

# Clinical Reviewer (MDR)

To strengthen our team, we are looking for a freelance

As a Clinical Reviewer, you will evaluate the clinical documentation of medical devices to ensure compliance with EU regulations and standards. For your work you will be supported by our clinical team and work close together with an internal clinical reviewer.

## Your Role

- Conducting reviews of Clinical documentation from manufacturers
- Assessing clinical data for safety, performance, and benefit-risk profiles of medical devices
- Verifying the alignment of clinical evidence with applicable harmonized standards and guidance documents (e.g., MEDDEV, MDCG)

## Required Education and Experience

- Board certified clinical specialist (MD, currently registered)
- Familiarity with MDR and relevant guidance documents is of advantage
- Notified Body experience is of advantage

## Job-specific competences

- Provide reliable, impartial and unbiased judgement
- Work to specified deadlines
- Ability to draw up detailed/accurate records/reports



# Join our TEAM!

## Because Patient Safety Matters

## 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

- A job with purpose
- A long-term perspective with development opportunities in a rapidly growing company
- Option to work from home with flexible working hours
- International work environment
- Dynamic, agile working environment

Please send your full application  
and Curriculum Vitae to

[recruitment@qmdservices.com](mailto:recruitment@qmdservices.com)

### Headquarters

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