Senior Manager Regulatory Affairs

Solothurn, Switzerland

Come and join this leading medtech company located in Solothurn! With a brand known globally, innovative devices from Class I-III and a care for quality, they have been recognised as one of the top employers in the industry.

Due to continued growth and investment in product development we are looking for an experienced leader who enjoys training, mentoring and managing a team of regulatory professionals working with Class III devices, to step into this newly created position. The position focuses on regulatory support to product development and is a great opportunity to shape the position in your own way - (you will not step into someone else's footsteps!

This is the chance to have a leadership role in a division that has been overperforming in the last few years. They provide great future prospects and have a focus on work-life balance and having fun at work - offering great social benefits.

As Regulatory Affairs Senior Manager you will have the following responsibilities:

- Train, mentor and manage a team of regulatory specialists
- Collaborate in multi-disciplinary teams from Quality, Regulatory, Marketing, R&D and technical on worldwide product launches
- Manage global registrations and maintain existing registrations following changes to MDR
- Deliver strategies to ensure compliance in all US, EU and international markets according to the relevant requirements

Do you have?

- 5+ years experience in the medical device industry in a regulatory affairs role
- Previous experience leading a team
- CE marking & 510(k)

Contact:

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