Quality Engineer R&D (all genders)
Full-time, Munich, Germany

At TRiCares, we are dedicated to providing innovative, minimally invasive treatment solutions for patients suffering from failing tricuspid valves. As a growing MedTech company based in Munich, we are looking for a proactive and skilled Quality Engineer R&D to join our dynamic team in Munich and support us in our mission to improve patient care.

This is a unique opportunity to join a growing medical device company, where you will be part of a passionate and international team working towards transforming the landscape of cardiovascular treatment through innovative technology.

By applying your expertise, you will drive the creation of comprehensive and compliant technical documentation with focus on writing and reviewing technical design and certification related documents and reports, ensuring compliance with internal specifications and industry standards. As part of the R&D team, you will find yourself close to the design and development of both the heart valve prosthesis and corresponding catheter systems. This enables you to have an impact on product development while keeping close collaboration with the departments for regulatory affairs and quality management.

Key Responsibilities:

- Lead creation, maintenance and optimization of technical documents within R&D in compliance with MDR requirements.
- Identify and implement improvements in the format, structure and content of existing Technical Documentation.
- Actively contribute to the creation of the design history files for the TRiCares product portfolio. Support R&D to ensure accuracy, completeness and compliance of documentation to all applicable regulatory and engineering requirements.
- Collaborate closely with Project and Quality Management teams to meet timelines and regulatory requirements.
- Train and advise colleagues on all topics related to technical documentation.
- Bring together different perspectives by scheduling and preparing for meetings with cross-functional teams to consider all stakeholder expectations.

Qualification and Experience:

- Degree in technical, scientific or related field.
- Minimum of 2 years of experience - preferably within the medical device industry – in creating and maintaining technical documentation.
- Knowledge of regulatory requirements (MDR and ISO 13485) is of advantage.
- Structured, solution-oriented with strong analytical skills and a team-oriented approach.
- Excellent communication skills
- Strong attention to detail, with excellent organizational and teamwork skills.
- Proficiency in English is required; German language skills are an advantage.

What We Offer:

- A dynamic and forward-thinking work environment, shaping the future of transcatheter heart valve therapy
- The opportunity to work in a truly international team, with talented and driven colleagues from around the globe
- A strong company culture rooted in collaboration, innovation, and purpose
- Excellent opportunities for personal and professional development in a fast-growing MedTech company
- A competitive compensation and benefits package tailored to your experience and contribution
- Regular team events and offsites to connect, celebrate successes, and foster team spirit
- An inclusive and respectful workplace where diversity is not only welcomed but valued
- Flexible working hours with remote work options

Ready to make a difference?

We look forward to receiving your application. Join us on our journey to transform patient care – one heartbeat at a time.

Apply now at: HR@tricares.de.