Analytical Lab (m/w/d)
Head of Quality Control (QC)
Analytical Development

Gen-Plus GmbH & Co KG
Staffelseestraße 6 · 81477 München

Tel.: +49 (0) 89 - 78 01 79 - 40
Fax: +49 (0) 89 - 78 01 79 - 44

Gen-Plus is a small medium-sized, privately owned company. Its aim is to discover innovations for our contract partners. We design formulation and technology concepts for the pharmaceutical industry. This includes solid and semisolid dosage forms as well as patches and thin films. The range is from the early idea to the medicinal product under GMP conditions.

Beside the conventional pharmaceutical technologies, Gen-Plus is investing in innovative technologies in area of personalized medicine.

To expand our research team, we are looking for a Head of Quality Control and Analytical Development and Validation Team.

Tasks and Responsibilities

- Lead of QC and analytical team
- Organisation of lab operations incl. equipment and lab infrastructure
- Development and validation of analytical methods suitable testing of APIs, excipients, intermediates and drug product (solids, semi-solids and liquids, and TTS) under cGMP and in R&D
- Planning, organisation and overseeing of analytical and QC lab activities (method development, validation, stability studies, pyhsico-chemical characterization of APIs, reference materials, method transfers, qualification of lab equipment)
- Review and approval of GMP relevant documentation (SOPs, validation plans, reports, methods, risk analysis, CAPAs, deviations, …)
- Approval of specifications, sampling procedures and testing procedures according to § 14 Abs 1 and ensuring testing comformity
- Testing and release of APIs, excipients, packaging materials, intermediates and drug products
- Selection of partner labs for external testings
- Participation in internal and external audits and inspections
- Budget and investment planning, ensuring permanent training of lab staff

Your Profile

- Master or PhD in a relevant discipline (e.g. pharmacy, analytical sciences, life science) or professional training with at least 5 years' experience in QC and analytical method development and valdidation, of which at least 3 years of team lead experience
- Knowledge in cGMP guidelines, experience in audits
- Problem solving attitude
- High flexibility, team leader and team player
- Positive, communicative and open personality helps you to understand other opinions and contributes to productive project discussions with internal and customer project teams
- Sound knowledge of MS Office applications and common chromatography software

We offer

- Flexible working hours
- Personal development
- Attractive company health management
- Committed team
- Attractive location
- Opportunity to work independently

Your Application

Are you interested in shaping the future with us?

Please send your application documents to

Barbara Schweikl
Gen-Plus GmbH & Co KG
Staffelseestraße 6
81477 München
089 7801 7940
Email: Barbara.Schweikl@gen-plus.de