JOB OFFER

Drug Development Director – Senior Researcher with Project Management Skills

A selective process is called to fill 1 Development Director position at SUCCIPRO (www.succipro.com), a small, dynamic and innovative spin-off company born from the DIAMET research team (www.diamet.org), which belongs to the INSTITUT D'INVESTIGACIÓ SANITÀRIA PERE VIRGILI (IISPV) (www.iispv.cat).

This is a full-time, indefinite contract position based between Barcelona and Tarragona, with flexible working hours and the possibility of remote work. The successful candidate will join a multidisciplinary team and play a key role in the development of SUCCIPRO's biologic therapy.

Job Responsibilities:

Prepare and lead the drug development process from lead candidate selection through to IND submission and early-phase clinical trials.

Design and oversee preclinical development plans, including pharmacology, toxicology, and safety studies in compliance with regulatory standards.

Coordinate and manage GLP-compliant safety and toxicology studies, ensuring data integrity and adherence to timelines.

Develop and implement regulatory strategies aligned with EMA/FDA guidelines to support successful IND/CTA submissions.

Prepare, review, and submit regulatory documentation including Investigator's Brochures, IND/CTA dossiers, and responses to regulatory agencies.

Collaborate with CROs and CDMOs to ensure quality and compliance in outsourced activities, including manufacturing and analytical development.

Supervise the production of clinical trial material under GLP/GMP conditions, ensuring traceability and documentation.

Maintain up-to-date knowledge of regulatory requirements and proactively assess their impact on development plans.

Work cross-functionally with internal teams (R&D, QA, Clinical) to ensure alignment and integration of development activities.

Contribute to the design of early clinical trials, including protocol development and regulatory submissions.

Ensure proper documentation and archiving of all development activities in accordance with Good Documentation Practices (GDP)

Qualifications:

PhD in Biochemistry or a related discipline is required.

Previous experience in preclinical regulatory drug development is essential (+3 years)

Experience in supervising and managing CROs and CDMOs contracts is highly valued.

Fluent professional English is required.

Proficient computer skills (normal user level).

We Offer:

A dynamic and collaborative work environment.

Opportunities for professional development.

Flexible working hours

Possibility to work from home.

Salary to be discussed according to the profile of the candidate

To apply, please submit your resume, a cover letter outlining your relevant experience, and contact information for two professional references to recruitment@succipro.com.

SUCCIPRO is an equal opportunity employer and encourages candidates from all backgrounds to apply.

SUCCIPRO
INNOVATIVE THERAPIES