

Join our **TEAM!**

Because Patient Safety Matters

Product Reviewer In Vitro Diagnostic Devices (IVD)

QMD Services is seeking an experienced **Product Assessor (IVD)** to join our **dynamic and international team**. The successful candidate will play a key role in conformity assessment activities under the IVDR, contributing to high-quality, compliant, and timely certification decisions.

This position is particularly suited to professionals with **prior Notified Body experience**, preferably with **audit experience** and/or acting as a **Final Reviewer** for IVD products.



 **M/F/D**

 **Fulltime**

 **Remote**
(willing to travel within Europe)

For Austrian applicants the annual salary for this position according collective agreement is **€ 47.546,94** gross per year. The actual overall compensation depends on your individual qualification, experience and legal requirements by your country of residence.

Who we are

QMD Services is the only Notified Body for the certification of in vitro diagnostics and medical devices in Austria. We are extending our team of professionals for this responsible role in a highly regulated environment. Our company language is English, as our specialists and employees come from all over Europe and we serve global clients.

We offer

- Competitive package (based on experience)
- Opportunity to work in a dynamic and international environment
- Engagement with a wide range of innovative IVD technologies
- Professional growth within a respected conformity assessment organization
- Collaborative culture with experienced regulatory professionals

Please send your full application and Curriculum Vitae to

recruitment@qmdservices.com

Headquarters

Zelinkagasse 10/3
1010 Vienna, Austria

Operations Office

Am Winterhafen 1
4020 Linz, Austria

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Key Responsibilities

- Perform conformity assessments of **In Vitro Diagnostic (IVD)** devices in accordance with IVDR (EU) 2017/746 and applicable standards
- **Review and assess technical documentation**, including performance evaluation, risk management, and quality management system documentation
- Act as **Final Reviewer/Decision Maker**, ensuring consistency, regulatory compliance, and robustness of assessment conclusions
- Participate in or support **QMS and product-related audits**, where applicable
- On-site audit and off-site review as part of Conformity Assessment for IVDR
- Release of batch test results of high risk devices (mostly class D products)
- Collaborate with internal teams (assessors, auditors, certification staff) in an international environment
- Maintain up-to-date knowledge of regulatory requirements, guidance documents, and best practices
- Maintain client specific administrative work (reports, contracts, conformity assessment programs)

Application Requirements

- The candidate's **CV must clearly indicate current code coverage** (e.g. IVDR scope codes / product codes covered)
- Relevant certifications, audit qualifications, or training records should be included where applicable

Required Qualifications & Experience

- University degree in relevant scientific discipline
- **Previous experience within a Notified Body** (preferably more than two years)
- Broad coverage of IVDR different type of devices in the design, manufacture, or testing of devices
- Solid understanding of IVDR classification rules, conformity assessment routes, and applicable harmonised standards
- Demonstrated experience in IVD product assessment under IVDR
- Audit experience (QMS and/or product-related) is strongly preferred.
- Experience acting as a Final Reviewer/ Decision Maker is a plus

Skills & Competencies

- Strong analytical and regulatory decision-making skills
- Excellent written and verbal communication skills
- Fluent in English (working language); German is desirable
- Ability to work independently while contributing effectively to a multicultural, international team
- High attention to detail and strong documentation skills