QA Specialist

°MEQU

About °MEQU

°MEQU has developed °M Warmer, a portable fluid warmer capable of heating infusion fluids and blood. °MEQU obtained CE marking for °M Warmer in October 2018. In accidents, there is often a need to administer fluids or blood to seriously injured individuals to restore normal bodily functions. By using °M Warmer, the chance of survival is increased, as infusion fluids and blood are heated to body temperature within seconds. This is essential in military field operations, civilian rescue missions, and, of course, in hospitals, including trauma centers and during surgeries. Among the company's customers are the Norwegian air ambulance, the British military, and most recently, Region Hovedstaden. °M Warmer stands out significantly from its competitors, being small, lightweight, portable, and a robust device that is easy to use. Its user-friendly design has won several awards. °MEQU currently employs 12 dedicated employees who, in addition to optimizing the existing product, are actively involved in developing new, exciting products.

About the position

As a QA Specialist at °MEQU, you will be part of a quality team consisting of a total of 2 QA Specialists and a RA/QA Manager. The team is responsible for ensuring that all of °MEQU's work meets applicable quality requirements.
Your primary responsibilities will include:

- Updating and maintaining quality documentation such as procedures, instructions, and forms
- Handling complaints
- Managing Field Action Assessments and providing support for Initiated Field Actions in collaboration with RA
- Performing trend analyses and evaluating them
- Preparing and conducting internal and/or supplier audits and participating in audits
- Handling NCR and CAPA activities
- Continuous process optimization
- Training new colleagues in both general and specific QA processes

You will report to °MEQU's RA/QA Manager and also work closely with your colleagues in both R&D and Operations. °MEQU's organizational structure is flat, and everyone works as a team towards common goals, helping each other across departments.

Over time, the position may evolve in line with the company's growth.

Your qualifications

You likely have a higher technical, scientific, pharmaceutical, or healthcare-related education. However, the most important qualification is that you have a minimum of 2 years' experience in QA within medical devices and experience working under ISO 13485, MDR, QSR, and 510(k) compliance.
As a person, you are dedicated and thrive in a dynamic organization where everyone works together and helps each other across functions and responsibilities. Moreover, it's important that you take responsibility, communicate effectively, and are proactive, while also working in a structured and systematic manner. Additionally, you find it appealing to be part of a growing company with the broad responsibility that it entails.

How to apply

°MEQU is collaborating with DEDENROTH on this recruitment, and Mia Danielsen is responsible for the recruitment process. To apply for the position, please send your CV to md@dedenroth-consulting.com.
You are also welcome to contact Mia at the above email address or phone +45 26178142.