D role.
- Able to travel up to 50%.
- Fluent in English and Italian.
- Software/Artificial Intelligence experience is highly desirable.

INTERESTED TO EXPLORE THIS FURTHER?

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

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