LAB EXPERTS AT YOUR SIDE

Over twentyfive years of experience
About us

For our clients in the pharmaceutical, biotechnology and cosmetics industries as well as for manufacturers of medical devices and novel foods SYNLAB Pharma offers a broad portfolio of analytical methods for clinical and preclinical studies from a single source.

We support you from drug discovery to release of the final product and offer you a global logistics network.

All services are monitored to the highest standards. Depending on the requirements we work according to DIN/EN ISO/IEC 17025/15189, GMP, GLP, GCLP or GCP.

Take advantage of our extensive experience from more than 3,000 clinical studies and projects.

As part of the SYNLAB Group, we are part of a worldwide network that is present in more than 40 countries on four continents and holds leading positions in most markets. SYNLAB offers a full range of innovative and reliable medical diagnostics for patients, practicing doctors, clinics and the pharmaceutical industry.

Over 20,000 employees contribute every day to the Group’s success across different geographies. SYNLAB carries out about 450 million laboratory tests per year, achieving sales revenue of about €2 billion.

Europe’s largest laboratory service provider.

From our 6 pharma lab locations in the heart of Europe, we offer you worldwide solutions in high and reliable quality. By networking our specialized laboratories, we can offer you all services from a single source, even for complex requirements.
Foods for Special Medical Purposes (FSMPs)

SYNLAB Pharma is the GLP, GMP and GCP lab expert during R&D and manufacturing of novel foods in accordance with the European guideline (1223/2009). Analytical services include: bioanalytical testing, hygiene, microbiology, toxicology and clinical trial services.

What we offer

Biopharmaceuticals
SYNLAB Pharma develops and implements strategies for the analysis of innovative products, biosimilars and biobetters. Only to name some areas of expertise: PK and PD, biomarkers and immunogenicity including assay development, optimization and validation. In addition, the drug production process is supported by our broad experience in running bioassays for release and stability testing.

Release & Stability
Another core service of SYNLAB Pharma is stability studies in accordance with ICH Q1A(R2), including consultancy, planning, storage and analysis. SYNLAB Pharma supports all stages of successful drug product registration: Long-term, intermediate and accelerated stability; ongoing and follow-up stability studies, stress testing (including photo-stability testing), and compatibility studies.

Cosmetics
SYNLAB Pharma supports all stages of the production process of cosmetics. We offer consultancy for analytics and quality control, physico-chemical methods, stability studies under defined climate conditions, microbiological and in-use stability tests. Quality controls are performed according to EP, USP or customer requirements.

Medical Devices
The development of medical devices requires specialized lab testing in order to cope with the increasing regulatory requirements. SYNLAB Pharma has the expertise in running biocompatibility testing (ISO 10993), leachable & extractable studies, toxicological and microbiological tests of medical devices for biological evaluation in development, registration and production.

Pharmaceuticals
SYNLAB Pharma provides a wide range of analytical services supporting our customers from drug discovery through preclinical and clinical studies to release and manufacturing of the final drug product. We have over 25 years of experience in supporting the pharmaceutical industry, from the measurement of single assays to the execution of complex and demanding projects across the entire life cycle of a pharmaceutical product.
Covering the whole lifecycle of your product

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Global Quality Management

SYNLAB Pharma offers a unique range of laboratory services always working in accordance with the highest available standards such as DIN/EN ISO/IEC 17025/15189, GMP, GLP, GCLP or GCP.

Quality implementation
In addition to our own comprehensive quality procedures, SYNLAB Pharma undergoes periodic audits from customers as well as from national and international regulatory authorities including DAkkS, FDA and Swissmedic. This ensures an ongoing review of all methods, procedures and results with continuous improvement of all established processes.

International quality system
SYNLAB Pharma guarantees that all services rendered are provided by continuously trained professionals who know the importance of each individual sample received. Periodic audits of all laboratory facilities are performed by our QA team. Rigorous cross-validation ensures comparability of global lab results.

Highest IT standards
Our IT platforms enable the dispatch of lab reports all over the world to display identical layout, content and flagging criteria. Secure online access is standard.

Central Lab Services

SYNLAB Pharma pairs the technology and the advantage of economies-of-scale of our reference laboratory with the flexibility and responsiveness of our clinical trials division to deliver the flexibility and high quality sponsors deserve. Global reach is available via our laboratory network with facilities in Europe, North and South America, South Africa, Near and Middle East, Australia, Asia and India. Periodic audits and cross-validations ensure data comparability through test correlations and evaluation of data accuracy and precision. Our dedicated project teams supervise and coordinate all global activities. This results in highly reliable shipping logistics and clean electronic data transfers.