

Regulatory Affairs Specialist

Tuttlingen, Germany

Come and join a leading global manufacturer as a Regulatory Affairs Manager!

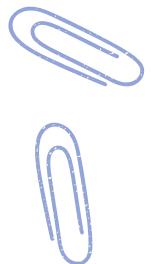
This family owned company is investing heavily into new product development to increase their ever expanding portfolio of 15,000 products. With their eye on the future, they are developing digital solutions to compliment and work alongside their current devices. They are looking for YOU to join their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

This is a fantastic opportunity to work in a company 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. We are looking for a Regulatory Affairs professional to join this team and build on their experience already gained in medical technology.

If you are the kind of person who sees beauty in complexity, challenges instead of obstacles and enjoys being hands-on this is the perfect position for you!

As Regulatory Affairs Manager you will:

- Oversee the entire lifecycle of the products from A - Z
- Collaborate in cross-functional teams on new product development projects; marketing, R&D, quality, regulatory, technical
- Support the transition from MDD to MDR
- Deal with exciting products including electrical and software devices



We are looking for:

- Bachelor's degree in science/engineering or equivalent qualification preferred
- 2+ years experience within Regulatory Affairs in a medical device context, MDR experience is a bonus
- Fluent English and German (written and spoken)
- Experience with mechanical engineering or software is a bonus



We'd love to hear from you!

**Please send your CV to paul@elemed.eu
for a confidential career discussion.**