SenTec is a Swiss medical device company headquartered in the Basel area. We develop, manufacture and market innovative solutions for noninvasive patient monitoring. Learn more about us – watch our company video: https://youtu.be/oMSD9QspHbE

To strengthen the Regulatory Affairs team, we are looking for a highly motivated

Senior Clinical and Regulatory Affairs Specialist 80%-100% (w/m)

Key tasks

- Responsible for establishing and managing Clinical Affairs
- Perform Post-Market Clinical Follow-up; request, manage and monitor clinical studies as needed
- Review, structure and revise existing Clinical Evaluation documents (over life cycle of products)
- Responsible for Post Market Surveillance according to current European Medical Device Regulation
- Analyze constantly data input channels for clinical, safety and performance data
- Support in managing Technical Files of existing and new products for international regulatory submissions
- Ensure compliance to the European Medical Device Regulation 2017/745 and be up-to-date with changes in international regulatory legislation and guidelines
- Proactive cooperation with Product Management and Development and other internal departments within the company

Our offer

- Engagement in a fast growing company
- Attractive compensation package
- You will work in an environment with flat hierarchies and short decision-making paths. In this way you can get involved and help shape the company
- You will benefit from an exciting and varied range of tasks
- A positive working atmosphere, a great team and good public transport connections (direct connection from Basel SBB)

Your profile

- Master in the field of technical, natural science or medicine
- At least 3 years' experience working in a similar position
- Open-minded person with hands-on attitude and good team player
- Excellent English skills, German is a plus

Keen on contributing to our growth?
We look forward to getting to know you!
Please send your complete application to jobs@sentec.com

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