Based in Gembloux, **Intressa Vascular** is a clinical-stage company developing the innovative solutions to address life-threatening **cardiovascular** conditions such as aortic dissection.

Their mission is to help physicians address the devastating effect of aortic dissections by providing next-generation **endovascular treatment**. For additional information about the company, please visit https://intressavascular.com/

In order to strengthen the clinical team in Gembloux, we are looking for a (m/):

Clinical Project Manager (M/F)

RESPONSIBILITIES

Reporting to the VP Clinical Affairs, and working closely with Medical and Regulatory Affairs, you will be responsible for the collaborative, compliant and timely management of the assigned **clinical projects** which are at the heart of the company's strategy and development roadmap.

Your main responsibilities are:

- Lead the cross-functional Clinical Project Team and represent it on the **Program meetings.**
- Manage clinical projects, including development, **review and approval of clinical documentation** (e.g. protocols, investigator's brochure), identification and selection of sites and service providers, EC/CA submissions, and regulatory reporting.
- Ensure continued **oversight** and management of the **sites and service providers**.
- Ensure timely and adequate safety documentation, review, and reporting (AEs and DDs).
- Keep the TMF up-to-date at all times, providing guidance to the Clinical Assistant as required.
- Ensure continued project compliance with laws and regulations, applicable SOPs, study protocols, and manuals.
- **Facilitate transversal communication** and cross-functional interaction within teams.
- Contribute to the company's inventive step to develop, secure, and protect company's know-how, trade secrets and intellectual property assets.

PROFILE

- Degree in Health Sciences, Life Sciences, or related medical/scientific field preferred. Relevant work experience may be considered in absence of degree.
- A minimum of **5 years of experience in the management** of medical device studies, including of **Contract Research Organization (CRO).**
- In-depth practical knowledge of **Good Clinical Practice** (ISO14155) and clinical research regulations.
- Keen interest in scientific and technical issues related to clinical study.
- Strong problem-solving and critical thinking skills, attention to details.
- Proactive, flexible, team player, with excellent organizational, time management and communication skills.
- Able to work and collaborate effectively with internal and external actors to achieve company objectives.
- Fluent in English, written and spoken. A good command of any other language is an asset.

OFFER

- A challenging position with a high potential innovative medical device company.
- Working in a human-sized, collaborative, and respectful environment
- A variety of interesting contacts with international exposure.
- An attractive compensation package in line with the position's responsibilities and your experience.

INTERESTED ?

Please send your CV together with an adapted cover letter via https://www.pahrtners.be/fr/offres-demploi/clinical-project-manager-mf/ or to recruitment@pahrtners.be.

Your application and related information you would share will remain strictly confidential.

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