GLOBAL HEAD OF REGULATORY AFFAIRS

- Multiple locations available: Baar, Switzerland; St.Gallen, Switzerland; Newbury, UK; Kiel, Germany; Valencia, Spain

- Contact: tamanna@elemed.eu

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Are you a regulatory affairs professional looking for a global challenge? Are you looking to have short reporting lines so you can make an impact? Are you looking to develop key strategy for multiple sites within RA internationally? If you answered yes to any of these questions then this is the role for you!

THE COMPANY

In this role, you will work for a mid-size medical device company, with a strong international presence in Switzerland, Germany, and China. This medical device manufacturer is the market leader in its field, enjoying strong partnerships with some of the world's most renowned MedTech companies, as well as designing and developing its own products.

Owing to a dynamic acquisition strategy, this company has enjoyed fast-paced growth and is entering the next (exciting!) phase of its expansion journey. You will join the leadership team within RA which has a great strong global presence.

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THE OPPORTUNITY

There are strong opportunities in the future to mentor and manage a team of regulatory affairs professionals. You will begin your journey as the Global Head of RA where you will lead international projects and make an impact on key strategy.

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

Management- You will develop skills to manage and mentor juniors within the team- you will play a critical role in driving the strategy for global registration processes.

MDR Project- You will play a critical role in the continual development and improvement of projects.

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

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QUALIFICATIONS

The ideal candidate for this role will have:

- Come from a strong regulatory background, with 5+ years of experience within regulatory affairs for medical devices
- Previous management or mentoring experience would be a bonus for this role, but not essential
- Excellent knowledge of the international registration process
- Expertise within MDR, MDSAP
- Excellent communication skills in English- fluency in German would be an advantage

Be the next manager of a growing medical device company. Provide key strategy and expertise across regulatory affairs, and leave a legacy internationally!

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

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

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