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History and Mission

CROMSOURCE is an ISO-certified international provider of outsourced services to the pharmaceutical, biotechnology and medical device industries, specialized in clinical development and staffing solutions. A well-established full-service CRO, **CROMSOURCE** is unique in offering an End-to-End Guarantee covering trial timelines, enrolment and contract price. **CROMSOURCE** operates through offices across all regions of Europe and North America and delivers a comprehensive breadth of services.

Highlights

End to End Guarantee
Quality data. On time. On Budget. Guaranteed. A unique concept in an environment where change orders and delays are common place with other service providers.

Expert Trial Rescue
The experienced CROMSOURCE team quickly assess the situation and implement tailored solutions which get such trials back on track.

Feasibility Plus
Feasibility Plus is provided without obligation at the proposal stage. Through direct contact with potential investigators, Feasibility Plus provides accurate country and site selection data, and allows precise budget and timeline forecasts.

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BVMA member since 2010
Audits passed in 2010, 2013, 2016 and 2019

www.cromsource.com

Services

Clinical Development Services
- Feasibility / Site Selections
- Clinical Operations: Project Management & Monitoring
- Regulatory Affairs and Medical Monitoring
- Medical Writing: Protocols, CIP
- Quality Assurance - Audits
- Pharmacovigilance / Materiovigilance
- Data Management and Statistics
- Drug and Vendor Management
- Regulatory for US/EU (eg. CER, PMCF, consulting for FDA Pre-IND/IDE)
- Legal Representative – US/EU
- Consultancy for customized solutions to optimize product development, minimize costs and time to market for novel therapies and medical device
- IT: Customized Tools & Resources (eg. THeCRF, TheDiary; TheSurvey)
- Staffing Solutions (FSP model or short- and/or long-term or permanent staff)

Early Phase Services
- ADME studies
- Bioavailability
- Bioequivalence
- Dose ranging/multiple dose tolerance
- Drug-drug interactions
- First in Human (SAD, MAD)
- Food effect studies
- Patient Studies
- Pharmacokinetics/pharmacodynamics
- Proof of Concept
- QTc studies