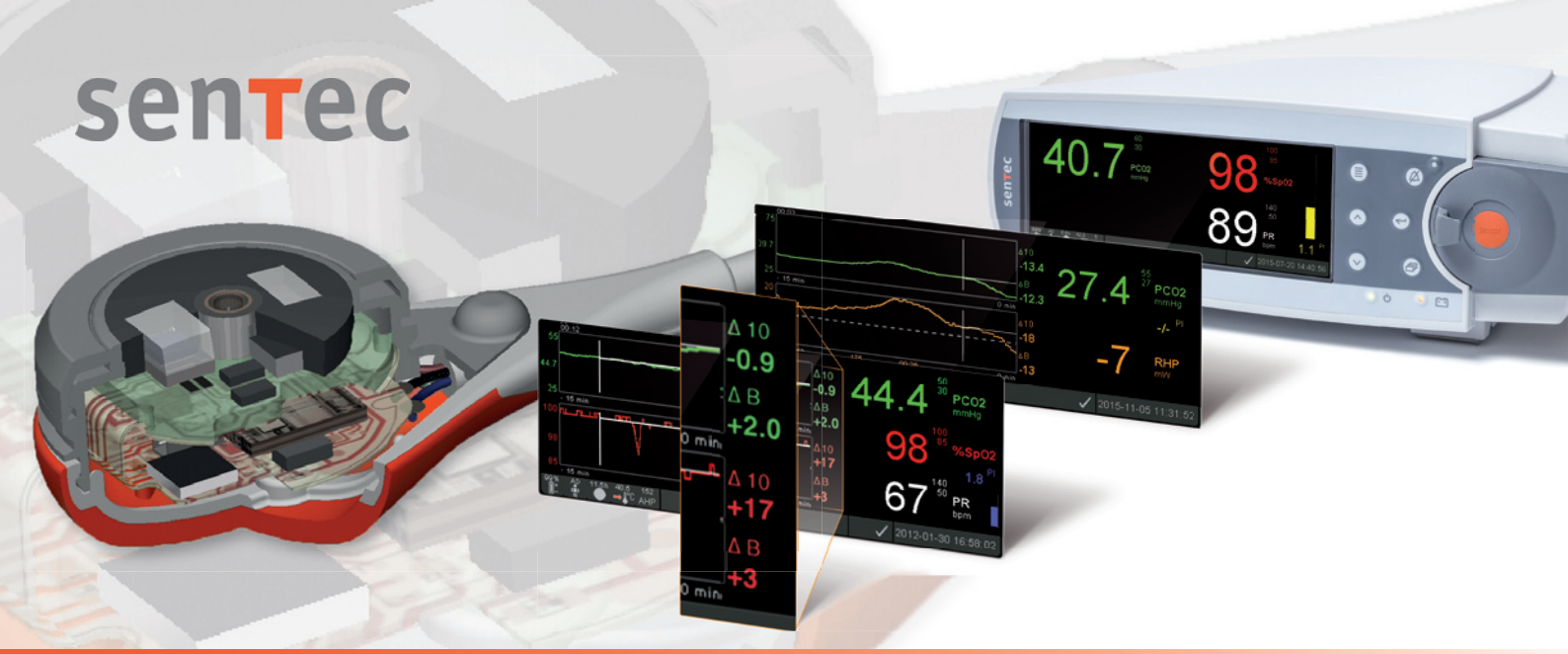

SenTec is a Swiss medical device company headquartered in the Basel area. We develop, manufacture and market products for noninvasive patient monitoring.

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

We are looking for a

Senior Regulatory Affairs Specialist (f/m)

Key tasks

- Developing Regulatory Affairs strategies for new products to be launched world-wide (USA, EMEA, APAC, LAT) 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)
- Support partner companies (distributors) with regard to regulatory issues
- Implementation of updated regulatory requirements and relevant standards for continous monitoring

Our offer

- Opportunity to plan, execute and control your own projects
- Form, motivate and lead interdisciplinary groups
- Flat hierarchy and short decision processes
- Expand your knowledge in a growing medtech field

Your profile

- Degree in life science and at least 5 years of relevant work experience in a Regulatory Affairs position for medical devices
- Practical experience with international submissions, preferably registrations in AP/MEA
- 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 and German

We looking forward to get in touch with you! Please send your complete application documents to: jobs@sentec.com

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