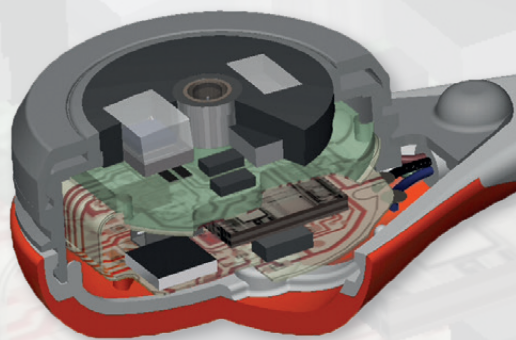


sentec.



Sentec - a „deeptech“ company - researches, 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 80 – 100% (f/m/d)

Key tasks

- Product registration world-wide (EMEA, APAC), keeping existing registrations up to date
- Compilation of submission dossiers in close collaboration with internal and external stakeholders
- Perform risk assessments and if required, report incidents to the authorities
- Follow up of national regulatory requirements and 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 preferred
- In depth knowledge of the regulatory framework of medical devices
- 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 complete application documents to jobs@sentec.com.

Sentec AG, Mrs. Corinne Gueldali, Human Resources, Ringstrasse 39, 4106 Therwil, Switzerland, Tel +41 61 726 97 66
www.sentec.com