Regulatory Affairs Specialist

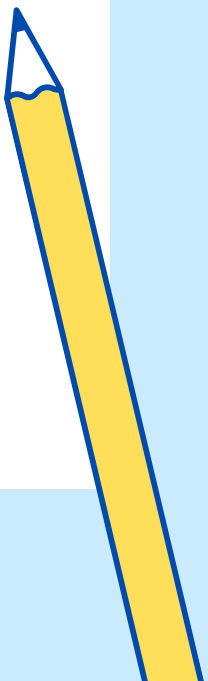
Zurich, Switzerland

THE COMPANY

Join a well established, growing company designing, developing and producing a Class III active implantable medical device that changes the lives of patients.

To further support the growth of their Regulatory and Quality division, our client have released an exciting newly created position as Regulatory Affairs Specialist.

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.



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

- Develop and create key technical documentation for product submissions.
- Monitor the landscape of MDR and global changes.
- Play a key role in monitoring and developing labeling tasks.
- Support in the submission process to key notified bodies internationally.
- Work closely in supporting the VP RA/QA with all maintenance activities for regulatory affairs.

Requirements

- 3+ years of experience within a Regulatory affairs.
- Expertise within the field of medical devices, with specific expertise within active medical devices.
- A key understanding of ISO 13485.
- Expertise in the creation and maintenance of technical files for active medical devices.
- Excellent communication skills in English.

Interested to explore this further?

Please send your CV to tamanna@elemed.eu to arrange a confidential career discussion.

Please note: Elemed has sole rights and a mandate to recruit for this role. Be vigilant of 3rd parties advertising or promoting this role as any 3rd party applications will not be accepted and will be withdrawn.

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