Miracor Medical is a class III medical device company specialized in improving clinical outcome of patients with impaired cardiac function (specifically acute infarcts), active in interventional cardiology. The company has developed a high potential proprietary technology that consists of hardware, software and catheter(s).

The technology is in advanced clinical stages for AMI/STEMI patients, with various potential additional therapeutic applications (among other high-risk NSTEMI and heart failure). The company expects to get CE Mark for its next-gen system in H1 2020 and approval from FDA to run a pivotal study in the next 12 months.

Miracor Medical (www.miracormedical.com) offers a culturally diverse English-speaking working environment in the heart of Europe, Belgium, a few hours away from Europe's capital cities. The offices are based in Awans.

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

Clinical Application Specialist

Responsibilities:

- Provide practical and technical guidance to HCPs on the product
- Provide onsite case support (mainly in UK and France) within clinical trials and commercial setting.
- Assist in clinical cases and ensure that HCPs are well trained and educated with regards to patient selection, treatment planning, optimum use of system, etc.
- Training and education for internal employees / representatives and external parties based on best practices of device usage and the latest clinical findings
- Monitor site performance and treatment outcomes at the sites.
- Facilitate cross functional and core team relationships with the team to enable achievement of goals.
- Ensures proper IFU and User instructions and where applicable clinical trial protocol adherence
- Be proactive in dealing with product-related site request and in solving issues
- Collecting information on product malfunctioning and deficiencies and reporting them via company systems
- Support the R&D function with respect to product development and / or product modifications
- Engage with HCPs, build relations and credibility to be a high level spokes partner
- Work with team to identify and continuously improve workflow standards that are compliant with regulatory guidelines, including internal operating procedures, oversight of study conduct, and any post market requirements necessary.

Profile:

- Previous cardiovascular cathlab experience and record of supported cases
- Willing to travel 80-100%

- Excellent command of English and French

Offer:

- A challenging and diversified position within a high-potential innovative medical device 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/email to Miracor at office@miracormedical.com . Ref: CAS. Your application and related information will remain strictly confidential.