



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.



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.

CTA for Clinical Trials/Projects/IRP

Legal Unit

VHIR offers a vacant position for a CTA with experience in various areas of the Healthcare environment to develop tasks within the Legal Unit. This unit ensures the regulatory compliance in all institutional acts, especially in biomedical research projects and IP.

Among the goals of this Units are: to ensure the regulatory compliance in all institutional acts, especially in IP and biomedical research projects; give advice to the scientific community at VHIR of the continuous legal developments that affect biomedical research and, therefore, the transfer of results to society; and to the different services and units at VHIR.

JOB DESCRIPTION

Education and qualifications:

Required:

- Certificate of Higher Education (CFGS/FP II) or Bachelor's Degree in Law, Health Sciences or similar or training in administrative clinical trials activities.
- Fluency in Catalan, Spanish, English, both written and oral (business level)

Desired:

- Master's Degree

Experience and knowledge:

- 2-3 years of experience managing Clinical Trials documentation/contracts.
- Specific knowledge in pharmaceutical and healthcare law and/or previous experience in the field of biomedical research (specifically in the field of clinical trials and EPAs) will be positively valued.
- Excellent organizational skills and client-orientated.
- Ability to work in teams.
- Ability to manage high workloads and work under tight deadlines.



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Main responsibilities and duties:

- Participate in the process of drafting, negotiating and reviewing clinical trials contracts and other clinical research contracts, amendments.
- Give support in any topic related to clinical trials and EPAs (EOMs) within the Legal Unit.

Labour conditions:

- Full-time position: 40h/week.
- Immediate incorporation
- Gross Annual Salary: (Remuneration will depend on experience and skills. Salary ranges are consistent with our Collective Agreement pay scale)

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.
- Flexible Remuneration Program (including dining checks, health insurance, transportation and more).

How to apply:

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