

## Remote in Europe

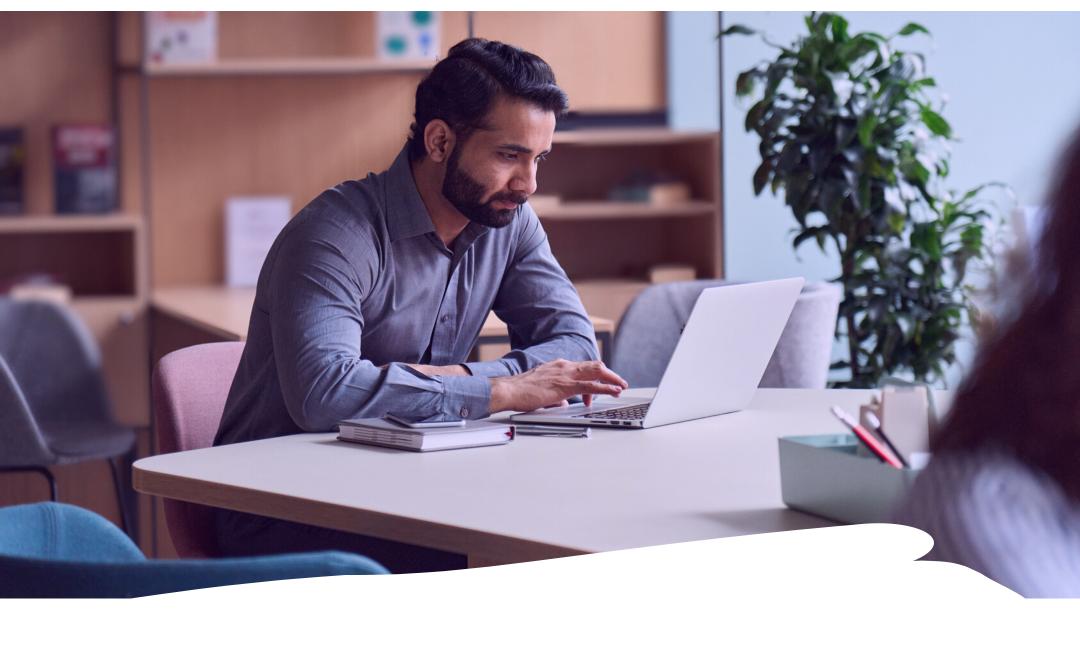
The medical device industry is currently experiencing huge challenges with changing regulations such as MDR and increasing costs, as well as exciting changes with amazing technological advancements. Would you like to have a seat at the table of how MDR is being interpreted and shape the industry? Are you an experienced leader who enjoys supporting your team's success and development? Is it time for you to challenge yourself and use your skills in a new context?



#### **THE COMPANY**

This is an opportunity for you to join a global organisation, with a strong brand and name that is helping the sector to navigate through the demanding times. This is a company with a long term vision, focussed on people, not just profits. One of the great things about a privately owned company is that decisions are made with the long term goal in mind, rather than just the next quarter. This means you'll benefit from excellent training and development as well as being part of a culture driven by ethics. You will receive some of the best training that the industry has to offer and be surrounded by some of the most highly qualified professionals, experts in their respective fields.

If you believe in creating a sustainable and safe future in the MedTech world whilst adding value to customers, this is the company for you!



#### THE OPPORTUNITY

Changes in the business to be more globally structured have created an excellent opportunity for a clinical leader (you) to step into a brand new role leading a highly successful team that has enjoyed year-on-year growth and work with professionals from a diverse background. You will be leading a team of up to 10 Clinical experts, building relationships internally and externally as well as remaining hands-on with clinical activities such as assessing clinical evaluations.

Having a seat at the table of how MDR is being interpreted means you will be challenging and using your skills in a new context all whilst being at the forefront of some of the most innovative products the industry has to offer.

#### **IN THIS ROLE YOU WILL:**

- Lead the clinical review team of up to 10 and manage them on deliverables set out by the global organisation
- Enable and develop the team to support their personal and professional development and ensure success
- Support internal training and track the teams progress
- Facilitate and/or negotiate solutions to bring closure to escalation processes if needed
- Continuously build and maintain a high level of expertise relating to present and pending regulatory requirements and standards, by attending internal and external training courses and reading scientific articles
- Take responsibility for managing multiple projects, and responding to questions from internal and external partners on standards, regulatory and technical requirements to achieve compliance with the relevant regulatory system
- Monitor and support the team on their respective projects and ensure that they meet the required deadlines
- Review and scientifically challenge the clinical data presented by the manufacturer in the clinical evaluation and any clinical investigations
- Assess the manufacturer's clinical evaluation reports and corresponding documents
- Develop strong communication skills by issuing quotes and supporting customers with relevant information and answering questions



### REQUIREMENTS

- Deep experience in Clinical Affairs for medical devices from research, hospital, industry or notified body
- Prior leadership experience managing teams directly (remote management is a plus)
- Scientific, engineering OR medical degree is a requirement for this position. Medical degree is a strong plus

# INTERESTED TO EXPLORE THIS FURTHER?

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