Head of Regulatory Affairs & Quality Assurance

Lausanne Area, Switzerland

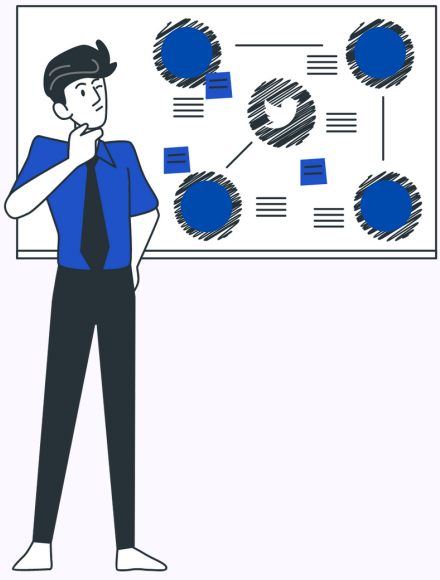
A fantastic new and exclusive role has come into Elemed, with the opportunity to be the next leader of a growing medical device company based in the Greater Lausanne region.

Elemed are searching for this innovative and internationally growing company's next Head of Regulatory Affairs and Quality Assurance, who will lead a team and develop key future strategies.

COMPANY

Be the RA/QA leader at our client's headquarters, based in a great location within the Lausanne area. You will report directly into the founders of the business providing for a fantastic opportunity to have short decision pathways and no red tape. Provide your own strategy and build the department according to your own ideas and expertise; thus allowing you to leave a legacy within a company at their exciting point of growth.

This company is developing novel devices on a global scale inclusive of the emerging markets. A great opportunity for you to provide leadership on international and developing markets.



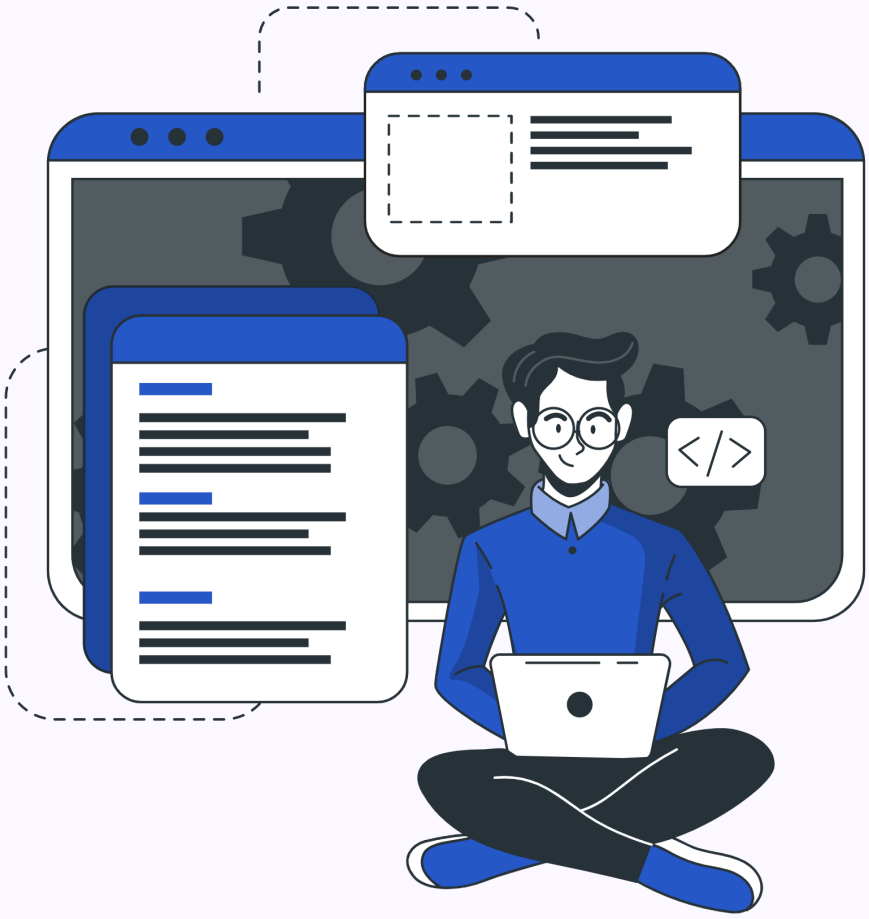
KEY RESPONSIBILITIES

As the Head of Regulatory Affairs and Quality Assurance, your responsibilities will include but not be limited to:

Management- You will be in charge of managing any employees within the RA/QA department. Also in the future hiring for your own team of experts.

Technical Documentation- You will be the process owner for all regulatory documentation activities, inclusive of CE marking and product development. You will manage and drive forward strategy for the product to be developed globally, inclusive of new market areas.

Owner of QMS- You will be in charge of all strategy and processes linked to the QMS. Inclusive of setting up the system from scratch.



THE IDEAL CANDIDATE FOR THIS ROLE WILL HAVE:

- Come from a strong regulatory and quality assurance background, with 5+ years within medical devices.
- A good working knowledge of QMS and audits. Ideally have set up a QMS system from the foundations up before.
- Previous management experience would be a bonus for this role, but not essential.
- Excellent knowledge of the international registration process, inclusive of the emerging markets like APAC.
- Expertise within MDR, MDSAP.
- Excellent communication skills in both English. French would be a bonus.
- Excellent stakeholder management exposure.

Be the next leader of a growing medical device company. Provide key strategy and expertise across both regulatory affairs and quality assurance, and leave a legacy internationally.

Interested to explore this further? Please send your CV to tamanna@elemed.eu to arrange a confidential career discussion.