Symeres
MAKING MOLECULES MATTER

S ymeres is one of the leading chemistry CROs in Europe, with over 500 scientists at six locations (Nijmegen, Weert, and 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 biopharmaceutical 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. All these fields of expertise are supported by highly experienced and well-equipped analytical teams.

At our location in Weert (NL), we are currently looking for a

QA Officer Quality Management System (QMS)

Full-time

The QA Officer QMS effectively performs activities to ensure compliance with applicable current Good Manufacturing Practice (GMP) regulations, corporate policies, and site Standard Operating Procedures (SOP).

Key job responsibilities:

- Provide support and guidance on qualification/calibration of facilities, production/analytical equipment and utilities related to the cGMP manufacture of active pharmaceutical ingredients (APIs).
- Review and approve qualification/calibration documentation of production and analytical equipment.
- Review and approve risk assessments related to production and analytical equipment qualification/calibration.
- Review and approve change controls, deviations, corrective and preventive actions (CAPAs) and effectiveness checks.
- Support writing/revising standard operating procedures (SOP) and work instructions (WI).

Qualifications

- At least HBO level or equivalent.
- At least 3 year working experience in an R&D function/chemical industry (*e.g.*, analysis, production).
- High level of initiative, flexibility, and problem solving.
- Awareness of the relevant procedures for their function in terms of quality, safety, health, and environment.
- Good written and oral communication skills (at least in Dutch and English).
- Proficiency with Microsoft Office (Word, Excel, PowerPoint).
- Working effectively with engineering and operational teams.
- Working knowledge of cGMP and computer system validation (CSV) in a pharmaceutical/regulated environment is preferred.

Job offer

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

Application

Please send an email with your CV and motivation letter to our HR Manager: karen.storms@symeres.com
For substantive questions, please contact our Director of Quality: chantal.rademaekers@symeres.com.
More information about the Symeres organization can be found on our website: www.symeres.com .

Acquisition based on this advertisement is not appreciated.