QUALITY ASSURANCE DIRECTOR

Nyon, Switzerland

THE COMPANY

Are you a real leader? Are you passionate about all Quality Systems aspects and looking for something where you will be empowered? Then, I have something for you!

My client is a top diagnostics company growing significantly year on year. It's a great chance to make your mark in a company that is growing, which means lots of opportunities for new product introduction into new markets!

This position has a strong overlap with product development!! We are looking for strong candidates that have experience leading Quality Systems activities globally and are looking for an opportunity to apply their skills in the IVD area. This is a unique chance to make the cross-over into another highly regulated environment and work with new challenges.

You will get the opportunity to work with cutting-edge products and a team of expert professionals. That will be your chance to become the right hand of the Head of RAQA, a member of the executive team!

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THE OPPORTUNITY

Your Quality department is responsible for the day-to-day management of the Quality Management System encompassing Non-Conformance, Corrective / Preventive Action, Internal / External Audit, Management of Change, and Supplier Management.

As a Head of Corporate Quality, you will be responsible for control and coordination of activities undertaken by your managed team including supplier management, statutory and regulatory requirements, and the principles of GMP.

You will also be entitled to supervise and coach all of the department at two different sites.

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YOUR RESPONSIBILITIES

- Supervision of the quality engineering function
- Ensuring compliance of the key process
- Ensuring departmental and business metrics are maintained to target.
- Identify and drive quality improvement, both departmental and company-wide
- Internal Investigations and Root cause analysis of complex problems
- Trend key business and quality critical parameters and respond proactively to these
- Perform any other duties as reasonably requested from the Company from time to time.
- Conduct all duties in compliance with Good Manufacturing Practice (GMP), Good Documentation Practice (GDP) and appropriate regulatory requirements.
- Ensure compliance and demonstrate acceptable personal and group performance against KPI for managed Quality Management System processes
- Facilitate all and, where necessary, lead investigations and root cause analysis for quality investigations to ensure compliance with internal and external regulations and specifications
- Interpret, analyse and trend data from key data sources including investigations to produce technical reports, recommendations and drive to completion to ensure improvement is achieved and that effectiveness of action can be measured and demonstrated
- Manage, coach, and develop direct reports

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YOUR QUALIFICATIONS

- Minimum of 8 years experience in Quality Management System within Medical Devices and/or IVD
- Experience in managing a team
- Fluent in French and English

Interested to explore this further?

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