



**MTZ**  
**CLINICAL**  
**RESEARCH**

*Quality is a member of our team*

Clinical Research Site

WITH FULL SERVICE

SINCE 2002



## Proven advantages:











-  2 locations
-  Experience: 260 performed early and late stage clinical projects, including first in human dosing (FIH), multiple application (MAD), drug drug interaction (DDI), proof of concept (POC) and others
-  Quality: ISO 9001 (2009–2018)  
ISO 9001: 2015 (obtained in 2018)  
3 inspections (FDA and Polish)
-  Security: Procedures complying with PN ISO/IEC 27001 and Company Policy
-  Investigational team: experienced, highly trained, reliable
-  Timelines: respected















[mtz-clinical.pl](https://mtz-clinical.pl)



## Offered services:

-  Conducting clinical projects at own Clinical Research Site (2 locations, 50 km apart, 62 beds offering flexibility and back up)
-  Preparation of study documentation (synopsis, protocols, ICF, CRF and final reports)
-  Obtaining legal-administrative (URPL) and bioethical (KBE) approvals
-  Clinical study management
-  Safety analysis at own certified medical laboratory
-  Medicinal Products (including infusions) processing at own Pharmaceutical Laboratory
-  Biological sample processing and storage
-  Data management
-  Biostatistical analysis (SAS)
-  Quality audits in research centers
-  Trainings on the principle of cGCP
-  Study monitorings

## Scope of studies:

-  Bioavailability/Bioequivalence (BA/BE) in healthy volunteers and in patients
-  Biosimilarity in healthy volunteers and patients
-  Early and late development in:
  -  Cardiology
  -  Dermatology
  -  Diabetology
  -  Gynecology
  -  Hematology
  -  Oncology
  -  Neurology
  -  Pulmonology
  -  Rheumatology

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