

## **Clinical Supply Manager, Clinical Supply**

We are looking for an experienced Clinical Supply Manager for a challenging and diverse position in Clinical Supply (CS). As our new colleague, you will be providing leadership for clinical supply activities in support of our global clinical phase I-IV studies. You will be responsible for a wide variety of exciting activities in close interaction with external (CRO/CMO`s) and internal Lundbeck partners and be part of a highly-dedicated team, solving daily challenges with a result-oriented mindset.

The department of Clinical Supply currently consists of 11 highly skilled and passionate colleagues who are experts in coordination of the outsourcing related to packaging and distribution of the Investigational Medicinal Product (IMP) as well as activities related to Interactive Response Technology (IRT).

Clinical Supply is part of the CMC area where we strive to develop new innovating treatments to help improve the lives of patients suffering from brain diseases. We work with both internal CMC development and GMP activities as well as external business partners in a dynamic and changing environment.

### **Your job and key responsibilities**

You will be part of a dynamic team responsible for the clinical supply chain from end to end.

Your essential tasks will be:

- Leadership of IMP deliveries as chair of a cross organisational IMP Working Group and by conducting project management for assigned studies and projects including oversight
- Fully Outsourcing of packaging, distribution and IRT related activities with focus on IRT related activities
- Representing the CMC area in several cross-organisational study management teams and providing challenge of and input to the clinical study design and protocols
- Develop and implement the clinical supply design, including planning and coordination
- Ensure efficient supply execution through a data founded and integrated approach
- Close collaboration and numerous daily interactions with internal and external collaborators
- Contribute to and drive the continuous improvement of departmental and cross-organizational processes

The job will require you to have creative and strategic skills to solve complex scientific challenges in an environment where prerequisites and timelines may change overnight. Furthermore, you will be expected to promote a proactive mind-set and create thorough and innovative solutions, as applicable. You can take leadership and are able to ensure progress

within given timelines and in accordance with study budget. At the same time, you can digest complex data output evaluate data quality and act accordingly during study conduct.

Your workplace will be in Valby.

### **We offer**

We value an open, trust-based and creative working environment, great team spirit and a high-quality mind-set. We offer a position where you will make rapid, disciplined decisions, have influence on the development of the clinical supplies area and be responsible for a variety of assignments while you continuously build upon your experiences and professional development.

### **Qualifications**

Our preferred candidate has the following personal and professional qualifications:

- A relevant master's degree in natural sciences or engineering and experience within the pharmaceutical industry or equivalent. Preferably experience in coordinating IRT, clinical supplies, clinical trial management, drug development or other areas of clinical trials
- A strong business understanding as well as excellent stakeholder management, communication and presentation skills
- Experience in project management
- Demonstrated problem-solving skills with the ability to prioritize, align, and simplify; used to manage multiple projects with changing priorities
- Flexible and adaptable, and able to work well in a busy and dynamic environment
- You sense the importance of details and have an interest in IT systems
- Work in a structured manner, introducing innovative solutions as needed
- A reliable person, inspiring confidence, and able to deal with people on all levels. You possess a positive attitude, political flair and a good understanding of foreign cultures combined with empathy and the ability to listen and cooperate
- You have strong business understanding as well as excellent stakeholder management, communication and presentation skills
- You are fluent in spoken and written English.

### **Further information**

For further information, please contact Director, Clinical Supply, Dorthe Lerche Berg, on [DLB@lundbeck.com](mailto:DLB@lundbeck.com). Your application and CV should not be sent via email.

We also recommend that you have a look at our website.

## Your application

Please click on the link below to apply for the job. Applications will be reviewed ongoingly and must be received no later than **August 26 2020**.