**Sponsor Oversight Lead**

Are you ready to join an expanding and innovative biopharmaceutical company and play a key part in our clinical development efforts?

In this newly established role, you will get to take on cross-disciplinary challenges and make a real difference in our research and development of new cancer treatments. You will join our experienced Clinical Operations team consisting of Clinical Project Managers and Clinical Project Assistants, currently 11 members. The team is part of Global Clinical and Regulatory Development.

**Welcome to Y-mAbs Therapeutics A/S**

Y-mAbs is a rapidly growing late-stage clinical biopharmaceutical company focused on the development and commercialization of novel, antibody-based therapeutic products for the treatment of cancer. The company has a broad and advanced product pipeline and had the first product approved by FDA in November 2020, which is expected to be launched early this year. Moreover, we have an interesting pipeline with several compound modalities entering clinical development.

We are planning to start several clinical trials in the coming year, and we are looking for a dedicated Sponsor Oversight Lead for our Clinical Operations team. With the mission to become the world leader in developing antibody-based cancer products that address clear unmet needs in pediatric oncology, we cannot wait to welcome you to our team.

As our Sponsor Oversight Lead, you will be responsible for the planning, conduct, and coordination of trial oversight activities with internal and external stakeholders to ensure quality and integrity of the data within and across Y-mAbs clinical trials. You will assist the Clinical Project Managers (CPM) in all operational aspects of one or more international clinical trials with the overall goal of developing new and innovative treatment for both pediatric and adult cancer patients.

We work in close collaboration with CROs and you will be responsible for securing trial deliverables in a proactive fashion, through clear communication and great teamwork at all stages of the trials. You will also ensure proper documentation, record management as well as sponsor engagement and sponsor oversight.

If you find the above interesting, join Y-mAbs and become part of our team in a fast-moving environment that offers a unique combination of scientific insight, entrepreneurship, and exciting challenges! Your work will have a palpable impact on our efforts to develop new and innovative cancer treatments as you will:

- Be an essential member of the clinical team supporting the clinical trial planning, execution, and finalization by performing data review tasks as well as wide communication with international stakeholders and collaborators
- Supporting the Clinical Project Manager (CPM) with identification of investigators, building relationships with investigators/trial sites
- Supporting the CPM in securing timely delivery and review of trial related documents, plans including but not limited to master PI/ICF, training records, and monitoring visit reports
- Visit sites to develop and maintain relationships when necessary, and conduct sponsor site visits to support recruitment and ensure compliance with the protocol, SOPs, ICH GCP, and relevant guidelines
- Ensure trial team members are adequately trained for responsibilities on trial conduct
- Oversight of CROs and other vendors involved in the clinical trials
- Contribute to the continuous improvement of Y-mAbs’ procedures and secure best practices

Make your mark on our great team

We offer you the chance to join a highly efficient team of carefully selected leading generalists at our great office location in the DTU Science Park in Hørsholm. Here, you will get to make a real difference.

Experienced Sponsor Oversight Lead with great problem-solving skills

To succeed in this role, you need good communication abilities as you will share knowledge with colleagues outside your field. You are a generalist clinical developer who can and want to work outside the silo with a service-oriented attitude combined with strong analytical skills and an eye for details. Additionally, you also bring a can-do attitude, and you meet any obstacles with a positive and solution-oriented approach.

Specifically, we imagine that you:

- Are BSc or MSc within the medical, biological, or pharmaceutical science or equivalent
- Have 3+ years’ experience and in-depth knowledge of executing early phase clinical trials
- Have experience in working within oncology and/or rare diseases
- Are structured, organized, and have excellent coordinating skills, while still thriving in an entrepreneurial environment
- Are communicating effortlessly with your stakeholders in- and outside Y-mAbs.
- Are result-oriented and committed to contributing to the overall success of Y-mAbs
- Speak and write English effortlessly
- You may expect to travel for 10-30 days per year

Interested?

If you want to know more about the position, you are welcome to contact Joris Wilms, SVP, Chief Operating Officer at: +45 2521 1647 or Søren Fabricius, Director Clinical Operations at +45 5388 0488.

You can apply for the position by sending an email to HR@ymabs.com no later than 19 February 2021. We will initiate interviews in a rolling manner as applications for the job are received.

Please note that applications must be submitted in English and will be treated confidentially. Mark your applications with position ID number: 1037.

Y-mAbs Therapeutics A/S is a Danish affiliate of Y-mAbs Therapeutics Inc., which is located in New York. Our mission is to discover, develop and deliver novel antibody therapeutics for the treatment of both pediatric and adult cancer patients. Please access the company web site www.ymabs.com for more information regarding the company and our development projects.