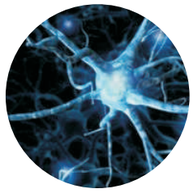
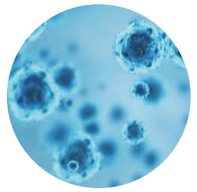
Your trendsetting CRO in the heart of the greater Nuremberg area



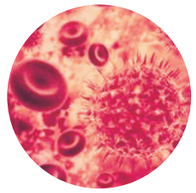
INNOVATIVE

Research wants innovation.
Together, your ideas and our ideas give rise to new research approaches and creative solutions for all study projects within the scope of clinical research.
With MEDICRO, you can be sure to have a constructive and valuable partner at your side.
Because innovation leads trials to success.



FOCUSED

Providing support for your trial, your project, your concern, is the focus of MEDICRO, a nationally and internationally operating contract research organisation.
We conduct clinical trials in the field of pharmaceuticals and medical technology with precision, in compliance with GCP requirements, without red tape and with the highest quality standards.



EFFICIENT

The costs for developing medicinal products and medical devices are steadily increasing.
MEDICRO is fully committed to supporting you throughout the duration of your project to ensure that delays can be recognised early on and prevented.
Our staff members' motivation will make your trial a success.

Contact us:

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Klostersteig 19
D-91580 Petersaurach
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Fax: +49 9872 9699029

MEDICRO
Clinical Trial Services and Medical Translations

Your Partner for Clinical Trials

MEDICRO GmbH
Klostersteig 19 • D-91580 Petersaurach • Phone: +49 9872 9699027
office@medicro.de • www.medicro.de

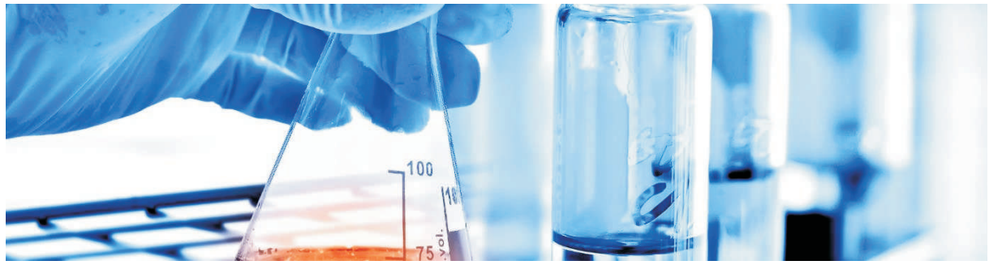
WHO WE ARE

MEDICRO GmbH has been conducting human drug and medical device trials as an independent contract research organisation for more than 15 years.

Depending on your individual requirements we perform a variety of short- or long-term tasks during your trial. From the planning and the submission all the way to the implementation and conclusion of your research project – we are the professional partner at your side.
High-quality translations and advanced training opportunities round off our range of services.

OUR TRIAL EXPERIENCE

In the fields of human drug / medical device / paediatric trials



- Oncology and haematology
- Neurology and psychiatry
- Cardiology and angiology
- Endocrinology and diabetology
- ENT diseases
- Rare or orphan diseases
- Rheumatology
- Gastroenterology
- Dermatology
- Radiology
- Respiratory diseases
- Virology and infections
- …

WHAT WE OFFER

Study preparation

- Project budgeting
- Drafting of informed consent forms
- Creation of documents and worksheets
- Creation and revision of SOPs
- Recruitment support (advisory service and information material)

Seminars

- In-house or as a webinar
- ICH-GCP training for on-site personnel including national regulations
  - Basic course: 2 days
  - Refresher: 1 day
- Study Nurse courses
- Current topics from clinical research

Translations

- Languages: ENG, DEU, FRA, SPA
- Adaptation to national regulations
- Documents:
  - Correspondence with authorities
  - Protocols, synopses
  - Informed consent forms, ….
- Proofreading of medical and scientific texts

Trial coordination

- Short- or long-term assistance, also as a temporary stand-in
- General documentation
- Data entry in CRF/eCRF
- Query resolution
- Coordination of vendors
- Auditing of inventory and material
- Archiving

Submissions

- Advisory service
- Support in obtaining essential documents
- Submissions to the higher federal authority and/or ethics committees
- Correspondence

Monitoring

- Pre-study, initiation, motivational, monitoring visits
- Remote Monitoring
- Accompanied visits with a quality assessment
- Close-out and archiving

Quality assurance

- Audit/inspection preparation
- Review of the Trial Master File
- Review of the trial file at the site
- Verification of monitoring quality
- Accompanied visits