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

CROLLL
individuell › schnell › flexibel

HISTORY AND MISSION

CROLLL is a full service CRO offering design, conduct, monitoring and analysis of clinical trials in cooperation with pharmaceutical and medical devices industry, universities and physicians.
Our first priority is the rapid and straightforward accomplishment of complete or partial solutions for clinical trials as well as NIS and other research projects in accordance with regulations in order to meet the ambitious timelines of our clients.

HIGHLIGHTS

Since the foundation of the company in 1998, our team increased to currently 49 highly qualified employees and a manageable number of freelancers with long-term experience in clinical research.
We focus on a very personalized service tailored to your individual needs for the efficient conduct of your project. Our excellent quality of service is based on the personal commitment and motivation of our employees directed to your project. Up to now more than 700 projects have been completed successfully.
The majority of our activities is placed in Germany with experience in the area of coordinating European business.
Individual projects and challenges are welcome especially.

SERVICES

Design / project management
- ▷ Design and conduct (industry / universities) of all kind of clinical studies
- ▷ Setup, coordination with and for the involved parties

Regulatory
- ▷ Submission to authorities and ethic committees
- ▷ Development / realization of amendments, submission of additional sites

Medical Writing
- ▷ Development and preparation of protocol, informed consent forms etc.
- ▷ Preparation of annual safety reports (DSUR) interim study reports and CSR
- ▷ Design / quality check / preparation of dossiers for benefit assessment (AMNOG)
- ▷ Preliminary work / preparation of scientific publications, presentations, and poster including literature research
- ▷ MedDRA and ATC/DDD coding

Monitoring – pre-trial and periodic
- ▷ Recruitment of investigational sites / review of site qualification
- ▷ Project specific training of team members and site staff
- ▷ On site monitoring and co-monitoring

Trial Coordination
- ▷ Maintenance of TMF / tracking / active pursuance of open issues
- ▷ Investigator payment according to German law / processing of fees / payments
- ▷ Day-to-day operations and quick reaction to changes
- ▷ Preparation of meeting minutes at scientific meetings

Data management / eCRF / Biometrics
- ▷ Data management, eCRF and Biometric services via cooperating companies