End-to-end service concept

Our services cover the entire value chain of medicinal products with small and large molecules

Drug development & technologies

- Solid, semi-solid and liquid formulations
- Parenteral, nasal, oral and topical applications
- Manufacture of clinical trial and small batches
- Secondary packaging and labelling
- Handling of high potent drugs
- High-throughput screening for poorly soluble APIs
- Micro- and nanosuspensions

Analytical development & services

- Method development and validation
- Purity testing including structure elucidation
- Release testing, Stress testing & ICH stability studies
- Extractable & leachable studies
- Root cause analysis of manufacturing & production issues (troubleshooting)
- Latest test methods in testing drug formulations, APIs, excipients, biologics

Pharmacovigilance services

- Pharmacovigilance services
- Automated literature search and assessment
- AI supported vigilance processes
- Pharmacovigilance system
- EU-QPPV, graduated plan officer, information officer

QP services & regulatory consulting

- QP batch release
- GxP services and audits
- Application for manufacturing, import and wholesale authorisation
- Implementation and maintenance of quality assurance systems
- Regulatory services
- QM officer, wholesale representative

Key Facts

- Full service provider for pharma and biotech in development and life-cycle
- 3 sites – more than 2,000 m² lab space and 600 m² clean room space
- EU-GMP, FDA, manufacturing licence
- 150 employees

Frankfurt Site

- Drug development
- Semi-solid and liquid formulations (including parenteral, intranasal)
- Small batches and clinical trial manufacturing

Ruelzheim Site

- Laboratory services
- Pharmacovigilance
- QP services & regulatory consulting

Appenweier Site

- Drug development
- Solid formulations
- Small batches and clinical trial manufacturing