Sensorion
The Inner Ear Diseases company

Montpellier, January 8th 2021

CLINICAL PROJECT MANAGER F/M

Company

Sensorion is a pioneering clinical-stage biotech company, which specializes in the development of novel therapies to restore, treat and prevent within the field of hearing loss disorders. Its clinical-stage portfolio includes one Phase 2 product: SENS-401 (Arazasetron) for sudden sensorineural hearing loss (SSNHL).

Sensorion has built a unique R&D technology platform to expand its understanding of the pathophysiology and etiology of inner ear related diseases enabling it to select the best targets and modalities for drug candidates.

The Company is also working on the identification of biomarkers to improve diagnosis of these underserved illnesses. Sensorion has launched in the second half of 2019 two preclinical gene therapy programs aiming at correcting hereditary monogenic forms of deafness including Usher Type 1 and deafness caused by a mutation of the gene encoding for Otoferlin.

The Company is uniquely placed through its platforms and pipeline of potential therapeutics to make a lasting positive impact on hundreds of thousands of people with inner ear related disorders; a significant global unmet medical need.

https://www.sensorion-pharma.com

Mission

Sensorion is looking for **an Clinical Project Manager F/M** based in Montpellier or in Paris.

The Clinical Project Manager (CPM) is the primary person responsible for the planning and execution of assigned clinical trials that may be local, regional or global in scope. This encompasses all aspects that impact the completion of the trial and include development of study project plans, problem solving all trial related issues utilizing available resources, initiation and monitoring of trials in progress, completion, reporting and publication of study results. The position ensures all studies are executed within predefined timelines & budgets and are conducted in compliance with GCP and all applicable standards & regulations in a timely manner.

- CPM is responsible for the overall day-to-day coordination and management of clinical trials from start up through close out activities.
- Directs the technical, financial and operational aspects of the projects, thus securing the successful completion of clinical trials.
- Works with the Clinical Research Director to identify and evaluate fundamental issues on the project, interpret data on complex issues and ensure solutions are implemented.
- The CPM, with support from the Clinical Research Director and any other internal staff, is accountable for ensuring that project deliverables meet time/quality/cost expectations.
- This is a full position based in Paris or Montpellier with a direct reporting to the Director of Clinical Research.

Profile

Experience

- Minimum of five (5) year experience in Clinical Development in pharmaceutical industry or CRO including demonstrated skills and competency in clinical project management tasks or supervisory experience
- Thorough knowledge of clinical studies management;
- Advanced knowledge of ICH/GCPs and current knowledge of FDA and EMEA regulations;

- Can demonstrate experience of successfully managing and/or leading multidisciplinary project teams;
- Teamwork skills and strong team spirit;
- Fluent in English, both written and verbal;
- Willingness to travel up to 20% of time including domestic and international travels.

Diploma/Training

- University/college degree (life science preferred) or certification in a related allied health profession from an appropriately accredited institution (e.g., nursing certification, medical or laboratory technology) or equivalent combination of education and experience that provides the individual with the required knowledge, skills and abilities;

Skills & Abilities

The CPM possesses a high level of the following skills and attributes:

- Sound knowledge of the key principles of cross functional project management (time, quality, cost);
- Financial acumen;
- Displays effective communication skills (listening, oral and written);
- Organizational skills and proficiency at multi-tasking with good attention to detail;
- Demonstrated ability to lead, coordinate teams as appropriate;
- The ability to prioritize workload;
- Cross cultural awareness and ability to adapt appropriately;
- Ability to work independently including adaptability and flexibility skills;
- Good computer skills.

Position to be filled as soon as possible

Send your resume and cover letter to :

- ✓ https://jobaffinity.fr/apply/58a226ng7bmcx1wfz8

Contact:

- ✓ Marie Michelle Conti, Human Resources Manager
- ✓ Email: marie-michelle.conti@sensorion-pharma.com