



## **Senior Clinical Research Associate/Start-up Specialist, Finland**

Are you an experienced Clinical Research Associate with submission experience and do you thrive in a dynamic, inspiring environment? Would you like the chance to work in a smaller company with a smooth decision-making process and valued employee influence?

**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 Denmark and we also have local offices in Sweden, Norway and Vantaa in Finland. We work with all indications and a wide range of customers.

**At Larix, you will be part of the Nordic clinical operations group** - consisting of 12 employees, primarily CRAs, Trial Managers and Trial Administrators, who are working either in-house at Larix or as outsourced consultants. As a Senior Clinical Research Associate, your main tasks will be to manage and monitor sites in Finland from submission to close-out.

**At Larix, we keep 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.

**You are an experienced Clinical Research Associate** – we will get to know you as a proactive and competent person who approaches projects with a team player's spirit. 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 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
- Experience with medical device studies is considered an asset
- You have experience from the CRO industry, the pharmaceutical industry and/or biotech companies
- You have good communication skills – and of course, you speak and write English and Finnish effortlessly

**At Larix, we look forward to welcoming you on board** – you are welcome to contact Dorthe Grønnegaard Mejer, VP Clinical Development, at +45 61 22 17 95 or [dgm@larixcro.com](mailto:dgm@larixcro.com) for more information. You can also visit our website at [www.larixcro.com](http://www.larixcro.com).

Communication ♦ Proactivity ♦ Quality on time

To apply for this position, please forward your application and CV by e-mail to our HR Director Lene Eskildsen at [info@larixcro.com](mailto:info@larixcro.com). Applications will be handled in the order they arrive.