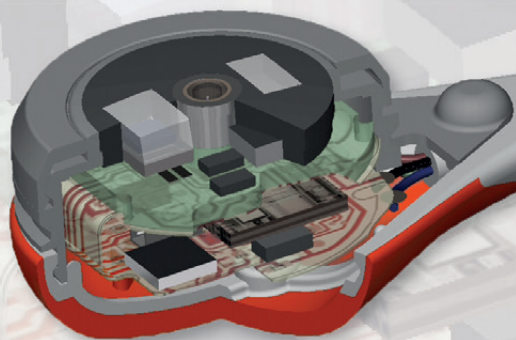


sentec.



Sentec is a Swiss medical technology company headquartered near Basel, Switzerland. We develop, manufacture and market innovative solutions for non-invasive patient monitoring. At Sentec Monitoring Ltd., a subsidiary in Budapest, we do research and development for the Sentec group with a wide range of service.

Learn more about us – watch our company video:
<https://youtu.be/oMSD9QspHbE>



Company Video

To strengthen our Team, we are looking for a highly motivated

Regulatory Affairs Specialist (f/m)

Key tasks

- Responsible for product registration world-wide (USA, EMEA, APAC) as well as keeping up-to-date existing registrations
- Compilation of submission dossiers in close collaboration with internal stakeholders and external regulatory agents
- Perform Risk assessments of the obligation to report incidents and; if required, reporting of incidents to the authorities and support implementing necessary measures internally (CAPAs) and in the market (FSCAs)
- Implementation of updated regulatory requirements and relevant standards for continuous monitoring

Our offer

- Meaningful activity on products that support the therapy of patients
- Flat hierarchy, short decision-making processes, interdisciplinary collaboration
- Opportunity to contribute and develop your skills in an agile company

Your profile

- Degree in life sciences and experience in a Regulatory Affairs position for medical devices
- In depth knowledge of the regulatory framework of medical devices in the legal and normative landscape
- Strong analytical thinking and ability to compile scientific data and summarize results
- Fluent in English, German is a plus

Are you keen on contribute to our growth? We look forward to getting to know you! Please send your complete application documents to jobs@sentec.com

Sentec Monitoring Ltd., Mr. Peter Krammer, Managing Director, Budapest, Hungary, Phone number: +41 81 330 09 69
www.sentec.com