



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.**

Clinical Research Monitor Rheumatology Research Group

VHIR offers **two vacant Clinical Research Monitor positions** looking for a self-motivated person with experience in clinical research and/or clinical trials data monitoring

JOB DESCRIPTION

Education and qualifications:

Required:

- Bachelor's degree in science
- Master's degree in science

Experience and knowledge:

- More than 4 years of experience in clinical research and/or clinical trials data monitoring
- Excellent interpersonal and communications skills
- Highly organized
- Ability to work under deadlines
- Ability to work with human study participants
- Proficient in word processing and database applications
- Knowledge of auditing
- Knowledge of good clinical practice and regulatory requirements (i.e. personal data protection)
- Fluency in spoken and written English

Main responsibilities and duties:

- Coordinating and executing all aspects of the clinical monitoring process in accordance with good clinical practice and regulatory requirements
- Reviewing data collection for accuracy and completeness
- Training and lead clinical research personnel at collaborating sites in protocol procedures and implementation
- Identifying problems in protocol implementation and conduct
- Reviewing/resolution of regulatory issues

- Completing a monitoring report for each visit conducted
- Interfacing with clinical investigators, medical staff, management, lab technicians and informaticians
- Report ongoing work and site-level problems to the Scientific Project Manager and the principal investigator
- Travelling to collaborating sites (if needed)

Labour conditions:

- Full-Time position (40 hours per week)
- Salary: 23.000-25.000 € (in accordance with qualifications and experience)
- Start date: June 2021

How to apply:

Applicants should submit a full Curriculum Vitae and a cover letter with the reference CV's CResearch2 to the following email addresses: elena.granell@vhir.org and (seleccio@vhir.org) .