Quality Team Manager

Occitanie, France

clarisse@elemed.eu

The company

Do you want to be ‘wowed’ by a Quality Team Manager opportunity? Are you looking for a challenging role in a unique environment?

If yes, you could join a multicultural team of quality experts in a medical device world leader in this exciting, cutting-edge role. Sitting alongside their innovation centre, you will work with state-of-the-art medical devices, benefit from excellent training, and make a difference to the lives of millions of people across the world.

This role is also part of a new department structure in order to change the culture of the business. A real chance to make impactful changes.

This position will give you the chance to join a collaborative and knowledgeable team where you will work on products that constantly change and evolve.

The opportunity

Under the responsibility of the Site Manager and in close collaboration with the VP RA/QA you will manage a team of 5 direct reports where you will provide support for each employee in terms of daily activities, technical skills development, and personal skills and personal development!

As a Quality Team Manager you will develop new processes to develop and simplify the QMS and play an important part in helping the company thrive.

Responsibilities

- Be responsible for developing, implementing, and maintaining the company's corporate Quality Management system

- Ensuring that the Quality Management System is effective and fit for purpose to meet the company's needs and future growth plans

- Identify and report on status, performance, and opportunities for QMS improvement to the company's management team

- Be responsible for post-market surveillance, vigilance, complaint handling, and CAPA processes and support the leadership in reporting to the competent authorities and Field Corrective Actions if required

- Lead training across the company to ensure the QMS is effective and quality is adopted within the company and manage the overall QMS training and competency program for the company

- Be responsible to generate, review and edit SOPs, and other applicable documentation within the QMS responsibility

- Undertake internal and 3rd party audits of the company as well as participate in the evaluation of suppliers

- Support new product development projects from the quality perspective

- Support the regulatory team with registrations or changes where required

Requirements

- Similar experience within the Medical Devices industry
- Minimum of 5 years experience as a Manager, where you have impacted their development and training
- Fluent in French and English

Get in touch

If you are interested in this exciting role, please send your application directly to **clarisse@elemed.eu**

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