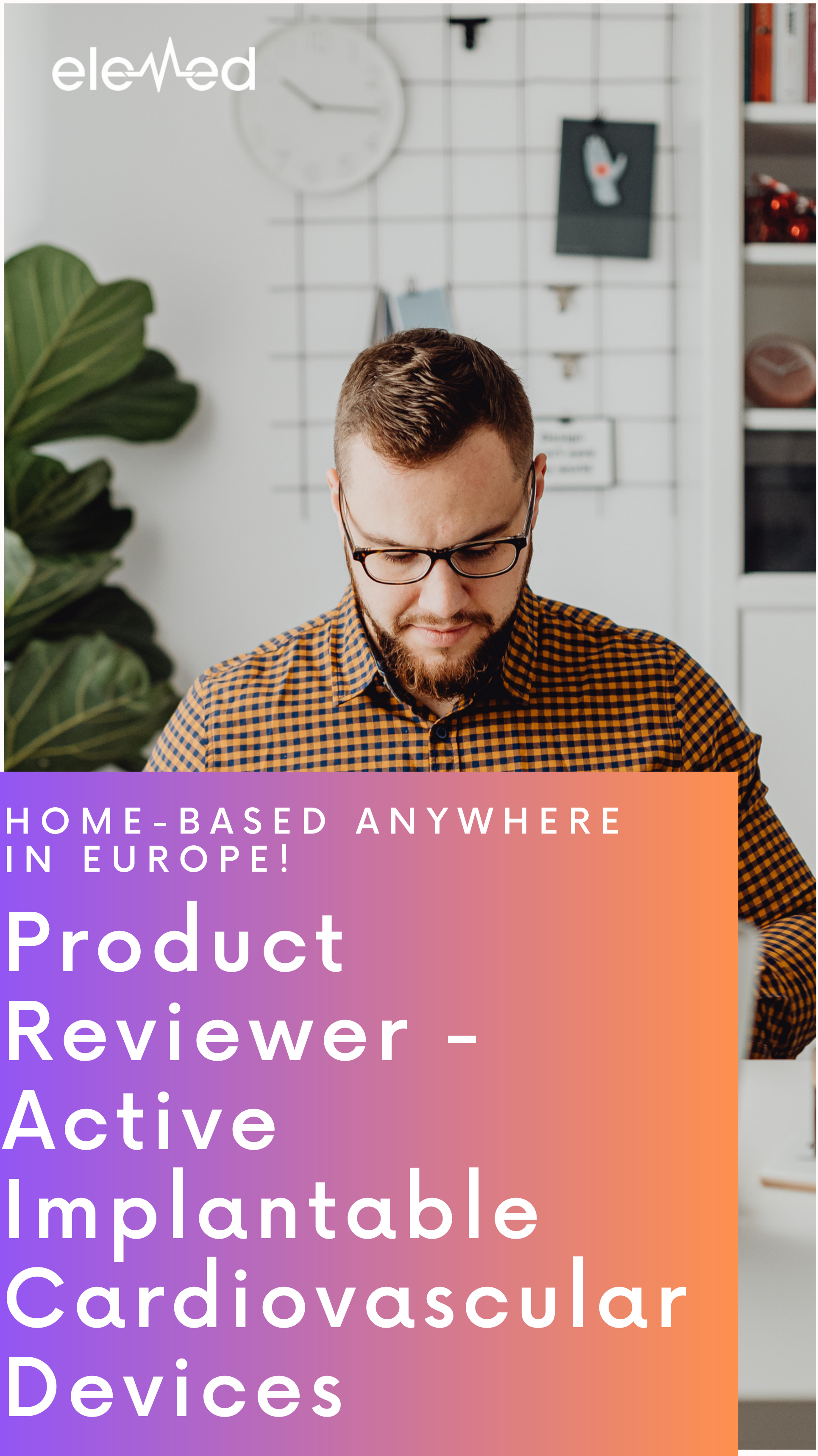

HOME-BASED ANYWHERE IN EUROPE!

Product Reviewer – Active Implantable Cardiovascular Devices

KRISTINA@ELEMED.EU

The company

Do you like working for notified bodies but are tired of the bureaucracy and slow-moving processes? Do you want to have an impact on processes in a stable environment? Are you already certified with the code to review Active Implantable Cardiovascular Devices devices? Yes..? Why not join a notified body that is trying to build efficient, fast-moving processes whilst being backed by a world-renown brand name?
We are offering a great chance for professional development in an organisation 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 and the possibility to impact the future of the business. This company is growing its Active Implantable Cardiovascular Devices team as an extension of its wider AIMD team which is driven by its goal to get the most life-changing/saving products to market as quickly as possible.

The opportunity

In this position, you will assess the most innovative Active Implantable Cardiovascular Devices products coming to the market: from startups and global corporations alike. You will be a part of a brand that is renowned worldwide and ensure its reputation remains unrivalled.

This is a chance to have the responsibility to aid manufacturers seeking a CE mark by performing conformity assessment activities and getting the life-saving Active Implantable Cardiovascular Devices to market as quickly and smoothly as possible. As well as working in these technical activities, you will have the opportunity to optimise processes internally making them more efficient and leaving a legacy within the business.

The role is uniquely structured so that there is very limited travel.

The responsibilities

As Product Reviewer - Active Implantable Cardiovascular Devices you will:

- Plan, oversee and execute the entire conformity assessment project plan from A-Z
- Assess manufacturer documentation for CE marking according to MDR 2017/745
- 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
- Optimise and improve processes internally to make them more efficient and effective

This is a great opportunity to be at the forefront of the newest, cutting-edge technology. If you have previously worked in industry in a **regulatory affairs/R&D role** with Active Implantable Cardiovascular Devices devices and want to widen your product scope instead of being limited to one company and work from home with limited travel then this could be the perfect opportunity for you.

Requirements

- Bachelor's, Master's or PhD in relevant science or engineering

- 4+ years of experience in the medical device industry

- Minimum 2 years of experience on product development projects for Active Implantable Cardiovascular Devices

- Previous experience working in a notified body

Get in touch

INTERESTED IN FURTHER CONVERSATION?

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

KRISTINA@ELEMED.EU

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