



# **Product Reviewer: Active Implantable**

**Home-based  
anywhere in Europe!**

**Contact: [monia@elemed.eu](mailto:monia@elemed.eu)**





# The company

Calling all regulatory affairs professionals! Do you want to oversee the newest innovations with 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 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 Cardiovascular and/or Neurology.





# The company

As a product reviewer, you will assess some of the most innovative cardiovascular 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 cardiovascular.

Your primary responsibility is to aid manufacturers seeking CE mark by performing conformity assessment activities across a range of different medical devices within Cardio devices- standalone and embedded. You will report directly to the head of the notified body.





# 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, 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





# Expectations

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

If you are interested in this exciting role, please send your application directly to

[monia@elemed.eu](mailto:monia@elemed.eu)

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