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.

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 local presence in Lund, Oslo and Helsinki.

At Larix, we maintain a friendly atmosphere – we believe having fun while working is important. We are a relatively small company with approximately 65 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.

Currently we have a number of new projects and need a medical writer to join our Medical Writing team.

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

- At least 7 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 asset:
    - Preparing abstracts, posters and manuscripts in a pharmaceutical industry setting
    - Preparing medical device documents according to MEDDEV 2.7.1 rev 4
    - Preparing summary documents for submission to regulatory authorities
- 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 info@larixcro.com no later than 16 February 2018. Applications will be handled in the order they arrive.