

## ■ COMPANY PROFILE



**Barbara Lopez Kunz**  
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### History and Mission

**Caidya's** mission is to help bring life-changing therapies to patients around the world through flexible, patient-focused clinical research. Founded in 2003 as Clinipace, the company grew into a full-service CRO through global acquisitions and a merger with dMed in 2021. At its core, Caidya believes in listening to clients, adapting to rapid industry change, and delivering personalized, efficient solutions that improve the drug development process and accelerate access to innovative treatments.

### Highlights

**Caidya** is a global, multi-therapeutic CRO conducting studies in 50+ countries and regions around the world. We deliver comprehensive clinical research solutions—from pre-IND planning through regulatory submission and post-marketing surveillance. Our team combines deep therapeutic expertise with proprietary technology to deliver high-quality data, ensure transparency, and support informed decision-making. With a strong culture of collaboration, agility, and customer focus, Caidya builds true partnerships with biopharma innovators, enabling complex trials to move forward efficiently and delivering tailored solutions that help bring new therapies to market faster.



www.bvma.de

### Caidya

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BVMA member since 2004  
Audits passed in 2004, 2006, 2009,  
2012, 2015, 2018, 2022 and 2024

# caidya®



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### Services

#### Clinical Development Phase I-IV

- Biostatistics and statistical programming
- Clinical data management
- Clinical operations
- Investigator recruitment and site selection
- Medical monitoring
- Medical writing
- Project management
- Protocol feasibility
- Quality assurance
- Safety/pharmacovigilance

#### Therapeutic Areas

Oncology, Rare Disease, Immunology, Pediatrics, Cell & Gene Therapy, Nephrology, Respiratory, Neurology, Infectious Diseases, Cardiovascular, Dermatology, Hematology, Ophthalmology, Endocrinology, Gastroenterology, Women's Health.

#### Regulatory Affairs

- Product development
- Portfolio management
- Regulatory strategy
- Manufacturing and quality support
- Medical and regulatory writing
- Regulatory publishing/submissions

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