CenTrial GmbH

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BVMA member since 2009
Audits passed in 2009, 2012
and 2015

HISTORY AND MISSION

CenTrial was founded in 2000 and is an independent CRO. We have long since expanded our business to industry-sponsored projects and work successfully with national and international clients from the pharma and medical devices industries. We provide high quality services and have been DIN EN ISO 9001 certified in all business segments since 2004. Our quality and reliability are key to our continued success in the field.

Ideen entwickeln –
Neues verwirklichen

develop ideas –
implement
something new

HIGHLIGHTS

In 2015, CenTrial has celebrated its 15th birthday. We can proudly say that the company has achieved a lot and continues to thrive:

Our vibrant CRO team maintain a comprehensive range of services for all aspects of clinical trials and beyond – and are renowned for their expertise and good communication skills, combined with fully GCP compliant processes and technical equipment.

Our continuing education group are offering well-received up-to-date seminars, and our Master of Science »Clinical Research« team have recently welcomed their 12th class of students.

SERVICES

Strategic Planning
- ▷ Consultancy for clinical development of pharmaceuticals and medical devices
- ▷ Networking with clinical and scientific experts

Clinical Operations
- ▷ Recruitment of study sites and Feasibility
- ▷ Development of study protocols and Informed Consent Forms
- ▷ Development and validation of questionnaires
- ▷ Submission to Ethics Committees
- ▷ Submission to Competent Authorities
- ▷ Study management
- ▷ Study registration in accordance with Declaration of Helsinki
- ▷ On-site study assistance
- ▷ Flying study nurse, patient home visits
- ▷ Monitoring including risk-based monitoring
- ▷ Site management
- ▷ Electronic Data Capture (EDC)
- ▷ Data management with fully compliant database systems
- ▷ Biometric and statistical analysis (contract statisticians)
- ▷ Pharmacovigilance and medical device vigilance
- ▷ Development of study reports and regulatory submissions
- ▷ Scientific publications
- ▷ Preparation of documents for the Common Technical Document (CTD)

Continuing Education Group
- ▷ GCP Trainings for medical staff and sponsor personnel
- ▷ Seminars on all aspects of clinical trials for pharmaceutical and medical device companies and for medical institutions / hospitals
- ▷ MSc “Clinical Research” in cooperation with Danube University Krems, Austria
- ▷ Customised inhouse seminars