For our location in <location>, starting <date|immediately>, we are looking for a

**Regulatory Affairs Manager**

# Your tasks

You will help us obtain approval for our innovative medical devices quickly and safely and master regulatory challenges in the long term.

To do this, you will, <together with your team|in collaboration with XY>,

* prepare, review, and submit approval documents,
* communicate with notified bodies and international regulatory authorities (e.g., the FDA),
* provide regulatory support for the new and further development of our devices,
* act as a consultant for our employees and business partners on regulatory issues,
* monitor and communicate regulatory changes
* < evaluating and reporting of incidents and recalls to the authorities if necessary>
* <assess our quality management system for conformity with national and international laws, regulations, guidelines, and standards>

# Your qualifications

As our ideal candidate, you

* have a degree in engineering or natural sciences,
* are very familiar with the national and international regulations relevant to regulatory affairs,
* have several years of experience in regulatory affairs with a medical device manufacturer,
* communicate and negotiate confidently and convincingly in English and <additional language>,
* have a structured and pragmatic approach to your work and
* are a team player who can balance the interests of regulatory affairs, quality management, and the development department.

# Your future environment

We are a <company description>

In addition to attractive, performance-related remuneration, we offer our employees a safe workplace in an innovative environment characterized by freedom and responsibility and many opportunities for professional development.

Interested? You can reach us at <contact details>.

No matter which form of communication you choose, we treat all forms of communication strictly confidential and look forward to hearing from you.