Clinical Research Associate (CRA) / Clinical Project Manager (CPM)
Full-time, Munich, Germany

TRiCares is dedicated to bringing innovative, minimally invasive treatment solutions to patients with failing tricuspid valves. We are looking for a proactive and skilled Clinical Research Associate / Clinical Project Manager to join our dynamic team in Munich and support us in our mission to improve patient care.

Unique opportunity for a Clinical Research Associate / Clinical Project Manager to join a small, expanding medical device company in Munich, Germany, where you will work closely together with the clinical management team. TRiCares GmbH is a manufacturer of cardiovascular medical devices for the minimally invasive treatment of tricuspid valve diseases.

Key Responsibili ies:

- Oversee clinical project management and day-to-day clinical operations throughout the entire study lifecycle, including start-up, execution, and close-out.
- Contribute to the development and writing of clinical study documentation.
- Support the submission of study-specific documents to regulatory authorities, including ethics committees and competent authorities.
- Ensure study conduct aligns with the clinical study protocol and relevant regulations.
- Maintain accurate and up-to-date study documentation and Trial Master File (TMF), including the collection, administration, and tracking of essential documents.
- Serve as the primary point of contact for study sites, ensuring prompt resolution of inquiries to maintain protocol compliance.
- Support timely clinical data collection, review, and cleaning activities to ensure high data quality and resolve any discrepancies.
- Conduct co-monitoring and monitoring visits, as well as other sponsor-initiated visits at study sites.
- Schedule and prepare for meetings and teleconferences.
- Monitor study progress and provide regular updates to Clinical Affairs leadership.
- Collaborate with cross-functional teams to ensure seamless execution of clinical studies.

Qualifications and Experience:

- Bachelor's degree in Life Sciences, Nursing, or a related field.
- Minimum of 2 years of experience as a Clinical Research Associate (CRA) or Clinical Project Manager (CPM) within the medical device industry or a Contract Research Organization (CRO), preferably with Class III devices.
- In-depth knowledge of ISO-14155, ICH-GCP, MDR, and associated guidelines.
- Experience with electronic data capture and tracking systems.
- Monitoring experience is advantageous.
- Knowledge of cardiovascular medicine is a plus.
- Strong attention to detail, with excellent organizational and teamwork skills.
- Exceptional communication abilities.
- Proficiency in Microsoft Office.
- Proficiency in English is required; German language skills are an advantage.

What We Offer:

A dynamic and innovative work environment with opportunities for professional growth.

Collaboration with a highly skilled and passionate team in a truly multinational company, with colleagues from all over the world.

Competitive salary and benefits package.

How to Apply:

Interested candidates are invited to submit their CV and a cover letter detailing their relevant experience and motivation for the role to Kalitvent@tricares.de.