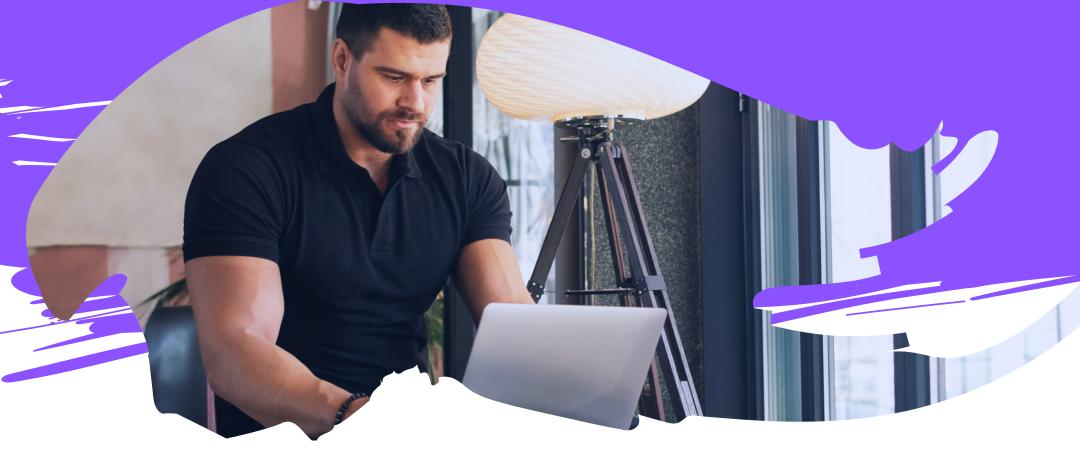


### THE COMPANY

The In Vitro Diagnostic Regulation is THE biggest change that the IVD industry has seen. Experts have estimated that over 80% of IVDs will need the input of a Notified Body, compared to just 20% that we see today.

This has created a great opportunity for IVD professionals to be part of a pioneering change impacting the industry. The clinical reviewer for IVDR is a newly created position at one of the world's leading Notified Bodies. This is a chance to be part of something new, in a company driven by ethics, not profits. You'll be surrounded by highly qualified professionals, and experts in their respective fields and also have the chance to see, firsthand, some of the most innovative technology in the IVD world!



### THE OPPORTUNITY

This is a fantastic opportunity offering regular hours, great stability, and a global working environment.

Recently named "Top employer in Germany", the leadership style is cooperative and really focussed on collaboration. In this role, you'll join a highly successful team, which has enjoyed 50% year-on-year growth.

You'll be constantly learning and expanding your area of expertise and have access to world-class indepth training on the new IVDR.



## THE RESPONSIBILITIES

- Checking and evaluating clinical evaluations for IVD products according to applicable regulations
- Creating reports on deviations and deficiencies in the assessments
- Supporting and advising employees and colleagues
- Being the key contact for medical device manufacturers
- Supporting during audits at manufacturers
- Review and scientifically challenge the clinical data presented by the manufacturer in the clinical evaluation and any clinical investigations
- Assess the manufacturer's clinical evaluation reports and corresponding documents
- Draw up reports demonstrating the result of the relevant conformity assessment activities



# YOUR QUALIFICATIONS

- Scientific academic degree
- 4+ years of professional experience with in-vitro diagnostics in industry or laboratory testing, with a focus on clinical risk assessment and performance evaluation
- Proficiency in English (German is a plus)
- Based in Germany

# INTERESTED TO EXPLORE THIS FURTHER?

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

Would you like to find out more about our open opportunities? Visit <a href="https://www.elemed.eu/vacancies/">https://www.elemed.eu/vacancies/</a>