EXPERTS AT YOUR SIDE
Over twenty-five years of experience
About us

SYNLAB Pharma provides laboratory services for our clients in the pharmaceutical, biotechnology and cosmetics industries as well as for manufacturers of medical devices and functional foods. Our broad portfolio offers analytical methods for clinical and preclinical studies as well as for quality control 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: GLP, GCLP, GCP, GMP, DIN EN ISO/IEC 17025/15189.

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

As part of the SYNLAB Group, we are a member 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. Through our vast network of specialized laboratories, we can offer you all services from a single source, even for complex requirements.
What we offer

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.

Technical Regulatory & Regulatory Affairs
Regulatory project design for early and late stage development including consultation of regulatory agencies (scientific advice & clinical trial applications).

Central Lab
SYNLAB Pharma is providing Central Lab Services since more than 25 years covering all relevant global regions incl. China. Lab results are comparable worldwide and lab report layouts are harmonized for all study sites involved. Our clients benefit from highly dedicated project management teams and cost-efficient logistics solutions.

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 approval: Long-term, intermediate and accelerated stability; ongoing and follow-up stability studies; stress testing (including photostability testing), and compatibility studies.

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.

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.

Biologics
SYNLAB Pharma develops and implements strategies for the analysis of therapeutic proteins, Advanced Therapy Medicinal Products (ATMP) and biosimilars. The drug production process is supported by our broad experience in running bioassays and physico-chemical analytics under GMP. Further area of expertise are: PK and PD, biomarkers and immunogenicity testing including assay development, optimization and validation.

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

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Covering the whole lifecycle of your product

<table>
<thead>
<tr>
<th>Discovery</th>
<th>Pre-clinical Development</th>
<th>Clinical Development</th>
<th>Market Phase</th>
</tr>
</thead>
<tbody>
<tr>
<td><strong>DRUG PRODUCT / SUBSTANCE</strong></td>
<td>Regulatory Compliance: Quality Control, Characterization, Release</td>
<td><strong>Analytical Development</strong>: Technical Evaluation</td>
<td>Global Central Lab Services: Global Logistics</td>
</tr>
<tr>
<td></td>
<td></td>
<td>Development of New Tests</td>
<td>Supply</td>
</tr>
<tr>
<td></td>
<td></td>
<td>Validation (EMA/FDA)</td>
<td>Visit Kits</td>
</tr>
<tr>
<td></td>
<td></td>
<td>Method Transfer from or to Client</td>
<td>Storage</td>
</tr>
<tr>
<td><strong>SAFETY / STRATIFICATION / EFFICACY</strong></td>
<td>Bioanalytics: TK</td>
<td><strong>Clinical Monitoring</strong>: Safety</td>
<td>Global Central Lab Services: Global Logistics</td>
</tr>
<tr>
<td></td>
<td>PK</td>
<td></td>
<td>Supply</td>
</tr>
<tr>
<td></td>
<td>Metabolites</td>
<td></td>
<td>Visit Kits</td>
</tr>
<tr>
<td></td>
<td>PD</td>
<td></td>
<td>Storage</td>
</tr>
<tr>
<td></td>
<td>Immunogenicity</td>
<td></td>
<td>Project Management</td>
</tr>
<tr>
<td></td>
<td>Bioavailability</td>
<td></td>
<td>Data Management</td>
</tr>
<tr>
<td></td>
<td>Bioequivalence</td>
<td></td>
<td></td>
</tr>
<tr>
<td></td>
<td>Biomarkers</td>
<td></td>
<td></td>
</tr>
<tr>
<td><strong>FULL SERVICE FOR CLINICAL TRIALS</strong></td>
<td>Genomics: Sequencing</td>
<td></td>
<td></td>
</tr>
<tr>
<td></td>
<td>Genotyping</td>
<td></td>
<td></td>
</tr>
<tr>
<td></td>
<td>Pharmacogenomics</td>
<td></td>
<td></td>
</tr>
<tr>
<td><strong>DIAGNOSTICS</strong></td>
<td></td>
<td></td>
<td>Clinical Monitoring: Safety</td>
</tr>
<tr>
<td></td>
<td></td>
<td></td>
<td>Efficacy</td>
</tr>
<tr>
<td></td>
<td></td>
<td></td>
<td>Therapeutic Drug Monitoring</td>
</tr>
<tr>
<td></td>
<td></td>
<td></td>
<td>Patient Stratification</td>
</tr>
<tr>
<td></td>
<td></td>
<td></td>
<td>Adverse Events</td>
</tr>
</tbody>
</table>
Global Quality Management

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Quality implementation
In addition to our own comprehensive quality procedures, SYNLAB Pharma undergoes periodic audits from customers as well as inspections from national and international regulatory authorities including FDA, Swissmedic and German competent authorities. In addition all our clinical laboratories are accredited by national accreditation services (DAkkS, SAS). This ensures an ongoing review of all methods, procedures and results with continuous improvement of all established processes.

International quality system
SYNILAB 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

SYNILAB 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.