CYTOKI

Clinical Project Leader/ Sr. Clinical Trial Manager – make a difference and join our mission to bring our IL-22 lead program to patients

Do you want to be part of a young organization and a team that is about to embark on an exciting journey to bring our IL-22 lead program to patients?

Cytoki Pharma is a biotechnology company focused on serious diseases. The company is based in Hellerup, Denmark and was founded in 2019 after licensing of a full IL-22 analogue program from Novo Nordisk. We are on a track to start phase 1 for our long-acting lipidated IL-22 in late 2022 enabled by a $45 M series A from a syndicate of strong international investors providing funding all the way to clinical proof-of-concept. To progress our lead candidate, we are looking for an experienced Clinical Project Leader/ Sr. Clinical Trial Manager to manage our clinical trial activities.

Driving the clinical trial activities in a collaborative and team oriented atmosphere

This is a unique opportunity to bring all your knowledge and experience into play in a young organization with a small, experienced and dedicated team. We work together in a transparent and collaborative manner with a high degree of trust and flexibility.

We are looking for a colleague who enjoys the teamwork of a small biotech and has the right competences and experience for delivering our early-stage clinical trials with our long-acting, lipidated IL-22 analogue. In return we offer a lot of freedom for you to fill the role, a pleasant and informal working atmosphere, and competitive compensation. You will be the main driver of the following priorities:

- Planning and managing Ph 1 and Ph 2 clinical trials from start-up to closure to ensure trial delivery to agreed timelines, budget, and quality
- Selecting, managing, and performing oversight of the clinical CROs and other appointed vendors
- Developing (or ensure delivery of) key clinical trial documents and plans
- Forecasting and managing budget and timelines for individual clinical trials

- Ensuring all clinical trial activities and deliverables are inspection ready and compliant with SOPs, ICH/GCP guidelines and regulatory requirements.
- Building and maintaining relationships with sites and KOLs

A proactive, self-driven team player with strong experience from managing clinical trials

We are looking for a colleague with a degree in Health/Life Sciences (e.g. MSc or BSN) and 7+ years of experience managing clinical trials from start-up to closure including experience with phase 1 and multicenter, international phase 2 trials (experience with trials in IBD is preferred but not a prerequisite). We expect that you have extensive knowledge of all aspects of the clinical trial process and documented experience in ICH GCP including risk management and sponsor oversight. From your previous positions you have experience with leading clinical trials through full outsourcing to CROs.

What your colleagues say about you:

- Proactive, self-driven and result-oriented with excellent problem-solving skills and a “can do attitude”
- Organized with great planning skills and attention to detail
- Able to think outside the box and handle (unforeseen) challenges on the way
- A team player with a positive mindset who gets energized by working with others and contribute to a fun and informal atmosphere

Caught your interest?
Please send your application and CV to recruitment@cytokipharma.com no later than 15 August 2022. Please note, that we review applications on an on-going basis and close the process when we find the right match.

If you have any questions, you are very welcome to contact Chief Development Officer Anne Louise Kjølbye at alk@cytokipharma.com or +45 3045 1043.