GlyCardial Diagnostics	Human resources	
Job offer 2019/01	rrhh@glycardial.com	93 4020259

About us

GlyCardial Diagnostics is a spin-off company of the IR-Hospital de la Santa Creu i Sant Pau and the Spanish National Research Council (CSIC) focused on the development of a novel in vitro diagnostic device for myocardial ischemia. The technology is based on the detection of Apo J-Glyc in blood as a biomarker for the early diagnosis of cardiac ischemia and the prediction of patient's evolution after an ischemic event.

The company was incorporated in September 2017, closed its first funding round of 2.4M€ in October 2017 and has recently received 1.9M€ from the SME Instrument H2020 funding program.

We are a small company willing to incorporate enthusiastic professionals that aim to be part of a disruptive project and to develop together with the company.

Job offer description

We are looking for a Clinical Development Specialist to drive biomarker clinical validation studies.

Academic training and job requirements

Education

- University degree in Health Sciences
- MSc in Clinical Trials

Knowledge

- Good knowledge of international IVD regulations (including EU directive and regulation and US regulation), of Good Laboratory Practice (GLP) and Good Clinical Practice (GCP)
- Computer skills: Office, database management
- High English level (demonstrable)

Experience

- At least 3 years of previous related experience in biomarker validation clinical trials and preferably previous experience in a biomedical start-up company

Competences

- High level of commitment and working capacity
- Methodical and organised
- Attention to detail and good problem-solving skills
- High planning and scheduling skills
- Ability to work under tight deadlines
- Ability to manage confidential information with discretion
- Ability to interact professionally with all organizational levels
- High communication and team working skills
- Highly motivated and proactive

Key Duties

- Design, review and/or conduct literature searches and complete literature reviews for design concept, protocol preparation, clinical risk assessments and/or clinical evaluation reports.
- Ensure national and country clinical/regulatory requirements compliance.
- Plan, track and monitor all clinical activities.
- Oversee the progress of clinical trial and be the contact person of the company with the CRO, the central laboratories and the study centres, and prepare reports.
- Train staff on protocol specification and assess local sites and laboratory.
- Administer and enter study data into electronic systems efficiently.
- Participate in site meetings if needed and prepare site reports.
- Evaluate documents for clinical processes, resolve discrepancies, evaluate procedures according to required protocols, and maintain an effective management control.
- Evaluate trial data, resolve queries, assist research scientists and audit teams.
- Maintain trial data and ensure accuracy of study materials.

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- Provide update on protocol issues, manage phone calls and provide status reports including updated patient lists.
- Collaborate with the regulatory affairs team of the company.
- Report to and support the Management team of the company.

The offer

- Estimated annual gross salary: We offer a competitive salary commensurate with the qualifications and experience of the candidate.
- Target start date: March 2019.

Application procedure

All applications must be sent by email to the following address rrhh@glycardial.com with the subject "**Job offer 2019/01".**

Applications must include:

- A motivation letter
- A complete CV including contact details
- Contact details of 2-3 referees

Application deadline

Please submit your application by **February 4th 2019**

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