REGULATORY AFFAIRS SPECIALIST

St Gallen, Switzerland OR
remote anywhere in Germany

tamanna@elemed.eu

contact

elemed

THE COMPANY

Join a well established, growing company designing, developing and producing innovative medical devices that changes the lives of patients globally. The company have offices around the world allowing you to be part of a global community of experts.

This is a great opportunity to join a close knit team with one vision, surrounded by other senior experts in their field. You will work with some of the top RA/QA experts, allowing you to develop your skills and further develop your career.

You can be sure that no two days will be the same! You'll enjoy a wide variety of responsibilities covering various elements within Regulatory on an international scale.

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RESPONSIBILITIES

As the Regulatory Affairs Specialist your responsibilities will include but not be limited to:

- Develop and create key technical documentation for product submissions globally inclusive of FDA, LATAM and APAC
- Monitor the landscape of MDR and global changes
- Play a key role in monitoring and developing labelling tasks
- Support in the submission process to key notified bodies internationally

REQUIREMENTS

- 3+ years of experience within Regulatory affairs
- Expertise within the field of medical devices
- A key understanding of ISO 13485
- Previous experience within LATAM or FDA would be a bonus for this role
- Excellent communication skills in English and conversational level German

INTERESTED?

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/