

COMPANY PROFILE



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History and Mission

PHARMALOG Institut für klinische Forschung GmbH, your professional partner for clinical studies. Our dedicated and experienced team of academics undertakes phase I, phase II-IV trials for you. We work for the pharmaceutical and biotechnology industries and for manufacturers of medical devices, food supplements and cosmetics. We are also happy to implement post-marketing and pharmacoeconomic studies for you. We undertake both complete research projects and special tailored services in all major indications. PHARMALOG: Transparent – Accurate – Flexible – Friendly

Highlights

Our customers/clients

appreciate our flexible and accurate services and our customised designs as well as our friendly staff and our transparency in every phase of the project.

Our company

- is experienced in managing studies in all major indications – as well as in paediatrics and phyto-therapeutics
- offers well-trained and experienced employees who are able to work nationally as well as in Western Europe and CEE
- provides a large investigator pool within Germany, Eastern and Western Europe



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BVMA member since 2011
Audits passed in 2011, 2014, 2017
and 2020



PHARMALOG®
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Services

From audit to report

As a full-service CRO we are in a position to provide you with effective and efficient support from the very start. We take a suitably flexible approach to each individual study to meet its specific requirements. We also offer tailored services, for example, assistance for specific project phases or consulting services.

Study preparation

- Study design, clinical and biostatistical concept
- Medical writing
- Submissions (ethics committees)
- CRF Development (paper or eDC)

Study management

- Study center selection
- Clinical project management, Initiation and training of the centers
- Monitoring
- Data input (GAMP 5 validated systems)

Study evaluation

- Statistical analysis
- Data validation
- Statistical evaluation
- Query management
- Preparation of an integrated study report according to ICH GCP
- Publication

Accompanying Services

- Audits
- Trainings
- Drug Safety in Clinical Trials