



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 Lund, Sweden.

Larix A/S is a Nordic Contract Research Organisation – we offer full-service solutions within the pharmaceutical 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. 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 are a relatively small company with approximately 55 employees, and we have all the benefits of being able to collaborate across borders, across functions, follow clinical trial processes from start to finish, and learn from each other.

You will be part of our Medical Writing group. Currently we have a number of new projects and need a medical writer to join our Medical Writing team, working from our office in Lund. You should also be prepared to work at a client's office in the Copenhagen/Lund area from time to time. The Medical Writing team comprises experienced and enthusiastic employees responsible for a wide variety of documents.

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, considered an advantage:
 - Performing literature searches and reviews
 - Preparing medical device documents according to MEDDEV 2.7.1 rev 4
 - 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

Medical writing at Larix involves working on a range of different projects, some only concerning medical writing, others including full-service functions and close collaboration with other functions within the clinical and medical device areas. Some tasks are carried out in-house at the Larix office, while others are completed as an in-house consultant at the client's location. The tasks are therefore diverse, and we expect that you will see this as an advantage.

You will join an experienced and growing team of medical writers and, as we are a relatively small group, you will be expected to be actively involved in producing and updating processes and tools to support further development of the Medical Writing function.

We look forward to welcoming you on board –we offer a competitive salary package including pension scheme, life and medical cover, ongoing education, and the opportunity to work at home from time to time.

For more information, please contact Angela Stocks, Director, Medical Writing, at ast@larixcro.com, +45 8177 8150
You can also visit our website at www.larixcro.com

To apply for this position, please forward your application and CV to job@larixcro.com. Applications will be handled by our Business Manager in the order they arrive.