HOME-BASED ANYWHERE IN EUROPE!

Product Reviewer Active and Non-Active Implantable Neuro

WORK FOR A NOTIFIED BODY!

MONIA@ELEMED.EU

THE COMPANY

Calling all regulatory affairs professionals! Do you want to oversee the newest innovations with active and non-active implantable medical devices?

Have you ever thought about joining a notified body?

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. The role is uniquely structured, so that there is very limited travel. This notified body is growing its non-implantable/active device team, and this is your chance to be part of that! We are looking for Regulatory affairs professionals with experience in Neurology..

THE COMPANY

As a product reviewer, you will assess some of the most innovative neurovascular products coming to the market: from startups and global corporations alike. 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 medical Neurological. Your primary responsibility is to aid manufacturers seeking CE mark by performing conformity assessment activities across a range of different medical devices within Neuro devices - standalone and embedded. You will report directly to the head of the notified body.

THE RESPONSIBILITIES

As Product Reviewer / Auditor you will:

- Plan, identify milestones, and oversee the entire conformity assessment project plan
- Assess manufacturer documentation for CE marking according to MDR 2017/745, caring 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

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 role with Neuro 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.

eleMed

EXPECTATIONS

- Bachelor's, Master's, or PhD in relevant science or engineering
- 4+ years of regulatory affairs experience with Implantable Cardiovascular and/or Neuro devices

INTERESTED IN FURTHER CONVERSATION?

IF YOU ARE INTERESTED IN THIS EXCITING ROLE, PLEASE SEND YOUR APPLICATION DIRECTLY TO MONIA@ELEMED.EU