■ COMPANY PROFILE



Prof. Dr. Leona Plum-Mörschel Chief Executive Officer info@profil.com



Sascha Heckermann Chief Executive Officer info@profil.com

History and Mission

Profil, founded in 1999, is the world's leading full-service CRO for metabolic research and focuses on early phase clinical trials in diabetes, obesity and NAFLD/NASH. This specialization, together with our scientific excellence and regulatory expertise, enables us to offer an unrivalled service and makes us the scientific partner who supports our clients to design the very best clinical trials and get the most out of their compound development.

Highlights

Sophisticated methodology

We provide a vast portfolio of methods and implement advanced techniques, typically only found at large academic centres.

Rapid subject recruitment

Due to the detailed information in our subject database with thousands of difficult-to-recruit subjects, **Profil** is able to offer the best-in-class recruitment speed.

Scientific excellence

With hundreds of scientific papers and oral presentations, **Profil** is the expert in its field.

IMP supply services

Profil has the capabilities to manufacture small batches of IMP for clinical trials.





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BVMA Member since 2022 Audit passed in 2022





www.profil.com

Services

Clinical unit with 65+ beds

- Phase Lunit
- In-patient and out-patient clinical research units
- Largest glucose clamp centre worldwide
- Fully automated glucose clamp system, ClampArt®

Specialized in metabolic disorders

- Diabetes
- Obesity
- NAFLD/NASH
- Related complications

Full service for early phase clinical trials

- Study design and protocol development
- Handling regulatory affairs
- In-house study execution
- Project management
- Subject recruitment
- Data management
- Statistics

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Scientific writing

Ethno-bridging studies

- Japanese subjects
- Chinese subjects
- Caucasian subjects

Consulting

- Clinical trial design and optimization
- Scientific advisory service

IMP and GMP clinical supply services

- Comparator sourcing (German, EU and US market)
- Manufacturing of small batches of IMP for clinical trials
- Sterile and non-sterile dosage forms
- Sterile injectables and infusions
- Liquid oral dosage forms and capsules
- Filling, packaging, blinding and labeling



