Symeres
Making Molecules Matter. Together.

Symeres is the leading mid-sized transatlantic contract research organization for drug discovery and development needs. With over 700 highly educated scientists and professionals in six locations in Europe and two in the USA, we offer best-in-class solutions for drug discovery and drug development, from small- to medium-sized molecule hits. We are large enough to matter, and small enough to care. Our services span across early stage hit finding all the way to the delivery of your early clinical drug substance API. Making molecules matter. Together.

Due to fast international growth and 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

The Quality Assurance Specialist manages the QA aspects of the production of small-molecule APIs for clinical trials or for commercial use. They write, review, approve, and archive documentation for the production and analysis of the products, equipment, facility, and personnel in close 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. They will be involved in the investigation of deviations during production, out-of-specification observations, change control procedures, risk analysis, and CAPA management. Furthermore, they will be part of the continuous improvement of our Quality Management System and involved in Quality Audits performed by customers and regulatory bodies and the qualification and auditing of subcontracting laboratories and service providers.

Qualifications

- 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 APIs. Experience with investigational products (ICH Q7, chapter 19) is a preference.
- Experience in the qualification and validation of facilities, equipment, methods, and processes.
- Technically minded, able to solve problems at the interface of QA and science and technology.
- Knowledge of IT/CSV and related QA requirements.
- Good command of the English and Czech languages.
- Advanced user of a personal computer (Word, Excel, Outlook, technical software).
- Good communicative and organizational skills, a team player.
- Reliable, accurate, and accountable personality.

Job offer

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

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Application

Are you interested in this position, and do you meet the job requirements? Please send your CV and motivation letter to our HR Manager, Dries Sauren (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 this advertisement is not appreciated