innate pharma®

JOB OPPORTUNITY

Innate Pharma S.A. is a clinical-stage biotechnology company with a focus on discovering and developing first-in-class therapeutic antibodies that harness the innate immune system to improve cancer treatment and clinical outcomes for patients.
The Company has three first-in-class antibodies in clinical development for treatment of cancer, including two checkpoint blockers in immuno-oncology, a new therapeutic field that is changing cancer treatment by mobilizing the power of the body's immune system to recognize and kill cancer cells. Innate Pharma also has several novel preclinical product candidates and research technologies.
The Company's expertise and understanding of natural killer cell biology have enabled it to enter into major alliances with leaders in the biopharmaceutical industry including AstraZeneca, Bristol-Myers Squibb, Novo Nordisk A/S and Sanofi.
As our research and development pipeline is expanding and advancing, INNATE PHARMA faces new exciting challenges and we therefore now seek one:

HEAD OF CLINICAL OPERATIONS M/F

Permanent Contract
Réf: DCO/PDO/2017/3

Mission

Within the Clinical Department and under the responsibility of the Chief Medical Officer (CMO), your main responsibilities will be the planning, execution and reporting of planned clinical trials according to the clinical development plan of each compound and company strategy.

You provide operational and strategic oversight of the clinical studies ensuring compliance with Good Clinical Practices and regulatory guidelines. You are responsible for consistent and successful execution of operational aspects of clinical trials, with a focus on quality of data collection, cost, and timelines.

You are managing the Clinical Operation team, ensuring recruitment, training and resource allocation.

You participate to the identification and selection of the CROs and establish and follow carefully the clinical studies budget.

You collaborate with other functions within clinical department for the strategic design of clinical studies.

Experience and qualifications requirement

- Bachelor's/Master's degree in a scientific discipline is required , oncology experience preferred
- 10+ years Pharmaceutical development experience
- Functional management experience required (managing CTAs, CRAs, CTMs, and CPMs)
- Well organized, excellent written and verbal communication skills are required.
- Experience and understanding of ICH, and GCP is required

- English (written and spoken) is mandatory and French (spoken) is desirable.
- The position is open to candidates with disabilities.

Please send your resume and a cover letter in support of your application with the reference « DCO/PDO/2017/3 » par email (diane.sherriff@srg.co.uk)