

CRA/CTA (Clinical Research Assistant/Associate)

VHIR offers a vacant position within its Academic Research Organization for a CRA/CTA to develop and manage the activities of monitoring of clinical research projects []

JOB DESCRIPTION

Education and qualifications:

Required.

- Degree or equivalent in Health Sciences.
- Training in monitoring clinical trials.
- It is essential a very good command of Catalan and Spanish, and intermediate-high level of English, both written and spoken.

Experience and knowledge:

- Minimum 1 year of proven experience in monitoring of clinical trials and other clinical research projects.
- Responsibility and work organizational skills.
- Independence and flexibility.
- Communication skills and teamwork.
- Availability to travel within the country.

Main responsibilities/duties:

- Making initial, follow-up and closing monitoring visits to the participating sites.
- Preparation of the visit's reports.
- Implementation of the established procedures (SOPs), and participation, if any, in the preparation and review of SOP in the activities involved.
- Main communicator between the sponsor and the investigator (if not together in the same person) in each site of the study.
- Support management activities of the projects in which the Unit is involved: insurance and contract

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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.**

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management, management of adverse events, medication management and/or other materials, making of monitoring and security reports, updating project monitoring tools, monitoring and supervision of the activities carried out in the sites that are monitored.

- Preparation and maintenance of the trial master files.
- Support and assistance in the preparation and organization of researchers' meetings.
- Detection, communication and support in resolving disagreements and incidents that may arise during the course of the trial.
- Support in the management of authorizations, modifications and notifications to the Ethics Committees and competent authorities.
- Collaboration in administrative functions necessary for the proper monitoring of the projects.
- Collaboration in the design of specific materials for each trial or study.
- Registration and control of the activities for each project.
- Collaboration in the organization of workshops, seminars and/or courses established by the Unit.

Labour conditions:

- Full-time position (40h/week)
- Gross Annual Salary: €23.000
- Immediate incorporation.

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.



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HOW TO APPLY

Applicants should submit a full Curriculum Vitae and a cover letter with the reference "CRA/CTA" to the following email address: seleccio@vhir.org

Please check on VHIR's official website the specific dates to submit your application. After this period the vacancy will be closed.

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