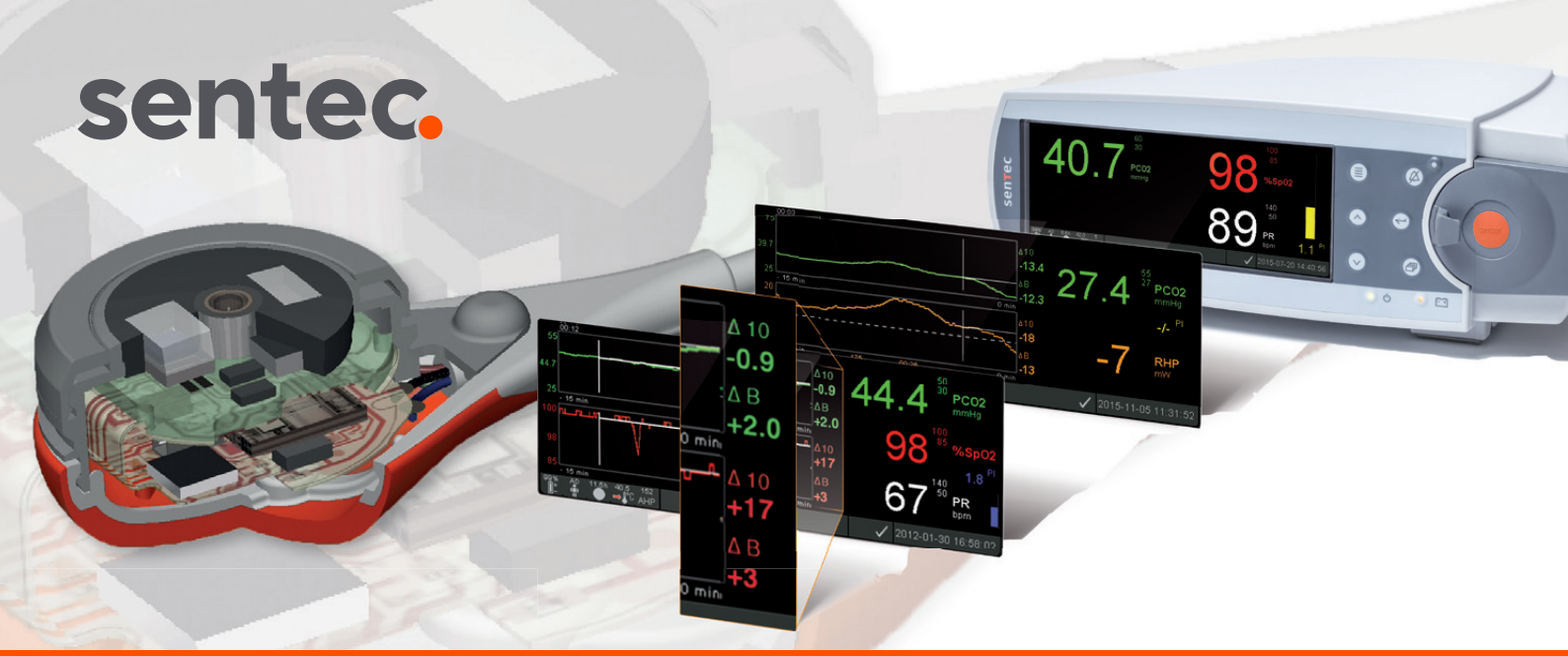
sentec.



Sentec, a „deeptech" company, develops, manufactures and sells non-invasive, digital sensor systems to specialized departments in hospitals worldwide.

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

Company Video

We are looking for a

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

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

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www.sentec.com