Regulatory Affairs Manager

Basel, Switzerland

Enjoy a wide range of responsibilities on an international scale within this leading medical device company.

Join a small team of highly qualified RA managers and benefit from their expert knowledge.

Best of all, this is a unique chance to have a personal working environment in a company that is financially robust and backed by one of Europe's best medical Device manufacturers.

The opportunity:

- Lead regulatory in new product development projects for new product introductions and changes to existing devices.
- Work alongside Marketing, Quality, R&D to devise the regulatory strategy for new product launch
- Support regional and local registrations across a range of countries: USA, Canada, MEA, Asia & Europe
- Driving the process of regulatory approval for challenging medical devices

Apply now

tamanna@elemed.eu

Essential Requirements

- Minimum 3-5 years experience in Regulatory Affairs activities in the Medical device industry
- Previous experience within translating MDD files to MDR.
- Fluency in both English and German.