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

We are looking for a Clinical Trial Supplies Manager to strengthen the CM&C team of 2 Bridge. The ideal candidate will be able to manage all aspects of clinical trial supplies for early stage up to late stage clinical studies covering set-up, management and close-out.
The desired expertise for the Clinical Trial Supplies Manager includes:

- Development and execution of clinical trial supply strategies for clinical studies in alignment with protocol requirements, key study parameters and milestones;
- Selection and collaboration with C(D)MOs for project execution;
- Technical knowledge of IRT systems;
- Ability to consolidate, maintain and track the clinical trial supply budget;

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

- Project management within clinical supplies;
- Preparation of project documentation (e.g. study specific pharmacy manual and other relevant guides);
- Development and execution of detailed clinical trial supply project plans in close collaboration with stakeholders and in compliance with GxP requirements, local regulations and client processes;
- Set-up of an IRT system;
- Collecting, registering and archiving information and documents in accordance with the applicable GxP guidelines for clinical studies ((e)TMF).

Qualifications/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);
- Quality conscience attitude; can do mentality;
- Project management skills, good organization and planning skills;
- Team-player with the ability to work independently;
- Analytical, pro-active, flexible and with an eye for detail;
- Interested in working in a multidisciplinary team;

- Enthusiasm with a keen interest to learn new things;
- Good communication skills;
- Interest in the overall process of health care development is a must;
- 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 hr@2bridge.be.