

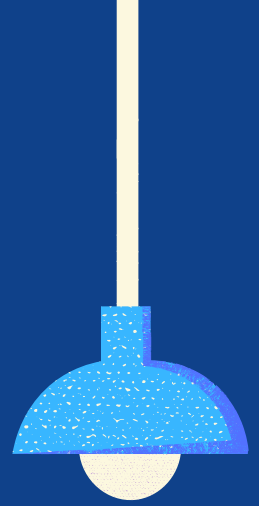
Baden-Wurttemberg, Germany

REGULATORY AFFAIRS MANAGER

Get in touch with kristina@elemed.eu



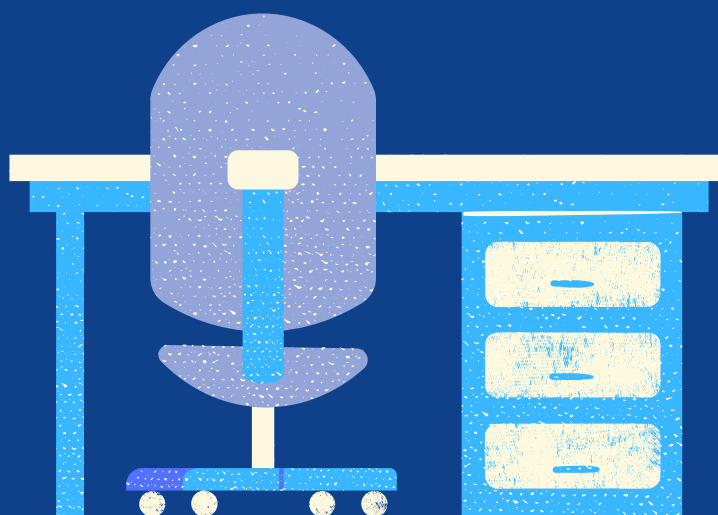
Are you looking for a new challenge that will allow you to have autonomy, variety and a visible progression pathway? Look no further as we have just the role for you!



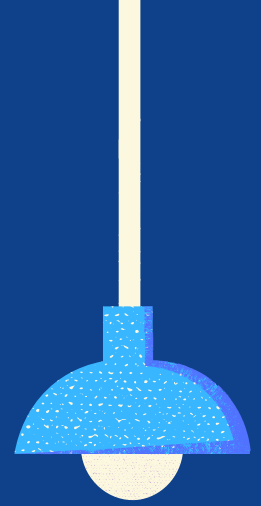
THE COMPANY

This family owned company is investing heavily into new product development to increase their ever expanding portfolio of 15,000 products. With their eye on the future, they are developing digital solutions to compliment and work alongside their current devices. They are looking for YOU to join their regulatory department and help change the future of medical technology.

This is a fantastic opportunity to work in a company that has safety and patient health as a core value and where you will be regarded as more than just a number. With exposure to international environments and a vast portfolio, you can ensure your personal and professional development.



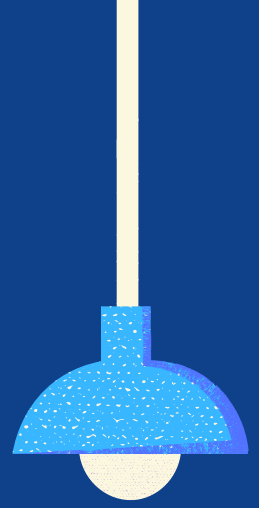
THE RESPONSIBILITIES



As Regulatory Affairs Manager you will:

- Oversee and manage the entire lifecycle of your portfolio of products from A - Z
- Guide and consult teams throughout product development projects and product lifecycle on standards and questions related to risk management/product safety
- Monitor and analyse worldwide registrations, trends and hot topics that may impact the company in terms of products and processes across all portfolios
- Collaborate in cross-functional teams on new projects; marketing, R&D, quality, regulatory, technical and provide support as the subject matter expert for the portfolio of medical devices
- Establish and maintain all relevant laws, regulations, guidelines, directives for all products in a system to support product compliance
- Support the transition from MDD to MDR
- Deal with exciting products including electrical and software devices





THE QUALIFICATIONS

We are looking for:

- Minimum bachelor's degree in science/engineering or equivalent qualification
- Minimum 3 years experience in Regulatory Affairs OR product management in the medical device industry
- Fluent English AND German (written and spoken)

ARE YOU INTERESTED IN THIS POSITION?

Please send your application directly to
kristina@elemed.eu

