Symeres is the leading mid-sized transatlantic contract research organization for drug discovery and development needs. With over 600 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 early clinical drug substance API. Making molecules matter. Together.

To increase our capacity, the Quality Management System (QMS) team at our Weert site is strengthening its team and is looking for an experienced and enthusiastic:

QA Officer Equipment Qualification, laboratory equipment

Full-time

The QA Officer Equipment Qualification effectively performs activities to ensure compliance with applicable current Good Manufacturing Practice (cGMP) regulations, corporate policies, and site standard operating procedures.

Key job responsibilities:

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

Qualifications

- At least HBO level or equivalent.
- At least 3 years' working experience in a QA function.
- High level of initiative, flexibility, and problem solving.
- Awareness of the relevant procedures for the 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 Information System team.
- Working knowledge of cGMP guidelines and computer system validation (CSV) requirements 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 to: recruitment@symeres.comFor substantive questions, please contact our Director 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.

www.symeres.com