



Job Opening: Preclinical and Regulatory Project Manager

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About Oniria Therapeutics

Oniria Therapeutics is a biopharmaceutical company in the field of precision oncology. It develops first-in-class targeted therapies to overcome cancer persistence. The company was incorporated in 2021 as a Spin-off of the Vall d'Hebron Institute of Oncology (VHIO), the University of Barcelona (UB) and Catalan Institution for Research and Advanced Studies (ICREA) by Dr. Héctor G. Palmer, Dr. Isabel Puig-Borreil, Dr. Xavier Barril, Dr. Carles Galdeano, Dr. Josep Tabernero and Esther Riambau. It is supported by national investors and organizations in the healthcare and life science sector. The company is located at the CELLEX CENTER, in the Vall d'Hebron campus. More information: www.oniriatherapeutics.com

Job description

We are seeking a motivated Project Manager in Preclinical Development and Regulatory Affairs to join our interdisciplinary team to develop new agents designed to modulate tumor dormancy to fight cancer progression. The candidate will participate in a highly translational project with interactions with academic institutions, industry, and clinicians by planning, initiating, and monitoring experiments, analyzing data and drawing conclusions. The role includes to procure services from CROs and coordinate the work with the CROs and other involved. The work involves many contacts both internally and externally, including international CROs/collaborators/research groups, which imposes high demands on communication skills and diplomacy ability.

It is a great opportunity to develop a product for transforming oncology treatments.

Key tasks:

- Work closely with the CEO, CSO and translation team to ensure alignment between translational goals and preclinical development plan.
- Responsible for organizing, overseeing, and managing Oniria's preclinical development activities.
- Primary point of contact for communication with external providers and collaborators.
- Applying project management skills in planning, initiating, risk assessing, executing and tracking of project activities, timelines and budget.
- Managing regulatory strategies and coordinating submissions to regulatory authorities in support of ongoing development programs. Maintaining relationships with regulatory agencies. Participation in the company's scientific meetings.
- Actively maintaining preclinical databases.
- Accurately writing preclinical protocols, study reports or other preclinical related documents to support clinical trials submissions according to project timelines.
- Applying regulatory guideless applicable to preclinical studies to daily work.

Requirements

Must have:

- PhD in biology, pharmacy, biochemistry, biomedical sciences or related discipline.
- Minimum 2-3 years of experience in regulatory affairs and project management working with cross-function teams, preferably including experience with preclinical-stage therapeutics.
- A solid scientific and technical background in early-stage development of drugs.
- Good teamwork and interpersonal skills, as well as independent, motivated, and collaborative person
- Excellent time management skills and ability to multi-task and prioritize work.
- Excellent organizational skills and attention to detail.
- Creative problem-solving skills.
- Fluency in English and Spanish.

Nice to have:

- Experience in the field of oncology drugs.
- Previous experience managing clinical stage therapeutics.

We offer:

- **Open-ended** (*indefinido*), full-time contract.
- **Salary:** competitive and commensurate with experience and qualifications.
- **Expecting starting date:** February 2024
- **Location:** preferably Barcelona, with possibility of remote work.
- Continuous training in a high-quality scientific environment.
- Good working atmosphere.

Application process:

Please send the following documents to info@oniriatherapeutics.com indicating "Preclinical and Regulatory Project Manager" in the email title:

- CV (Name_Surname_CV.pdf)
- Cover letter (Name_Surname_CL.pdf)
- 2-3 contacts for references

Confidentiality in document handling and equality of opportunities policy are guaranteed by Oniria Therapeutics.

Deadline to submit applications:

Submit your application as soon as possible and no later than 15th December 2023

Contact:

Esther Rimbau