iTeos Therapeutics is a Belgian innovative biotech company based in the South of Brussels (Gosselies). This human-sized structure is focused on expanding the benefits of immunotherapy for cancer patients by developing a proprietary pipeline targeting IDO1, A_{2A}, immune checkpoints and non-immunogenic ("cold") tumors.

iTeos Therapeutics' competitive edge is in the combination of expertise in drug discovery, translational tumor immunology and early clinical trial design. The company uses a unique platform to identify rational combinations of immunotherapies and novel targets. For more information, please visit www.iteostherapeutics.com.

In order to support this challenging growth, we are actively looking for a (m/f):

Senior Clinical Project Manager

Responsibilities :

As a Senior Clinical Project Manager, you autonomously manage global clinical trials in interaction with CROs. You are in charge of all aspects of multiple global clinical studies (Phase I/II), in particular all sites and vendors related issues. You manage the study operational plan and the CRO activities, including project timelines, quality of deliverables and follow-up of approved trial budget.

Your main responsibilities are as follows:

- You develop study plan and manage study timelines.
- You coordinate internal and external clinical development activities.
- You participate in the selection of investigational sites and CROs for assigned studies.
- You provide clinical study teams with trainings on assigned protocol.
- You review and refine clinical operational plans including the study monitoring plan.
- You oversee clinical trial sites' adherence to pertinent regulations.
- You follow-up CROs' activities to ensure agreed deliverables.
- You proactively identify clinical trial risks and suggest mitigation plan.
- You identify and provide solutions to clinical trial issues.
- You contribute to relevant study documentation including clinical protocols, statistical analysis plan, clinical study reports as well as operational plans.

Profile :

- Master or PhD in life sciences or closely related field.
- Minimum of 8 years of clinical research experience in the biotech/pharma industry, with minimum 5 years in the clinical study coordination.
- Experience with Phase I/II clinical trials.
- Thorough understanding of FDA, EMA, ICH and GCP guidelines.
- Previous experience negotiating vendor/site contracts and managing the budgets.
- Autonomous, committed, flexible and problem solving oriented.
- Highly collaborative, strong team spirit and excellent interpersonal skills.
- Excellent oral and written communication skills in English. The command of French is a plus.
- Willingness to travel 20% of your work time internationally and domestically.

Offer :

- A stimulating clinical position within a high-potential innovative biotech company.
- To work in a human-sized, dynamic, respectful and professional environment.
- The opportunity to take part in a challenging scientific and business growth.
- Varied contacts inside the company, the biotech/pharma sector and the scientific world.
- An attractive salary package in line with the position responsibilities and your experience.

Interested ?

Please send your CV together with an adapted cover letter to recruitment@pahrtners.be or via http://www.pahrtners.be/job/senior-clinical-project-manager/

Ref: SCPM.

Your application and related information will remain strictly confidential**.**