Senior /Principal Medical Writer

Are you an experienced medical writer looking for a new challenge in a dynamic and responsible position? Would you enjoy working with different customers and projects either in-house or based at the client's office?
If so, Larix offers you the opportunity to work in a stimulating environment with the possibility to influence your work and responsibilities. We are now seeking a Senior or Principal Medical Writer to join our team in Herlev, Denmark or Lund, Sweden.

What you will do

Medical writing at Larix involves working on a range of different projects, some only concerning the Medical Writing department while others are full-service projects, i.e., in close collaboration with the other departments at Larix, such as Data Management, Statistics, Programming and Clinical Operations. Some tasks are carried out in-house at the Larix office, while others are completed at the client's location or remotely. Thus, you should also be prepared to work at a client's office in the Copenhagen/Lund area from time to time.

At Larix, we work with a wide range of clients spanning from the smallest biotech companies to the largest pharma companies. Therefore, we get to work with a wide range of therapeutic areas. Furthermore, the types of medical writing tasks are diverse comprising regulatory tasks as well as publications, medical device projects as well as medicinal products. Thus, the medical writer role at Larix is broad, and we expect that you will see this as an advantage.

We have a number of upcoming projects and need an experienced medical writer to join our Medical Writing team. You will join a growing team of enthusiastic and skilled medical writers. As we are a relatively small group (currently six), you will be expected to be actively involved in producing and updating processes and tools to support further development of the Medical Writing function.

Who you are

The preferred candidate will be proactive, flexible, service-minded, focused on high quality and timely delivery and will have:

- At least 5 years of experience as a medical writer of clinical regulatory documents (clinical trial reports, protocols, investigator's brochures)
- A university degree in health or life sciences or equivalent, preferably a PhD
- Experience in the following is considered an advantage:
    - Performing literature searches and reviews
    - Preparing medical device documents according to MDR
    - Preparing abstracts, posters and manuscripts in a pharmaceutical industry setting
- Exceptional ability to plan, drive and coordinate complex medical writing projects in collaboration with many stakeholders

- Broad knowledge of several therapeutic areas as well as a general understanding of regulatory requirements, drug development processes, statistical methods and clinical research concepts
- Ability to work independently in a structured, proactive way with a quality mindset
- Strong communication skills and excellent spoken and written English and Danish (preference)

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 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 Organization (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 an office at Medicon Village in Lund and a local presence in Helsinki. Larix is a part of Alten Group and we work closely with our sister companies (Aixial and CMED), 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 75 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.