Senior Clinical Research Associate / Clinical Trial Manager

Do you want to be part of a dynamic team that drives clinical activities in an inspiring environment? Would you enjoy to work with different projects and with the ability to influence your job? Larix Clinical Operations is now seeking an experienced Clinical Research Associate / Clinical Trial Manager to join our Herlev office.

Larix A/S is a Nordic CRO – we offer full-service solutions within the pharmaceutical, biotech and medical device areas. Our headquarters are located near Copenhagen in the middle of the Medicon Valley region, and we have strong ties to the thriving pharmaceutical and biotech activities in this region.

Clinical Operations consists of experienced and enthusiastic Clinical Research Associates and Clinical Trial Managers responsible for the planning and conduct of clinical activities in the Nordic countries.

At Larix, we maintain a friendly atmosphere – we think having fun while working is important. We are a relatively small company with approximately 55 employees, and we have all the benefits of being able to collaborate across functions, follow clinical study processes from start to finish, and learn from each other. You will have a great influence on the development and on future tasks.

At Larix, you'll be part of our Clinical Operations group in Denmark. You will work from our offices in Herlev but should also be prepared to work at a client's office in the Copenhagen area from time to time.

Your main tasks will include:

- Perform and handle all aspects of trial & site management of sponsor projects
- Perform monitoring (from Pre-study, Initiation, Monitoring and Close-out visits)
- Submission to regulatory authorities and ethic committees
- Perform and handle trial management of small or larger studies
- Prepare or contribute to the update of SOPs and procedures within Clinical Operations

We are looking for an experienced Clinical Research Associate /Clinical Trial Manager – we will get to know you as a proactive and competent colleague who approaches projects with a team players' spirit. At the same time, you are very good at working independently and are able to plan, structure and drive your own tasks. You are an open-minded, co-operative and service-minded person. Moreover:

- You have a relevant background from life science
- You have 5+ years of experience with on-site monitoring (site feasibility, selection, initiation, monitoring and closure) of clinical trials according to ICH-GCP
- You have solid experience with submissions of clinical trials to Ethics Committees and Regulatory Authorities
- You have extensive knowledge of relevant national and European guidelines and regulations for clinical trials
- You have the flexibility to travel
- Experience with medical device studies is considered an asset
- You have experience from the CRO industry, the pharmaceutical industry and/or biotech companies

- You are a dedicated team player with a high-quality mind-set, meet your deadlines and know how to prioritise between different tasks in a dynamic environment which requires a high degree of flexibility
- You have good communication skills – and of course, you speak and write English and Danish effortlessly

At Larix, we look forward to welcoming you on board - we offer an attractive salary package including pension scheme, and the opportunity to work from home from time to time. You are welcome to contact Lene Eskildsen, Vice President QA &HR at lhe@larixcro.com or call for further information: +45 23 72 88 63. You can also visit our website at www.larixcro.com

To apply for this position, please forward your application and CV to info@larixcro.com no later than 14 March 2019. Applications will be handled in the order they arrive.