

COMPANY PROFILE



Ralf Freese
Medical Director
r.freese@ctc-north.com



Saskia Borregaard, PhD
Director, Clinical Trial Management
s.borregaard@ctc-north.com

History and Mission

CTC North is a privately owned full-service CRO located at the University Medical Center Hamburg-Eppendorf (UKE). Its objective is to guarantee the professional conduct of clinical trials in accordance with ICH-GCP, the German Drug Law (AMG) and the German Medical Device Law (MPDG) within a university environment.

Since 2006 **CTC North** has been providing a comprehensive range of professional services for clinical Phase I-III trials.

Highlights

Dedicated Phase I research facility, including 35 overnight beds as well as 10 beds with intensive monitoring capability and 18 outpatient places. Emergency infrastructure (ICU) within the University Medical Center to ensure qualified emergency aid, especially for first-in-man trials.

Modular concept of projects, offering individually tailored project designs from full-service packages to clinical conduct only.

Certified according to ISO 9001:2015, a standard focusing on quality management.



CTC North GmbH & Co. KG

Martinistr. 64
20251 Hamburg, Germany

Tel. +49 40 524719-0
Fax +49 40 524719-199

BVMA member since 2014
Audits passed in 2014, 2017
and 2020



www.ctc-north.com

Services

General Services

- Complete clinical trial management for the Pharmaceutical, Biotech and Med-tech Industry as well as for Investigator-Initiated Trials
- Professional recruitment (patients, healthy volunteers and special populations)
- Monitoring and Safety Management
- Data Management
- Medical Writing
- Quality Management
- Advisory and consulting services for all aspects of clinical development and conduct
- Support services for clinicians managing European Research Grant applications
- Training courses for all study personnel (e.g. GCP, AMG, MPDG)

Phase I Services

- Planning, clinical conduct and analysis of Phase I trials from first-in-man up to proof-of-concept providing a link between healthy volunteer and early patient trials
- Full Phase I trial setting with a highly experienced core team and state-of-the-art facilities and equipment

Phase II-III Services

- Multi-center Study Management and Monitoring
- Study site activities in Phase II to III trials in a professional university setting
- Close collaboration with clinical and research specialists
- A large pool of highly experienced study nurses