REGULATORY AFFAIRS MANAGER

Are you an experienced regulatory affairs medical device specialist wanting to develop your career?

Renowned for its excellent work-life balance, cosmopolitan cities and stunning natural beauty, Switzerland also hosts one of the biggest medtech hubs in Europe.

And, with its central location, travelling to anywhere in Europe only takes a couple of hours!

THE COMPANY

This is a very unique chance to join a privately owned, growing company of 2000+ people, currently going through a transformation.

In this role, you'll join a close knit team that is responsible for regulatory management of medical devices globally. The company is present in over 80 countries across EMEA, North America, LATAM and Asia.

This is a really varied role, you'll have the chance to work on new product development projects and see the whole lifecycle of a device, from concept to market.

You'll have exposure to global markets and get to learn about the regulatory environment in each of those countries.

You'll also support the transition to the new MDR!

Another great point, is that this is a Swiss company, and you'll be based in their HQ. You'll be at the centre of where all the decisions are made, meet senior leadership in the corridors, and have lots of face to face contact with your partners.

Swipe up >>>

As Regulatory affairs manager you'll have the following responsibilities

- Guide other partners as the subject matter expert for your own portfolio of medical devices, on a global scale.
- Closely collaborate in multi-disciplinary teams from Regulatory, Quality, Technical, Marketing & R&D on new product development projects and international product launches
- Drive key relationships with European Authorities, notified bodies and International competent authorities such as Canada, Japan, China and Brazil
- Maintain existing registrations globally, follow the changes to the MDR and be responsible for the full lifecycle management of your device portfolio

Requirements

- **2- 6 years experience Regulatory affairs in the medical device industry**
- **The complexity of the portfolio you'll manage will be adapted subject to your experience! There is something for everyone.**
- **English**

Contact us! hello@elemed.eu