Our client is an **innovative medical device company.** The company develops and manufactures a breakthrough solution that improves **patient care.**

The company commercializes its devices in **European hospitals** and targets a global business development.

To ensure the growth of our client, we are looking for a:

QA Manager (M/F)

RESPONSIBILITIES

As a **QA Manager**, you will ensure the **compliance** and **efficiency** of **quality processes** in line with ISO 13485 standards. Based in **Liège**, Belgium, with occasional EU travel, you will implement and maintain **quality management systems** and work closely with other departments to meet quality requirements.

Your main responsibilities are:

- Maintain and monitor Quality objectives.
- Ensure compliance with **ISO 13485:2016** and other applicable **QMS standards**.
- Act as the management representative for quality matters (ISO 13485:2016).
- Promote Quality Systems, Objectives, and Policies.
- Manage **internal** and **external** audits, including supplier audits.
- Host **third-party** audits (notified body, competent authority and customer).
- Oversee the complaint and feedback process.
- Collaborate with operations to establish quality agreements with suppliers.
- Identify and implement **process improvements**.
- Analyze risks and trends in complaints, deviations, and audits to define **corrective** and **preventive** actions.

PROFILE

- You hold a Bachelor's or Master's degree in a relevant technical field.
- You have at least **5 years of experience** in a **highly regulated sector**, knowledge of ISO 13485:2016 is an asset.
- You understand the regulatory environment for medical devices.
- You bring strong communication, organizational, and problem-solving skills.
- You are able to plan, organize, and execute tasks effectively in your field.

- You are practical, self-disciplined, and work well independently.
- You demonstrate **team spirit**, adaptability, and a customer-oriented mindset.
- You have a strong attention to detail and are committed to **regulatory compliance**.
- You communicate fluently in **English** (both written and spoken), with French or other languages considered a plus.
- You are open to travel up to **20%** and have a valid driver's license.

OFFER

- A challenging and diversified position within a **high-potential fast-growing innovative medical device company**.
- The opportunity to participate in the development of the company.
- To work in a **human size**, dynamic, respectful, and professional environment.
- **International** exposure, with learning and development opportunities.
- An attractive compensation package in line with the position responsibilities and your experience.

INTERESTED ?

Please send your CV together with an adapted cover letter via URL or to recruitment@pahrtners.be.

YOUR APPLICATION AND RELATED INFORMATION WILL REMAIN STRICTLY CONFIDENTIAL.