Symeres
MAKING MOLECULES MATTER

Symeres is a leading chemistry CRO in Europe, with over 500 scientists at six locations (Nijmegen, Weert & Groningen in the Netherlands, Prague in the Czech Republic, Oulu in Finland and Södertälje in Sweden). We provide a range of services to support small-molecule drug discovery and development projects for (bio)pharmaceutical companies in the USA, Europe, and Japan. Our key areas of expertise are synthetic chemistry, medicinal chemistry, parallel chemistry, and chemical process R&D for clinical trials. As part of an increasing demand for our services at **our location in Prague CZ** (Běchovice) we are looking for a skilled and enthusiastic

Quality Assurance Specialist - Full-time

Function description

The Quality Assurance Specialist manages the QA aspects of the production of small molecule API´s for clinical trials or for commercial use. He/she writes, reviews, approves and archives documentation for the production and analysis of the products, the equipment, facility and personnel in narrow collaboration with the Head of QA, Head of QC and Head of Production. The activities include the release of raw materials, intermediates and products produced under GMP. He/she will be involved in the investigation of deviations during production, out of specification observations, change control procedures, risk analysis and CAPA management. Furthermore He/she will be part of the continuous improvement of our Quality Management System and is involved with the Quality Audits performed by customers and regulatory bodies and the qualification and auditing of sub-contracting laboratories and service providers.

Key requirements

For this function we are looking for an enthusiastic candidate who meets the following requirements:

- University education in analytical chemistry, organic chemistry or a related discipline.
- At least 5 years' experience in the pharmaceutical industry in a related position. Experience in early development of small molecules is an advantage.
- Strong knowledgebase of GMP-requirements for small molecule API's. Experience with investigational products (ICH Q7, chapter 19) is a preference.
- Experience in the qualification and validation of facilities, equipment, methods and processes.
- Technical type of thinking, able to solve problems at the interface of QA and science and technology.
- Knowledge of IT/CSV and the related QA requirements.
- Good command of the English and the Czech language.
- Advanced user of a Personal Computer (Word, Excel, Outlook, technical software).
- Good communicative and organizational skills. Team player.
- Reliable, accurate and accountable personality.

Job offers

An exciting position in a dynamic organization with an attractive remuneration package and opportunities for learning and development.

Application

Are you interested in this position and do you meet the job requirements? Please send an email message with your CV and motivation letter before April 15th, 2022 to our HR Manager Thea Scherpenborg (hrm.hrm@symeres.com).

Further information on the specifics of this position can be obtained from Renata Čichoňová, Head of QA (renata.cichonova@symeres.com). This vacancy is open for EU residents only.

More information about the Symeres organization can be found on our website: www.symeres.com

Acquisition based on the occasion of this advertisement is not appreciated