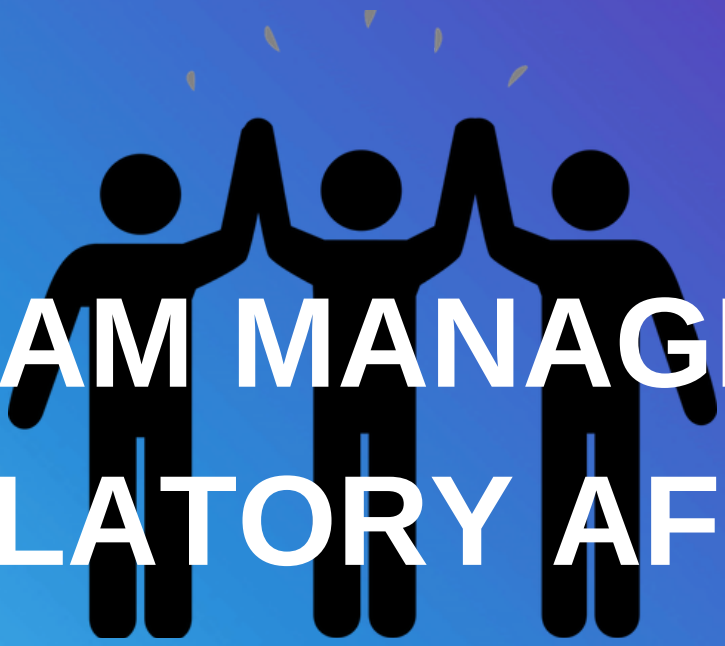

TEAM MANAGER REGULATORY AFFAIRS

ZUG, SWITZERLAND

Are you interested to use your experience in Regulatory Affairs and Medical Devices in a new context?

Want to learn something new?
Ready to take a step up in managing a team?
This is a unique opportunity to do all those things.

THE COMPANY

Join a privately owned, growing company of 2000+ people. In this role, you'll be responsible for managing a young, motivated and international team of 3 regulatory specialists.

This team manages a really varied portfolio: medical devices, consumer products, and more! So you'll have the chance to learn a about a variety of different standards including (electrical safety, electro magnetic compatibility, environmental packaging testing, sterilization, biocompatibility, food contact and more.)

The company is present in over 80 countries, and all the major markets, including: EU, USA, Canada, Brazil, Australia, China, Japan and more! So from a regulatory perspective you'll have exposure not only to Europe, but also to the rest of the world!

We are looking for either an experienced specialist, looking to take a first step into management, or someone experienced that enjoys management, coaching, mentoring and supporting a young team.

hello@elemed.eu

COMPANY

As Regulatory affairs manager you'll have the following responsibilities

- Coach, guide, train and mentor a team of regulatory specialists
- Provide regulatory technical and strategic support to innovation projects
- Build regulatory awareness amongst other departments
- Management of worldwide country registrations, labelling & marketing reviews/releases and scientific lobbying
- Closely collaborate in multi-disciplinary teams from Regulatory, Quality, Technical, Marketing & R&D on new product development projects and international product launches for medical and non medical devices
- Maintain existing registrations globally, follow the changes to the MDR and be responsible for the full lifecycle management of your varied portfolio of products.
- Drive regulatory strategy for product approval and maintenance for your team's portfolio of products globally.

REQUIREMENTS

- 6+ years experience Regulatory affairs in the medical device or consumer products industry
- Intermediate German

Get in touch!

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