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



## HISTORY AND MISSION

The company was established in 2000 as the independent contract research organisation of the University Hospital in Tübingen. CenTrial GmbH has been certified in compliance with DIN EN ISO 9001 for all its business segments since 2004. We conduct your clinical trials according to international standards, such as the Good Clinical Practice guidelines (ICH-GCP).



**Ideen entwickeln –  
Neues verwirklichen**

*develop ideas –  
implement  
something new*



## HIGHLIGHTS

A broad spectrum of services:

As a CRO CenTrial GmbH supports both clinical trials on behalf of the pharmaceutical and medical devices industry and Investigator Initiated Trials (IIT).

We support clinical trials in all indications. Our scope of performance includes consulting and services for all types of trials as well as training of study personnel.

Being a part of the University Hospital Tübingen we have direct access to the experts and medical facilities based there.

## SERVICES

### Ready to help and advise you

- ▷ Consulting in the development of pharmaceuticals and medical devices
- ▷ Cooperation with clinical and scientific partners

### Professional implementation

- ▷ Recruitment of study sites
- ▷ Feasibility
- ▷ Protocol development
- ▷ Medical writing
- ▷ Submission to Ethics Committee
- ▷ Submission to Competent Authorities
- ▷ Study management
- ▷ On-site study assistance
- ▷ Flying study nurse
- ▷ Collection and recording of data
- ▷ Monitoring
- ▷ Site management
- ▷ Electronic Data Capture (EDC)
- ▷ Pharmacovigilance
- ▷ Medical Device Vigilance
- ▷ Data management
- ▷ Biometric and statistical analysis carried out by collaborating partners
- ▷ Supervision of study sites
- ▷ Reports and publications

### Qualification of personnel involved in the trial

- ▷ GCP training of study nurses and investigators
- ▷ Master of Science Clinical Research
- ▷ In-house courses