

The Vall d'Hebron Research Institute (VHIR) is a public sector institution that promotes and develops the research, innovationand 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.



HR EXCELLENCE IN RESEARCH

In April 2015, the Vall d'Hebron Research Institute (VHIR) obtained the recognition of the European Commission in 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).

VHIR embraces Equality and Diversity. As reflected in our values we work toward ensuring inclusion and equal opportunity in recruitment, hiring, training, and management for all staff within the organization, regardless of gender, civil status, family status, sexual orientation, nationality, religion, age, disability or race.





# **Study Coordinator**

## **Allergology Department**

The Allergy Research Group (Immunomediated Diseases and Innovative Therapies Area, Systemic Diseases Team) is geared toward the goal to look after people living with allergic diseases and improve their quality of live, according to their needs and also their own decisions, through a comprehensive care and personalized treatments. More information about the department could be found on the following link: <a href="https://www.vallhebron.com/es/especialidades/alergologia">https://www.vallhebron.com/es/especialidades/alergologia</a> and from the research group here.

The Allergy Research Group (Immunomediated Diseases and Innovative Therapies Area, Systemic Diseases Team) at Vall d'Hebron Hospital wants to incorporate to its team one Clinical Trial Study Coordinator and Data Entry to give support on specific tasks.

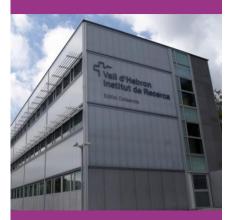
#### JOB DESCRIPTION

#### **Education and qualifications:**

- Bachelor Science degree (Nurse, Biomedicine, Biotechnology, Biology, Biochemist, Pharmacy, or other science bachelor degree).
- Training in good clinical practice and clinical trials methodology is desired.
- Good communication skills and very fluency in written and spoken English.
- Computer user level (Office package, mail).
- Knowledge on SPSS and other data base are desire
- Organized and methodical person with high motivation and initiative
- Quick responsive to time requested by the team and sponsor.

### **Experience and knowledge:**

- Experience with SAP management program (not mandatory).
- Ability to work independently as well as in a team environment



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#### Main responsibilities and duties:

Study Coordinator for clinical trial with the following tasks:

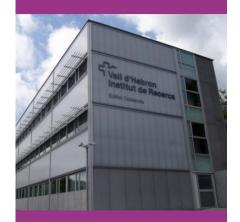
- Coordinate and promote patient recruitment for clinical trial.
- Data entry for a phases II, III and IV studies.
- Keep up-to-date clinical data from source documents to eCRFs.
- Keep up-to-date the Clinical Trial Management System (CTMS)
- Answer queries and give quick feedback to sponsor and clinical team.
- Give support to the clinical team and report to the Clinical Trial Management Unit.
- Ensure maximum adherence to the protocol avoiding deviations.
- Assist the research team in the tasks necessary for the development of studies (vital signs, EKG, etc..)
- Attend site monitoring visits, review and resolve queries in accordance to GCP.

#### **Labour conditions:**

- Full-time position: 40h/week.
- Starting date: immediate
- 6 month of contract, renewable and negotiable.
- Gross annual salary according to salary scale: 26.000€

#### What can we offer?

- Incorporation to Vall d'Hebron Research Institute (VHIR), a public sector institution that promotes and develops the biomedical research, innovation and teaching at Vall d'Hebron University Hospital (HUVH), the biggest hospital of Barcelona and the largest of Catalan Institute of Health (ICS).
- A scientific environment of excellence, highly dynamic, where high-end biomedical projects are continuously developed.
- Continuous learning and a wide range of responsibilities within a stimulating work environment.
- Personal training opportunities.
- Flexible working hours.
- 23 days of holidays + 9 personal days.



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• Flexible Remuneration Program (including health insurance, transportation and more).

#### How to apply:

Applicants should submit a full Curriculum Vitae and a cover letter with the reference "Study Coordinator Clinical Trials in Allergology group" to the following email addresses: Mar Guilarte <a href="mailto:mguilarte@vhebron.net">mguilarte@vhebron.net</a> and Victoria Cardona vcardona@vhebron.net and seleccio@vhir.org before May 15<sup>th</sup>.



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Diversity. As reflected in our values we work toward ensuring inclusion and equal opportunity in

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