

COMPANY PROFILE



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History and Mission

CRS is one of the leading full-service CROs (Phase I-IIa) in Europe with more than 40 years of expertise in clinical development. We provide expert services in fast and efficient conduct of high-quality trials to help customers bring safe and effective pharmaceutical products to market - all realised by our team of more than 250 dedicated professionals. Our outstanding quality and reliability is reflected by the high number of satisfied international clients.

Highlights

CRS. Top Class in Early Phase.

- More than 40 years of experience
- More than 2.000 successfully completed trials
- Various CRO Leadership Awards

CRS. Service Portfolio.

- From FIM to POC in patients
- All types of Clinical Pharmacology Studies
- Studies in renally and hepatically impaired patients
- Core therapeutic segments: Cardiometabolic/Cardiovascular, Women's Health, Dermatology and others
- Single-Site or Multi-Site (Site+Concept) setup
- Maximum operational and financial synergies

CRS. Experts. Early Phase.

First in Human to Proof of Concept in Patients.



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BVMA member since 2019
Audits passed in 2019 and 2022



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Services

Full-Service

A complete service solution out of one hand under one contract.

- Consultancy
- Medical Writing
- Project Management (Single-Site, Multi-Site)
- Clinical Conduct
- Quality Assurance
- Clinical Trial Supply Management
- Data Science
- Monitoring by external partners
- Pharmacy Services
- Other Services (Monitoring, Pharmacovigilance, etc.) provided by external partners

Extensive Expertise

Patients in our core therapeutic segments

- Renal & Hepatic Impairment
- Cardiometabolic / Cardiovascular Diseases
- Women's Health
- Dermatology
- Respiratory

Healthy Volunteers

- FIM
- SAD | MAD
- Biosimilars
- BE | BA
- PK | PD
- DDI | FDI
- QTc
- Vaccination studies
- Others