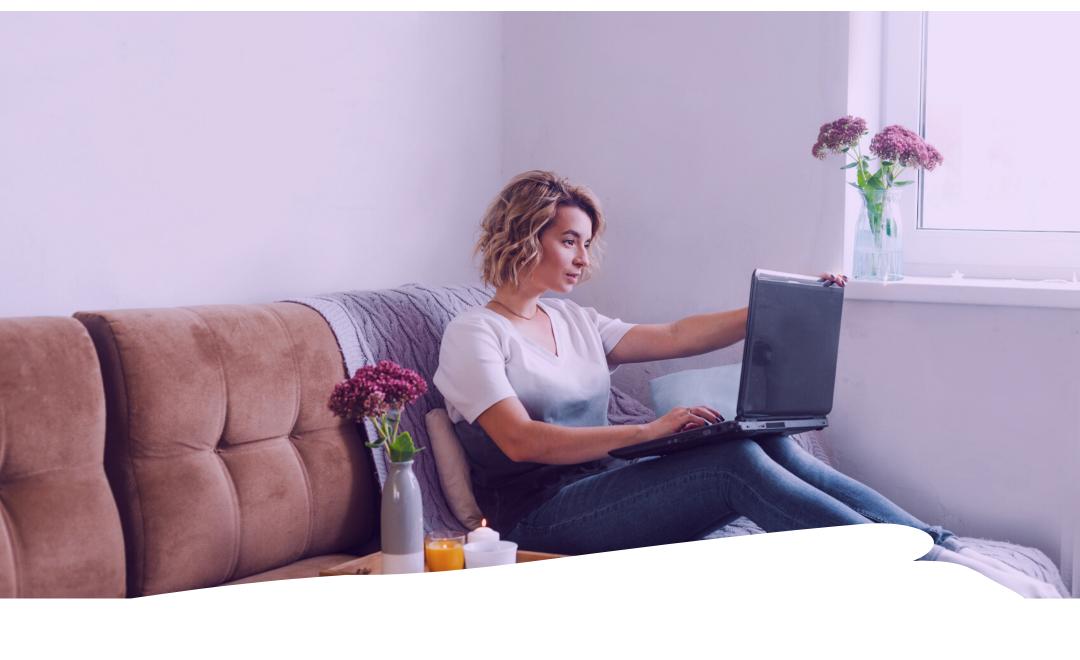


THE COMPANY

Do you want to oversee the newest innovations in IVD technology? Do you want to work for a company that will leave a legacy within the MedTech industry? We have just the opportunity for you!

This company leads the way globally for assessing whether diagnostic manufacturers meet the requirements set out in European legislation, ensuring their products are safe and effective. This is a great chance to really be at the centre of the most cutting-edge IVD technology entering the market and be surrounded by the top experts in the field.

You will get access to advanced training and learn about the new In Vitro Diagnostic Regulation, and you will be part of the biggest change affecting the Diagnostics industry in the last 50 years!

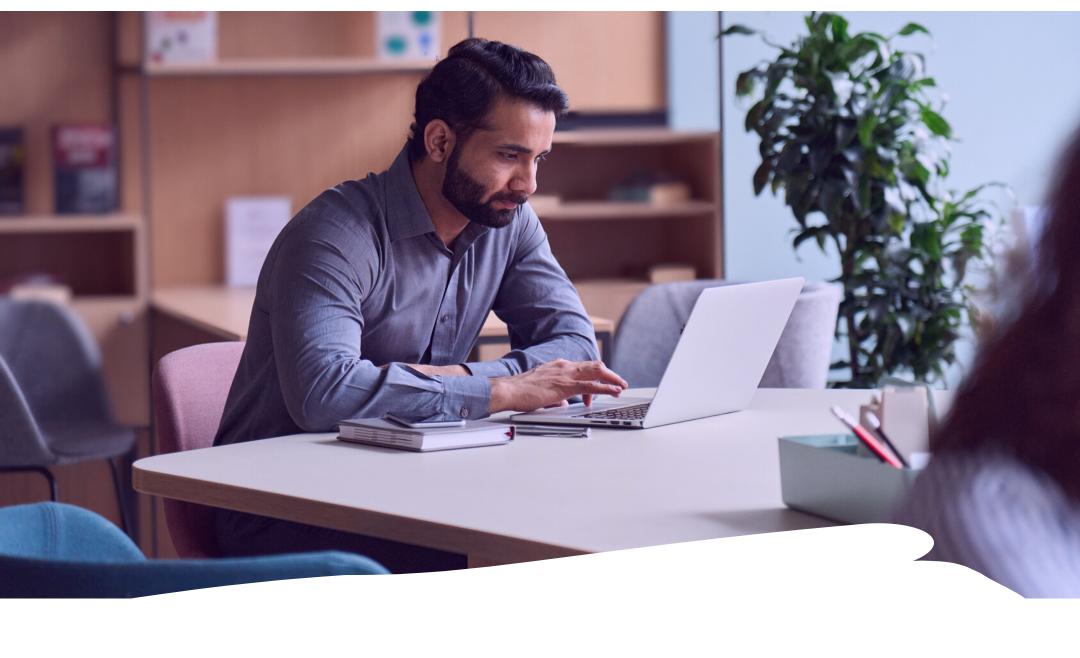


THE OPPORTUNITY

Are you looking for a dynamic role that will allow you to travel around Europe? Are you a people person and your communication skills are top-notch? This is the opportunity for you!

This is a great chance to be part of a growing and expanding company that is the leader in its field, joining as the QMS Manager. You will have a very versatile role within the business as it doesn't involve just auditing all the time - you will be responsible for handling the communications with the clients!

If you're somebody who thinks in pragmatic solutions rather than restrictions, if you are ambitious, motivated, and have the ability to apply critical thinking to challenge the norm, we want to hear from you!



YOUR RESPONSIBILITIES

- Evaluating the Quality Management Systems and technical documentation during audits and inspections at IVD manufacturers according to EU directives and quality management standards
- Evaluating technical documentation as well as product dossiers and test results of production batches in the area of in-vitro diagnostics of the highest risk class, mainly at the office
- Continuous support with evaluation and certification projects, including creating the relevant documentation and communication with customers
- Cooperation and independent project work in the context of looking after customers



REQUIREMENTS

- Technical background
- Minimum 4 years of experience auditing Quality Management Systems of IVD manufacturers (externally from notified body or internally)
- Fluent written and spoken English (any other EU language is a plus)
- Willingness to travel up to 50% in Europe

INTERESTED TO EXPLORE THIS FURTHER?

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

elewed