**MaganaMed GmbH**
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# *MaganaMed GmbH*

## Business Strategies

MaganaMed GmbH develops and distributes software solutions for the medical sector. Our flagship product is a system for online documentation and management of clinical studies. Our software fits perfectly into the daily workflows of creating and managing clinical studies. Ease of use is especially important to us, because good software stays in the background and supports your workflow.

The system is ideally suited for use under the regulatory requirements in medical device, IVD or pharmaceutical development. All processes are GDPR-compliant, all data is protected by modern encryption technologies (TLS 1.3 and AES-256) and the entire documentation and execution is implemented in compliance with GCP.

Whether academy or industry – Magana Trial Manager ensures smooth and complete documentation: from small individual studies, research consortia to product approval (e.g. CE-marking, IVD-R, ...)

Study software Made in Germany: intuitive, high-performance and secure!

## Core Technologies and Services

Our core competencies relate to

- Software as a Service: no installation or maintenance effort on the customer's premises
- Documentation and management of clinical trials of any size and complexity
- Support of regulatory requirements for clinical studies – e.g. ISO13485 / CE-marking, GCP, IVD-R, ...
- Data security (e.g. according to GDPR)
- Intuitive creation of electronic CRFs
- If desired, we can create the CRFs and study structure for you
- Support of hybrid documentation (electronic plus paper)
- Support in management and documentation of biosamples (plasma/serum/urine)

## Distribution

Direct sales to universities, hospitals and companies in the fields of MedTech, diagnostics, pharma, biotech and medical devices in Europe.