Pharmbiotest Poland LTD
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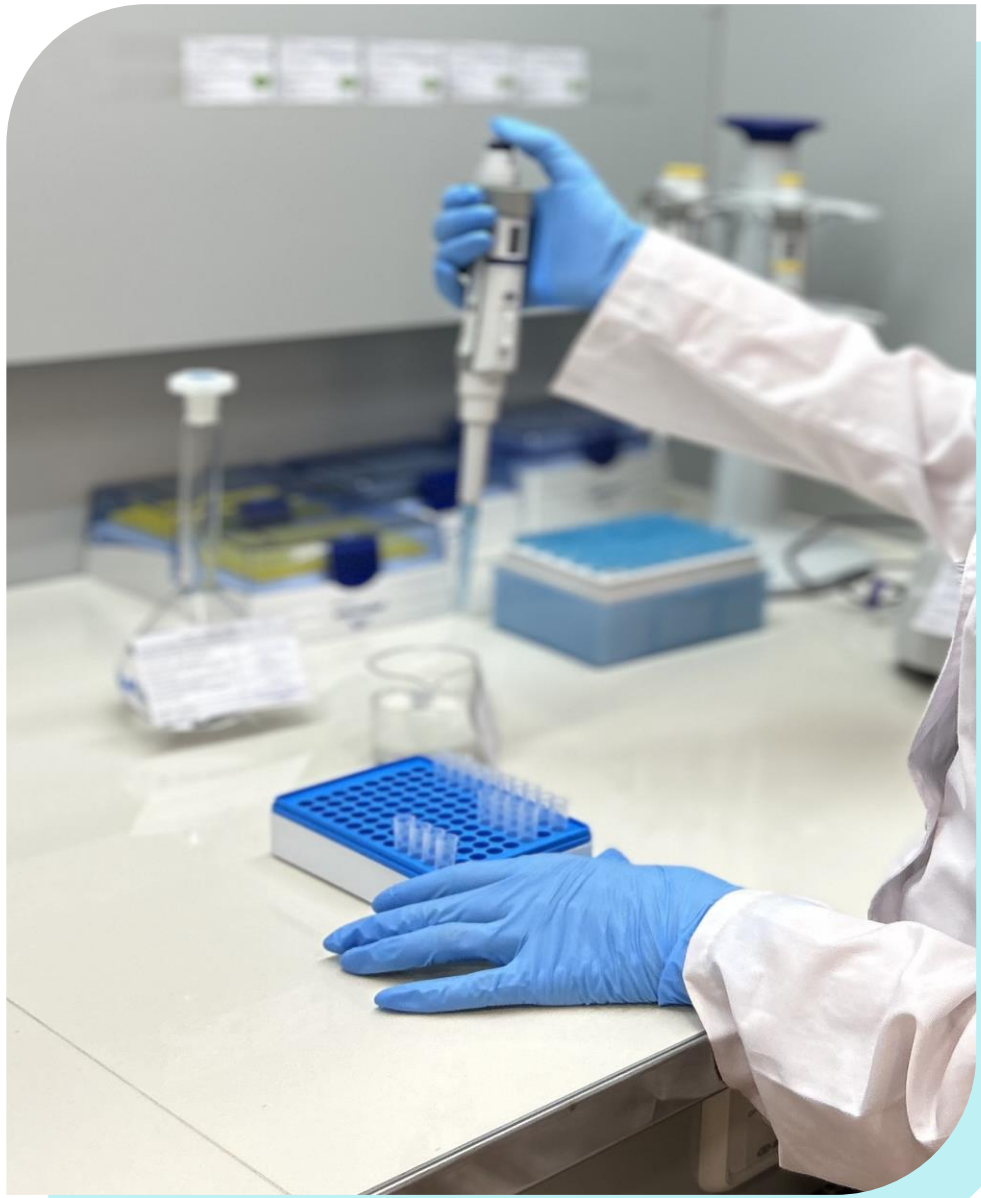
PHARMBIOTEST Poland LTD

