JUNIOR PROJECT MANAGER PRODUCT DEVELOPMENT

2 Bridge company

2 Bridge is a Belgian-based 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, 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 junior Project Manager Product Development with a keen interest to learn about managing projects within the domain of CM&C and with an eye for quality within an early-stage product development setting.

The CM&C and Quality project manager will be involved in the execution of early-stage product development and/or support the development of initial Quality Management Systems (QMS) for startups, with support and under the guidance of a senior CM&C/Quality professional.

As representative of the CM&C team, you will manage a wide variety of CM&C activities covering selection of Contract Manufacturing Organisations (CMOs) and management of oversight on Drug Substance or Drug Product activities at the selected CMOs and will thereby get the opportunity to broaden experience quickly.

Key activities of a CMC and Quality project manager are:

- Support the creation and implementation of CM&C related aspects of a development strategy;
- Execute agreed CM&C strategy under support of a senior CM&C leader;
- Support the identification and selection process of CM&C service providers;
- Prepare scientific and regulatory documentation (IMPD/IND) within the area of CM&C;
- Support identification and mitigation of CM&C risks;
- Support the delivery of clinical drug supplies in accordance with development plans;
- Ensure compliance with applicable regulations and guidelines throughout activities;
- Manage the implementation of new QMS at early-stage development start-ups

What we offer

Be part of a multidisciplinary team that values versatility, adaptability and effective cross-functional collaboration, in the spirit of our values: ‘we care, we adapt, we drive to excel’. By being exposed to a multitude of projects and functions, you will quickly gain a broader view of drug development, thereby enhancing your professional growth.

You will work in a non-hierarchical environment that highly values teamwork and where you will have the freedom to shape your role. We believe in fostering a culture of learning and continuous feedback, where everyone feels supported and empowered to succeed.

2 BRIDGE
Science to Life

We build a safe work environment where people feel comfortable speaking up, taking accountability, and balancing their personal boundaries with business expectations. Our team collaborates and supports each other, creating a dynamic and inclusive workplace where you can thrive.

Qualifications/Desired profile

- MSc/PhD in Industrial Pharmacy, Bio-medical sciences, Chemistry, or equivalent.
- Ideally 1-3 years of prior experience within product development, registration and/or within a GMP production environment of large/small molecules highly recommended, although industry-starters with the right mind-set will also be considered.
- Some knowledge and/or interest in pharmaceutical legislation (ICH/GMP,...).
- Enthusiasm with a keen interest to learn new things.
- 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.
- Good communication skills.
- Fluency in English (especially in writing) is a must.

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.