CLINICAL TRIAL SUPPLIES MANAGER, CM&C

2 Bridge company

2 Bridge is a Belgian-based consultancy company that provides advice and support on all key disciplines of healthcare product development (discovery, pre-clinical, clinical and product development, registration and life-cycle management). We work globally with startups, biotech, medtech, pharma, and investors. 2 Bridge typically operates via flexible and cross-functional teams, aligned to the project need. Our broad and multidisciplinary expertise allows to address the most complex and challenging tasks during development. For more information, please visit: www.2Bridge.be.

Job description

Are you interested to make a difference and contribute to realize our growth plans? We are looking for a Clinical Trial Supplies Manager to strengthen the CM&C team of 2 Bridge. You will work in a non-hierarchical environment that highly values teamwork and where you will have the freedom to shape your role. You will report to and work closely with the Head of CM&C of 2 Bridge.

The desired expertise for the Clinical Trial Supplies Manager includes:

- Technical knowledge of IRT set-up
- Ability to manage and execute clinical trial supply strategies for clinical studies in alignment with protocol requirements, key study parameters and milestones;
- Ability to select and work with C(D)MOs for project execution.

Projects within the Clinical Trial Supplies group can vary but usually fall within the following scope:

- Preparation of documentation within the area of Clinical Trial Supplies.
- Develop and executes detailed clinical trial supply project plans in close collaboration with stakeholders and in compliance with GxP requirements, local regulations and client processes;
- Support with set-up of IRT system.
- Project management within clinical supply group.

Desired profile

- MSc/PhD in (Industrial/Hospital) Pharmacy, Bio-medical sciences, Bio-engineering, Biology, Chemistry, or equivalent.
- 1 – 3 years prior-experience within clinical trial supply management is highly recommended, although industry-starters with the right mind-set will also be considered.
- Knowledge and/or keen interest in pharmaceutical legislation (GxP).
- Team-player with the ability to work independently.
- Project management skills, good organization and planning skills.
- Analytical, pro-active, flexible and with an eye for detail.
- Interest in the overall process of health care development is a must.
- Interested in working in a multidisciplinary team.
- Enthusiasm with a keen interest to learn new things.
- Good communication skills.
- Fluency in English is a must.

What we offer

- You will be contributing to our ambitious growth plans.
- You will be part of an enthusiastic team where human interactions, teamwork and bringing together different perspectives are highly valued.
- You will work in a small, but growing organization with an informal and non-hierarchical way of working.

How to apply

CV, motivation letter where you:

1. Introduce yourself
2. Explain why you are interested in the job
3. Outline why you think you are the right candidate

Interested? Please send your motivation letter and CV to info@2bridge.be.