Location: Gothenburg

Clinical Development Director - to the exciting biopharma company Alzinova

We are looking for our next team member at the position of Clinical Development Director

Alzinova is an exciting biopharma company that develops a treatment for Alzheimer's disease with the aim of making it possible for Alzheimer's patients to live an independent and active life. We are now searching for a new team member in the role as Clinical Development Director (CDD). The CDD will report to CEO and be part of the management team.

We are looking for a CDD who will provide leadership and direction for the company's pipeline of clinical development programs. The CDD will, in partnership with the Chief Medical Officer (CMO) and Development Project Director, drive effective and quality execution of the allocated clinical programs. In this role, the CDD supports the development of the clinical program strategy in compliance with the overall program goals and regulatory requirements. The CDD responsible to ensure operational delivery of the clinical program as per Standard Operating Procedures (SOPs), International Conference on Harmonisation-Good Clinical Practice (ICHGCP), applicable regulations and agreed company goals and values. The role is covering both operational and strategic aspects of clinical development.

What you'll be doing: Your responsibilities include, but are not limited to:

- Lead clinical projects of Alzinova's drug candidates ALZ-101 and ALZ-201.
- Responsible for clinical development plans, activities, and risk management.
- Manage the development of key clinical documents such as the Clinical Development Plan (CDP), protocol synopses, the clinical sections of the Target Product Profile/Claim (TPP/C) documents and the clinical sections of other regulatory documents.
- Actively lead internal clinical project and cross-company study meetings.
- Responsible for protocol development and study execution to time, budget and quality and ensure appropriate risk management and contingency plans are in place to fulfill the clinical program goals.
- Lead CRO/vendor selection (incl Request for Information (RFI)and Request for proposal (RFP) process and vendor management (vendor oversight).
- Administrative tasks such as documentation, applications, budget follow-up and contracts.
- Actively participate in cross-functional project meetings.
- Identify and participate in relevant and value-creating external collaborations e.g. investigators, vendors and scientific leaders and Key Opinion Leaders (KOLs).
- Ensure internal working procedures and processes are in place and implemented for operational compliance in executing the clinical development program.
- To collaborate with other management team members in developing aligned clinical study strategies for country, site and Key Opinion Leader selection and develop the clinical program level recruitment feasibility.
- To contribute to project publications and abstracts strategy and support the coordination of clinical related publications.

The profile we are looking for:

We are looking for an experienced, ambitious, self-motivated, solution oriented, CDD with high energy and the ability and drive to develop, accomplish and follow up on different clinical projects and tasks. You should have experience in drug development with a preferred clinical background in neurology (preferably Alzheimer's Disease) and solid experiences in clinical trials and health authority and KOL interactions. Experience from immunology is an advantage. You should be open to work in an entrepreneurial and

growing organization, where ownership, collaboration and drive is of greatest importance and have a hands-on approach as well as strategic capability. The scope for the role is broad and requires an interest for working with many different tasks and duties. We value solid relation building skills, collaboration skills, project management skills. A proven track record of working in start-ups or in small biopharma companies or from headquarters, not only big pharma would be an advantage.

Other information about the position

- Travel, both international and domestic will be required, from time to time.
- Office in AstraZeneca BioVenture Hub, Mölndal. Possibility to work occasionally from home.

We offer:

- An open environment with a strong and driven team working together towards our joint goals.
- A dynamic workplace being part of an exciting journey in developing new treatments against Alzheimer's.
- Flexible start date.

Application:
Is this a perfect match for you? Send your application today including resumé and a cover letter to recruitment@alzinova.com. We process all applications on a continuous basis.

About Alzinova

Alzinova was founded by scientists at the University of Gothenburg, Sweden, in 2011 after the invention of a unique technology platform, AβCC. The AβCC technology specifically stabilises disease-driving oligomers of the peptide amyloid-β (Aβ), thus enabling the development of disease-modifying therapeutics against Alzheimers Disease. Alzinova is dedicated to developing novel disease-modifying treatments to enable patients to live an independent life free from Alzheimer's disease. Alzinova's treatment concept focuses specifically on the neurotoxic oligomeric forms of amyloid-beta and is thus completely different from antibody candidates in clinical development. Alzinova's lead candidate, ALZ-101, is in clinical development as a therapeutic vaccine for the treatment of Alzheimer's disease. The company has designed a first-in-human clinical study to evaluate the safety and immunogenicity of ALZ-101 and the results of the study are expected to be available in the second half of 2023. Alzinova's antibody, ALZ-201, is in preclinical development. The project portfolio for the development of disease-modifying treatments is broadened by the company in preparing the antibody so that it can also be taken into the clinical phase.
Alzinova is listed on NASDAQ First North in Stockholm, Sweden (ticker: ALZ).