# 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?**
If so, Larix Clinical Operations is now seeking an experienced Clinical Trial Manager.

## What you will do

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

Your main tasks will include:

- Perform and handle all aspects of trial & site management of sponsor projects
- Lead the trial team by driving trial activities to ensure execution and delivery according to agreed timelines, cost, and at high quality
- Handle all aspects of trial oversight
- Perform and handle trial management/project management of small and larger studies both national and international
- Prepare or contribute to the update of SOPs and procedures within Clinical Operations
- Support junior colleagues
- Work in compliance with, ICH-GCP, national and international regulations and standards as well as Larix' quality system

## Who you are

We are looking for an experienced 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
- Strong project management skills
- You have experience with monitoring of clinical trials according to ICH-GCP
- Advantage if you have experience with submissions of clinical trials to Ethics Committees and Regulatory Authorities, but not a must
- You have extensive knowledge of relevant national and European guidelines and regulations for clinical trials
- 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 priorities between different tasks in a dynamic environment which requires a high degree of flexibility
- You have a quality mindset and able to prioritize your work in a fast paced and changing environment
- You have good communication skills – and of course, you speak and write English and Danish effortlessly

## Are you interested?

We would love to hear from you. To apply for this position, please forward your application and CV to **job@larixcro.com** or by applying here on **LinkedIn** using EasyApply. Applications will be handled by our Business Manager in the order they arrive.

Please note that vacancies will be filled on a rolling basis after opening and this job posting is also for our candidate pool for future job openings. Therefore, we highly recommend you submit your application as early as possible to be considered for the opportunity of your choice.

## Who is Larix?

Larix A/S is a Nordic Contract Research Organisation (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. We also have a smaller office at Medicon Village in Lund and a local presence in Oslo and Helsinki. Larix is a part of Alten Group and we work closely with our sister company (Aixial), which gives us all the advantages of being a small, agile and flexible company combined with support from a larger company when it comes to e.g. systems and resources.

At Larix, we maintain a friendly atmosphere – we think having fun while working is important, and we prioritize creating a work environment with a healthy work-life balance. We are a relatively small company with approximately 70 employees, and we have all the benefits of being able to collaborate across functions, learn from each other and follow clinical study processes from start to finish.