camovis.

FOR SPONSOR

POWERING TRIAL SUCCESS

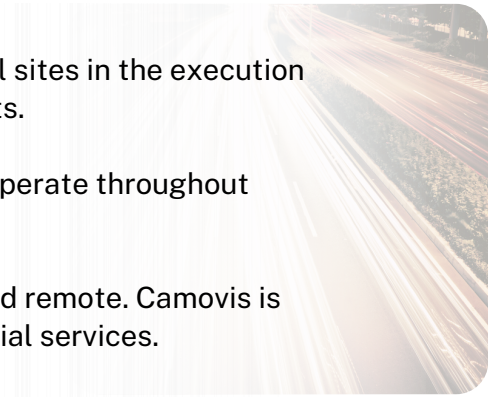
Your Partner for Sponsor Services

Why together

Clinical trials are complex, full of interdependencies, and sponsors carry overall responsibility for quality, compliance, data integrity, and the success of the study. Even when partnering with experienced CROs, one critical factor remains: the true outcome of a study is determined at the site. It is often here that delays, inefficiencies, and performance variability occur, jeopardizing project timelines, increasing costs, and raising risk.

Camovis closes this operational gap. We stabilize site performance and streamline critical processes — from study start through patient recruitment to close-out. Precisely where trials need to succeed and impact is greatest: we act directly on patient flow and site workflows.

We enable. We speed up.



- **Since 2014,** we have been supporting sponsors, CROs, and trial sites in the execution of clinical trials according to the latest regulatory requirements.
- **Over 350** carefully selected Camovis research professionals operate throughout Germany, Switzerland, Austria and beyond.
- **We are dedicated to delivering.** Anytime, anywhere, on-site and remote. Camovis is your partner for high-performance, patient-centered clinical trial services.

Our Commitment — We plug in

“Camovis complements existing operational structures and accelerates study execution. We are committed to be a trusted partner for Sponsors and their teams to deliver clinical trials smooth and successfully.”

Carolin Kurth I CEO I Camovis

- Close stakeholder collaboration
- Deep integration in site operations
- Focus on timing and quality
- Acceleration of operational workflows
- Bridging HR & ressource gaps
- Engaged, agile, and patient-centric

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Camovis Services

We are where clinical trials take place: on-site at the trial centers. At the same time, we support teams in study organization, empower staff through training, and provide consulting when processes or recruitment face obstacles. In this way, we combine operational services, training, management, and consulting to ensure that studies are not only planned but also successfully executed.

Site Coordination: Operational management of study workflows.

Patient Recruitment: Targeted identification, pre-screening, and follow-up.

Clinical Operations: Conduct of study visits, medical examinations, and sample logistics.

Data Handling: Data transfer, eCRF entry, query management, and preparation of archival data.

DCT Services: Home visits, remote support, virtual visits, and patient-centered care.

Project Training: Practical, study-focused training for site teams.

Courses: Webinars, e-learning, and in-house training for study coordinators and study nurses.

Consulting: Site performance enhancement, audit & inspection readiness, and personnel coaching.

Let`s get to work

Camovis GmbH

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Feel free to contact us — we're here and happy to help! :)