Contract Research Organization

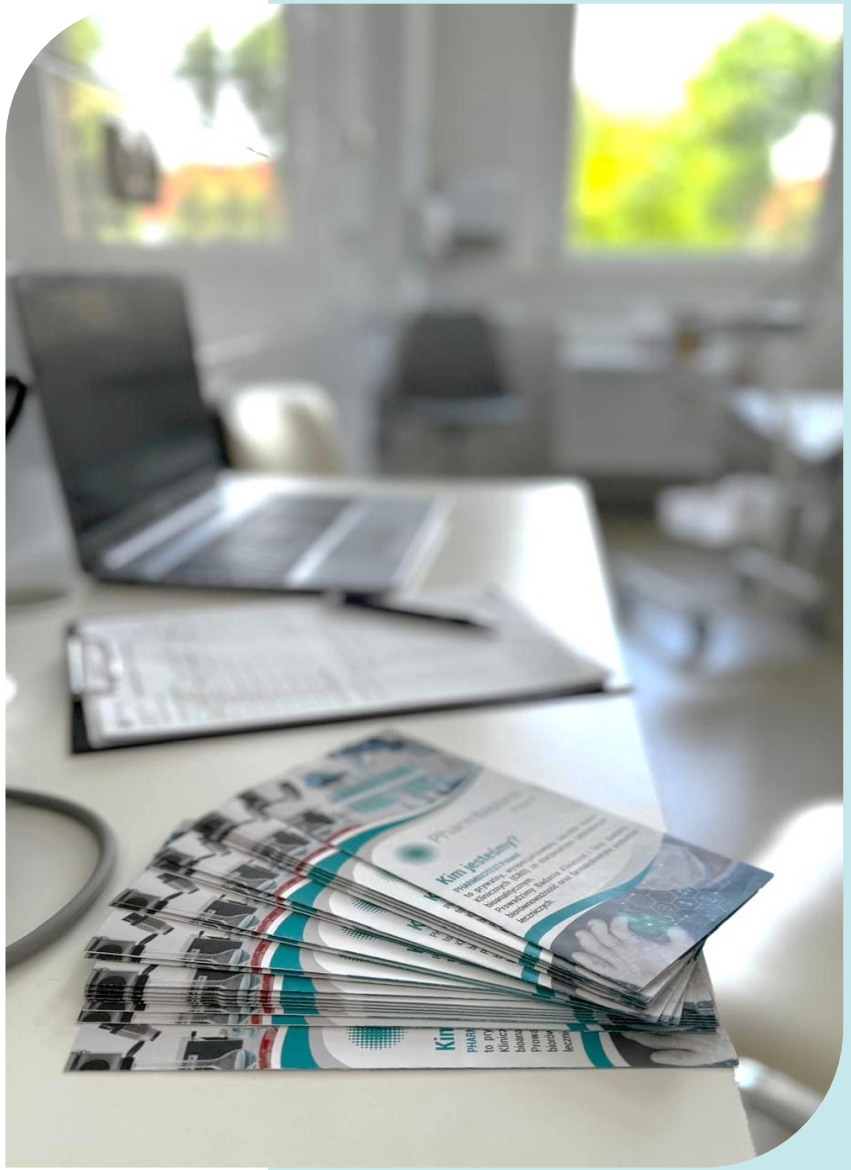
End-to-End Solutions: From R&D to Clinical Trials and Regulatory Submission



Pharmbiotest Poland

Who we are

PHARMBIOTEST Poland is a contract research organization (CRO) with its own R&D, clinical and bioanalytical facilities, providing a broad range of development and research services for pharmaceutical, generic drug and healthcare companies. Our headquarters are in Grudziadz, Poland.



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Our services

PHARMBIOTEST Poland offers comprehensive clinical research services for Phase 1 clinical trials, specializing in pharmacokinetic, bioavailability, and bioequivalence studies. In addition, we conduct R&D activities, including the development of generic drugs, offering expertise from early formulation to clinical trial.

