Clinical Evaluation Leader

Bavaria, Germany OR Remote in Europe

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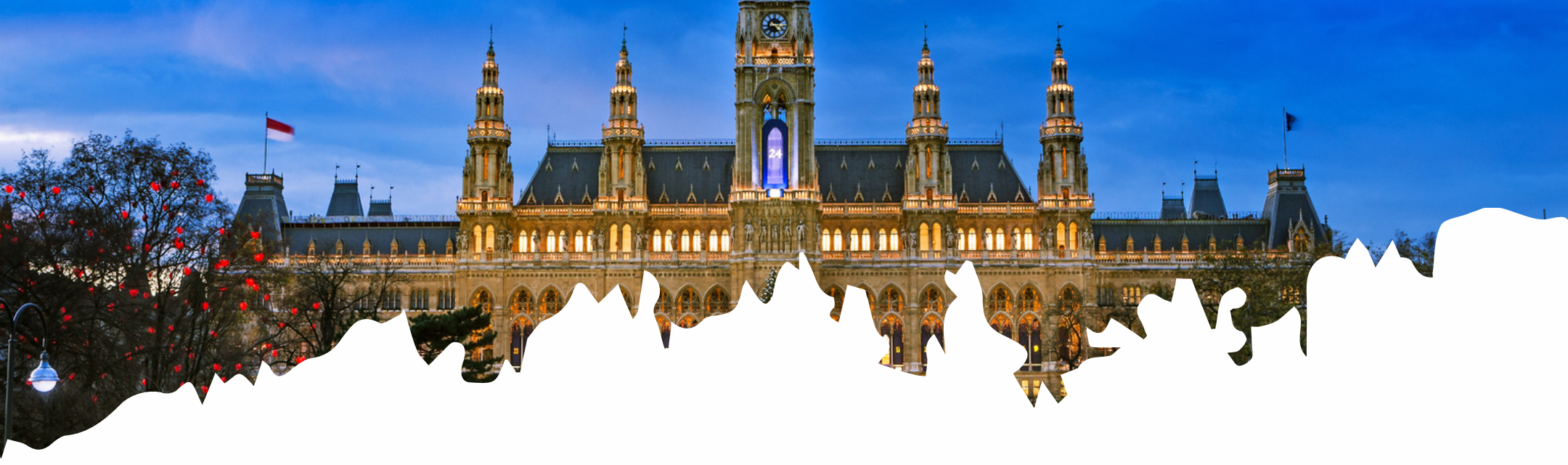
The company

This company is the definition of small and mighty, they are by no means an unstable start-up but if you are looking to join a large, slow-moving corporation with thousands of employees, this isn’t the place for you. If, however, you enjoy working in a business where you can have an impact and shape the future of the organisation, read on.

With a focus on providing regulatory affairs, quality and clinical solutions to manufacturers globally, this company has seen a huge amount of success in the MedTech industry. Each day looks different when you work here with new challenges arising from the variety of customers and products so you can be sure you will not get bored.

Working collaboratively in a solution-oriented way is the key to this company’s success and will allow you to continually develop your skills and knowledge whilst maximising what you already know. Everyone knows each other by name and enjoy attending events together at their office in the heart of Bavaria.



The Opportunity

You will have the opportunity to lead the clinical evaluations team and shape your role as a Clinical Evaluation Leader, this is a newly created position so you can really put your stamp on it. As well as remaining hands-on with Clinical Evaluation activities, you will also have the opportunity to lead and develop the team of 4 clinical evaluation consultants and define the strategic direction for the department. You will report directly to the company's CEO and have direct access to senior management in the company.

This position can be based remotely anywhere in Europe and does not require frequent travel.

Responsibilities

Your responsibilities include but not limited to:

- Write, develop and review high-quality clinical evaluation documents and reports for medical devices and ensure compliance with the Medical Device Regulation and other relevant requirements
- Create technical documentation such as clinical evaluation plans, CERs, PMCF plans, PSURs and more
- Provide customers with support and advice
- Manage and develop the clinical evaluations team (4 direct reports)
- Coordinate cross-functionally with other teams to ensure alignment and compliance to the company's goals and necessary regulations
- Manage relevant timelines required internally and for customers from a clinical perspective
- And more...

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Your qualifications

- 4+ years CER writing experience in the medical device industry
- Fluent speaking, reading and writing in English. German is a bonus.
- Solution-oriented/problem solver

Interested in further CONVERSATION?

Please send your CV to kristina@elemed.eu
to arrange a confidential career discussion.

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