The Vall d'Hebron Research Institute (VHIR) is a public sector institution that promotes and develops the research, innovation and biosanitary teaching of the Vall d'Hebron University Hospital. Through the excellence of our research, we identify and apply new solutions to the health problems of society and we contribute to spread them around the world.

In April 2015, the **Vall d'Hebron Research Institute (VHIR)** obtained the recognition of the European Commission **HR Excellence.**
This recognition proves that VHIR endorses the general principles of **the European Charter for Researchers and a Code of Conduct for the Recruitment of Researchers (Charter & Code)**.
Thus, there are no restrictions of gender, national origin, race, religion, sexual orientation or age and **candidates with disabilities are strongly encouraged to apply.**

Research technician

Cardiovascular Diseases research Group

VHIR offers a vacancy/vacant position for a University Graduate to work as Pharmaceutical research within Cardiovascular Diseases Department. More information about our group can be found here:

http://www.vhir.org/portal1/grup-lines.asp?s=recerca&contentid=186716&idrefer=186717

JOB DESCRIPTION

Education and qualifications:

Required:

Graduated in pharmacy and in possession of a master's degree in clinical trial monitoring with proven experience in cardiovascular diseases (minimum 10 years).
Post-graduate degree in the pharmaceutical field and specifically in the legal framework of conducting clinical trials and post-authorization studies (of cardiovascular diseases) will be highly valued.
Authorship in publications of scientific articles in the cardiovascular field will be valued.

Experience and knowledge:

Desired:

- Good degree of knowledge en la monitorización de ensayos de enfermedades cardiovasculares
- Experience in monitorization of cardiovascular disease trials (10 Years at least)
- Good degree of knowledge in the legal framework for the conduct of clinical trials and post-authorization studies of cardiovasculars disease
- Strong sense of responsibility, initiative, self-motivation and social skills as key personal abilities
- Ability to work independently and as a team
- Good level of English
- Postgraduate or course in the pharmaceutical field about cardiovascular diseases
- Author of publication (about cardiovascular diseases) in journeys with high impact will be valued

Main responsibilities and duties:

- Inclusion monitoring, randomization, data quality, preparation of queries, problem solving, pharmacovigilance tasks and trial management, including the preparation of all necessary documentation for the completion of all administrative and legal

requirements for the incorporation of new centers, including start-up and follow-up visits. If necessary, of all participating sites until the inclusion is completed.

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- Monitoring tasks such as: preparation and sending of reports, management of study documentation for each center, management of study medication, patient registration, start-up and follow-up visits if necessary, monitoring visits (100% monitoring of 100% patient data) and recruitment status reports. Pharmacovigilance tasks such as: management of SAEs (reception, communication to Authorities and CEICs) and safety reports.

Labour conditions:

- Gross annual salary: 29.271,61
- Start Immediately
- Permanent contract

What can we offer?

- Skillful and social colleagues in a dynamic environment.
- Challenging tasks and a wide range of responsibilities
- Personal training opportunities.
- Flexible working hours.
- 23 days of holidays + 9 personal days.
- Flexible Remuneration Program (including dining checks, health insurance, transportation and more).
- Annual teambuilding events.

How to apply:

Applicants should submit a full Curriculum Vitae and a cover letter with the reference CV´s Farma to the following email addresses: mteresa.fernandez@vhir.org and (seleccio@vhir.org) .