Clinical Trial Supply Coordinator

Manage the clinical trial supplies of a NASDAQ listed Biotech Company

As Clinical Trial Supply Coordinator at Ascendis Pharma you will be responsible for managing the Clinical Trial Supply in compliance with the regulatory requirements of GMP and GDP**.**

You will work closely with devoted colleagues throughout the organization, who all share the overall goal of developing and bringing new, innovative medicine to patients. The supply chain relies entirely on third party manufacturing, transportation, and warehousing. The Clinical Trial Supply Coordinator reports to the Director of Clinical Trial Supply Management and will be part of a dedicated and highly experienced team.

Your main tasks are:

- Be responsible for all aspects of the Clinical Trial Supply Chain from CMOs to clinical sites for clinical trials from phase 1 to 3
- Planning, sourcing, preparing, surveilling, and facilitating clinical supplies for clinical trials of Ascendis Pharma
- Elaboration of label text, standard operating procedures, and quality documents
- Setting up IWRS for larger clinical trials
- Collaborating with and managing subcontractors and CMOs
- Review of documentation from subcontractors
- Elaboration and review of instructions and manuals used in the conduct of the clinical trials
- Overview of subcontractors
- Playing an active role in communicating and collaborating with internal Ascendis Pharma departments e.g. CMC, Clinical Operations, QA, and the Logistics Manager

Ideally, you hold a Master's degree in Pharmacy or have an education as a Pharmaconomist or the like, and more than 4 years of relevant work experience, preferably from biotech or the pharmaceutical industry.

You must have a minimum of 4 years of experience working in a GMP/GDP regulated environment and in managing clinical trial supplies. You have proven working experience in managing complex projects with several internal and external stakeholders. You have a thorough understanding of regulatory requirements. You are proficient in English and Danish at a professional level, both written and spoken, and you master Microsoft Office. Knowledge of IWRS is an advantage but not a requirement.

You have a positive attitude and possess good communication and cooperation skills, which enable you to establish warm and respectful professional relations internally and externally. You are comfortable in a dynamic environment in which priorities may quickly change. You like working in a highly regulated area. Finally, you have a personal commitment to delivering results and succeed.

Ascendis Pharma offers you an exciting and challenging position in an entrepreneurial and international NASDAQ listed company with a short line of command. You will be involved in activities that are central to Ascendis Pharma's strategy, and you will work with highly skilled and experienced colleagues to advance the company's exciting product pipeline.

Traveling: 10 – 20 days/year.

Place of work: Ascendis Pharma resides in a wonderful office facility in Tuborg Havn in Hellerup.

For more details about the job or the company, please contact Recruiter, Louise Nielsen, UniqueConsult on M +45 29 74 24 24. All applications must be submitted in English and are treated confidentially.

Ascendis Pharma A/S is an international company with offices in Copenhagen, Germany and the US. Ascendis Pharma is building an integrated biopharmaceutical company to advance its pipeline of long-acting prodrug therapies. They employ their proprietary TransCon technology platform to generate therapeutics with best-in-class profiles that address large markets with significant unmet medical needs. Ascendis Pharma has a diversified and balanced high-value product pipeline, including internal programs and partnerships with market leaders.

For further information, please see www.ascendispharma.com