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History and Mission

ACM's clinical trial service platform includes Study Management, Site Monitoring and Management, Laboratory Sample Management, Drug Supply Management and Medical Writing Support. Our highly trained and accomplished clinical research professionals manage and conduct clinical studies according to the latest GCP and regulatory requirements. ACM has experience in the study conduct of phase I-IV clinical trials and post-marketing surveillance in pharmaceuticals and medical devices.

Highlights

Since its founding in 2004, ACM has provided clinical trial operational services to large and small sponsors of clinical studies throughout the „DACH" region.

In 2006 ACM changed its structure to a GmbH and in the following year moved its operations office to central Berlin allowing for future growth and more efficient service for our clients.
Since 2009, ACM has been an audited member of the BVMA.

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BVMA member since 2009
Audits passed in 2009, 2012, 2015, 2018 and 2021

ACM®
ALLIED CLINICAL MANAGEMENT

www.allied-clinical.com

Services

Clinical Trial Management
- Study set up
- Site feasibility and selection
- Risk assessment
- Investigator meetings
- Vendor management
- EC and Central Authority submissions
- Study newsletters

Site Management
- Site initiation
- On-site and remote monitoring
- Investigator training
- Drug accounting
- Site close out

Medical Writing Support
- Providing support to medical writers and study management through document content management systems

Biological Sample Management
- Management of the collection, shipment and processing of biological samples
- Developing the logistics plan for the shipment of biological samples
- Training of investigational sites regarding the proper handling of biological samples according to trial specifications

IMP Management
- Management of the timely supply of IP to trial sites
- Tracking of IP from delivery to destruction
- Tracking the validity and expiration of IP
- Creating IP labels and packaging according to applicable laws
- Training of IP use at investigational sites