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BVMA member since 2018
Audit passed in 2018

kfgn KLINISCHE FORSCHUNG GRUPPE NORD
EXPERTS IN CLINICAL TRIALS

HISTORY AND MISSION

The Klinische Forschung Gruppe Nord (kfgn) is a network of seven clinical trial sites throughout Germany, which deal exclusively with conducting clinical studies on patients and volunteers. Our clients are pharmaceutical companies, Contract Research Organizations (CROs) or manufacturers of medical devices. Our goal is to support medical progress and to do everything we can for the well-being of our participating patients.

HIGHLIGHTS

The company was founded in 1995 by Prof. Dr. med. Hanns-Gerd Dammann as the Klinische Forschung Hamburg GmbH.

Since then, the kfgn has constantly been growing and today it is a network of seven clinical trial sites throughout Germany, which deal exclusively with conducting clinical studies on patients and test persons.

Since 2013, the management and service department (kfms) has been part of the company. The kfms undertakes superordinate tasks for the test centers in the areas of project and feasibility management, contract and quality management as well as in patient recruitment. It is the central contact for sponsors and CROs.

SERVICES

Your benefits at a glance

- Seven centrally managed clinical trial sites in Germany
- Highly experienced in-house investigators and clinical research staff
- More than 20 years of experience
- Database of over 55,000 patients across all sites for more than 65 different types of symptoms
- Own online recruitment portal: www.patientenstudien.de
- High quality and standardized study-planning, evaluation and documentation
- Over 100 partners from the pharmaceutical industry and CROs
- Strong and reliable commitments to our partners

The Services for sponsors and CROs

- Study Feasibility Analysis (Pre-Feasibility / Feasibility)
- Preliminary identification of suitable patient groups
- Professional compilation of the required documents for the ethics submission
- Fast, standardized study preparation and planning
- Identification of the study-specific quality characteristics
- Central performance management and contract management
- Advice on compensation structuring in clinical trials
- Timely patient recruitment via own patient recruitment portal
- Study and Site-Specific Advertising Strategies
- Extensive patient database
- Generation of study data at the highest quality level
- Study adapted inspection data and Audit-Robust worksheet data
- Comprehensive quality management: quality control / quality assurance during implementation and documentation
- Work in accordance with the SOPs and observance of all applicable ethical and regulatory principles as well as ICH standards