R&D OF GENERIC DRUGS

BA/BE STUDY

PHASE I STUDY

CLINICAL TRIAL DOCUMENTS DEVELOPMENT

BIOSTATISTICS & BIOANALYTICAL SERVICES

THERAPEUTIC DRUG MONITORING

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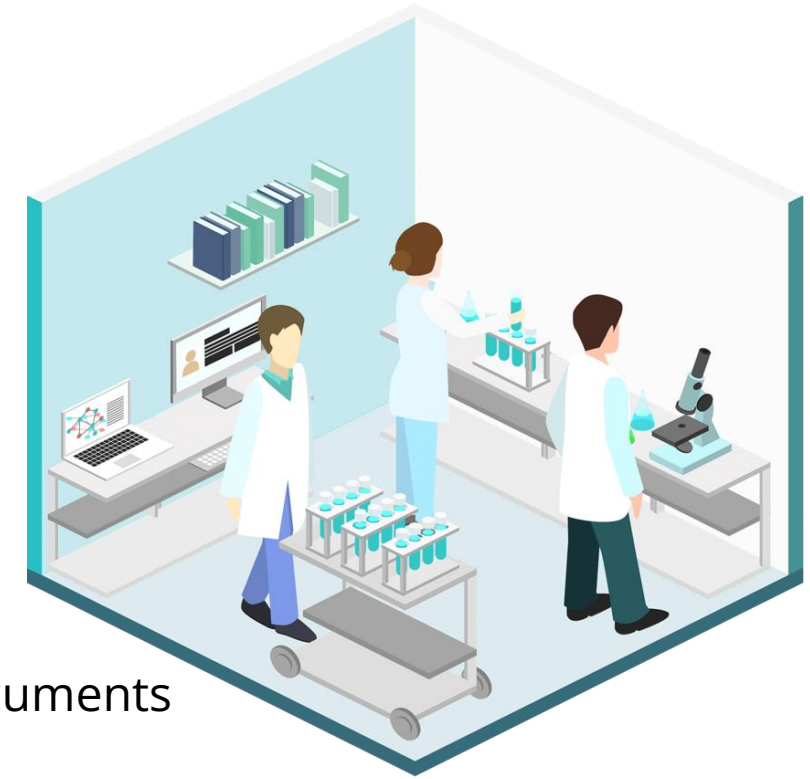
End-to-End Solution

Our company offers a full-service package, covering the entire process from drug development (R&D) to clinical trials and bioanalytical studies, including BA/BE/PK evaluations, study reporting, and regulatory document submission to CTIS. Additionally, we provide standalone services for formulation development, clinical trial execution, and bioanalytical analysis, tailored to our clients' needs.

- R&D and Formulation Development
- Physicochemical Characterization
- Study Design and Planning
- Clinical Trials (BA/BE/PK)
- Bioanalytical Analysis
- Data Processing and Analysis
- Preparation of Regulatory Documents



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Our R&D Services

We offer formulation development and the continued development of existing formulations.

Research and development of innovative dosage forms and drug combinations.

Formulation of generic and hybrid pharmaceutical products, including composition, dosage form selection, and production technology.

Establishing quality requirements for active pharmaceutical ingredients (APIs), excipients, and final drug products.

Evaluating excipient interchangeability across various dosage forms.

Scaling and implementing production technologies for industrial manufacturing.

Our advantages

- Wide experience in developing cardiovascular and neurological drugs
- Over 30 successful Phase I clinical trials
- A full range of Phase I trials (turnkey studies)
- Clinical unit & Bioanalytical laboratory (GLP- certified) in one facility
- Any complexity of bioanalytical methods
- Staff trained in accordance with GLP and certified in GCP
- Wide experience in difficult designs (full-replicate, two-stage etc.)

We look forward to partnering with you!

Our contact information:

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