# Corporate Factsheet


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**CROMSOURCE Quality**
ISO 9001:2015
multi-site certified quality management system.
ISO 14155:2020
conformity confirmed.

## Company Profile

**CROMSOURCE** is an ISO-certified international provider of outsourced services to the pharmaceutical, biotechnology and medical device industries, specialising in clinical development and staffing solutions.

**CROMSOURCE** was founded in 1997, more than 25 years ago. Since that time, our successful growth has been built on stability, integrity, and high levels of customer satisfaction. These factors all contribute to a high rate of repeat and referral business, powerful evidence of the quality of our services. We have grown steadily, but responsibly, to become an organisation of over 350 organised and well-trained experts.

A well established full service CRO, **CROMSOURCE** is unique in offering an End-to-End Guarantee covering trial timelines, enrolment and contract price. This guarantees our clients that their trials are delivered on time and within the contract price without CRO-initiated change orders. **CROMSOURCE** operates through offices across all regions of Europe and North America and delivers a comprehensive breadth of services.

**CROMSOURCE** supports the full spectrum of clinical development via our Pharmaceutical, Medical Device and Staffing Solutions divisions. We seamlessly move biopharmaceutical products from first-into-human through all subsequent phases of pre- and post- approval research internationally. We also support medical device projects through regulatory planning and execution, to pre- and post-market clinical investigations in Europe and North America.

## Global Reach

**CROMSOURCE has the global reach to conduct biopharmaceutical and medical device studies around the world. CROMSOURCE**, with world headquarters in Verona, Italy, is a leading CRO in Europe and the US with a solid infrastructure and operational subsidiaries in Belgium, Germany, Poland, Spain, Switzerland, the UK, the Netherlands, and the US.

From our office locations across Europe and North America, **CROMSOURCE** employs experienced field-based teams in locations across the globe to provide expert capabilities in regions including the Middle East Africa, APAC, and South America.

### GLOBAL PRESENCE, LOCAL EXPERTISE

CROM SOURCE
advise | agree | deliver

# Corporate Factsheet

## Services

- Feasibility and Site Selection
- Clinical Operations: Study Start-Up, Project Management and Clinical Monitoring
- Data Management and Biostatistics
- Medical Monitoring
- Medical Writing
- Quality
- Pharmacovigilance/Materiovigilance
- Vendor Management
- Regulatory
- Strategic Consulting
- Legal Representation
- Technology Solutions
- Flexible Resourcing Solutions

### Pharmaceutical

- From single site to complex global trials.
- 300,000+ patients studied.

### Medical Device

- All sizes of projects conducted.
- 200,000+ patients studied.

### Flexible Resourcing

We are equally comfortable acting in a functional service provider setting or simply providing short-and long-term or permanent staff for your organisation.

### Therapeutic Areas including but not limited to:

Autoimmune/Inflammation
Cardiovascular
CNS - incl. Pain and Psychiatry
Dermatology
Endocrinology/Diabetology
Gastrointestinal
Haematology Imaging
Infectious Diseases incl. HIV
Nephrology/Urology
Oncology
Ophthalmology
Rare Disease
Respiratory
Rheumatology
Vaccines
Women's Health

### Therapeutic Areas including but not limited to:

Cardiovascular
CNS - incl. Pain
Dermatology
Endocrinology/Diabetology
Gastrointestinal
In-vitro diagnostic
Oncology
Ophthalmology
Orthopaedics
Respiratory
Women's Health

### Services:

- Functional Solutions (FSP)
- Hybrid Delivery Model
- Insourcing / Staff Augmentation
- Combined Solutions Provision
- Permanent Placement

## Why CROMSOURCE?

CROMSOURCE is the only CRO offering a real "End-To-End Guarantee" for your clinical trial:

- Guaranteed Study Start-Up Time
- Guaranteed Enrolment
- Guaranteed Price with no CRO initiated Change in Scope
- Guaranteed Date for Database Lock

## How we do it:

It starts with our Feasibility Plus™ program. We directly contact investigators in all countries under consideration, identify enrollable patients, and assess both competing trial and in-country regulatory situations. Our process is so thorough that we can guarantee both study start-up time and enrolment. With feasibility assured, we offer our One Trial One Price™ program – our pledge that you will not be subjected to multiple post-contract change orders. The price agreed upon at contract signature will be the only price you pay. And with more than 25 years of vetting, we are so confident in our processes that we will guarantee the date for Database Lock.

CROMSOURCE will write this into your contract with penalties that actually mean something.