

## SUMMARY

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- JOIN OUR RESEARCH NETWORK

# ABOUT US

**Clinica CCBR Clinical Research Network** is a privately-owned life science company that provides **first - class** clinical trial management services to pharmaceutical, biotech, or medical companies.

**Clinica CCBR Clinical Research Network** plays an **essential role in the clinical trial's overall success:**

- is a the **bridge** between the sponsors, CROs and the sites that are conducting research;
- is an **integral part** for a trial to achieve accelerated timelines, streamlined processes, improved data management, and more accurate results.



**Our goal** is to successfully complete clinical trial and deliver best results on time and on budget.

We aim to provide all that is necessary to get studies up and running faster, while a dedicated patient engagement team drives identification and recruitment of suitable patients, using an extensive database of patients and targeted campaign.

Combining our data and experience, along with ensuring the right site and investigator mix, are what allow us to bring greater certainty and speed to the enrollment process.

**Track record: Over 1500** subjects randomized in **over 30 clinical trials** in **over 15 years.**

**Our Team** is highly experienced in the medical research industry and is committed to providing high-quality service in the business. The specialists are certified with clinical study expertise and a grasp of customer demands.

Our country expertise, supplemented by a network of partnering sites, ensures we provide a truly top quality service to researchers, CROs and Pharma companies.

We have an extensive experience in a wide range of therapeutic areas.

## Ethics and Compliance

**Clinica CCBR Clinical Research Network** delivers high quality services, certified through the implementation of ISO 9001:2015 standard and SOPs developed in-house.

We implement consistent standard operating procedures and quality control throughout the sites.

All services provided are in accordance with current EU Legislation, FDA regulations and the applicable national laws.

# ABOUT US

**OUR VISION** is to be the preferred SMO partner that drives the evolution of the industry, enabling pharma, biotech's to accelerate the delivery of innovative, life-changing treatments.

**OUR MISSION** is to enhance the quality of clinical trials by partnering with and providing to pharma and biotech with forward thinking, cost-effective methods combined with performance driven sites.

## OUR VALUES:

- **Commitment:** we are passionate about supporting partners improve results and patient outcomes.
- **Integrity:** business is conducted with integrity and according to ethical standards.
- **Ownership & Accountability:** we are acknowledged for our scientific expertise, and long track record of efficient and innovative clinical trials.
- **Performance:** we always strive to improve our performance and the performance of our clients.
- **Sense of urgency:** we approach every opportunity, every challenge with a sense of urgency and recognize that our proactivity is critical to success.
- **Teamwork:** our best performance depends on internal and external partnership and collaboration.

## OUR FOCUS:

**Understanding client needs:**  
the importance of time, cost and quality, and are able to guarantee these metrics are met. We are offering a comprehensive portfolio of services which can be tailored to meet the individual needs of each client.

**Being always flexible and oriented to final goals.**

**Network Power:**  
we aim to provide professional, high-quality clinical research through a dedicated network of geographically distinct clinical investigative sites with large readily available patients database.

**Communicating with all stakeholders openly and in a timely manner.**

**Improving the outcome for patients.**

# OUR CENTRALIZED E-SERVICES

What are the benefits of working with us?

We connect Sites with Sponsors and CROs in a paperless environment, via platform that ensures streamlined workflows with a “one system, one source” approach to clinical data capture.

We accelerate trials by ensuring adequate resources for speed site selection, patient recruitment and retention; clinical monitoring.

Our systems enables electronic data capture and remote monitoring with powerfully integrated solutions that protect data integrity and compliance across 21 CFR 11, ICH-GCP, GDPR, HIPAA.

- You can rely on the quality of our data



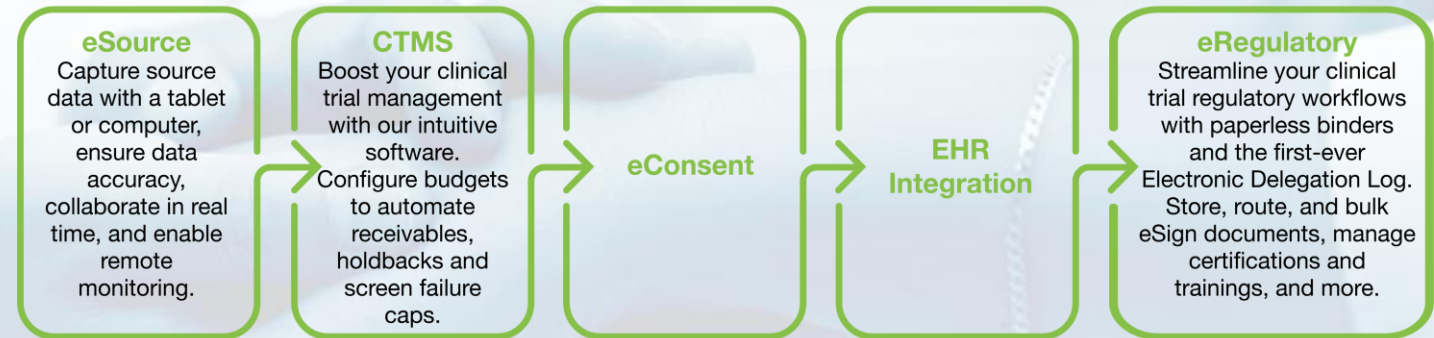
- Standard launch processes to deliver uniform conduct and quality through all the sites



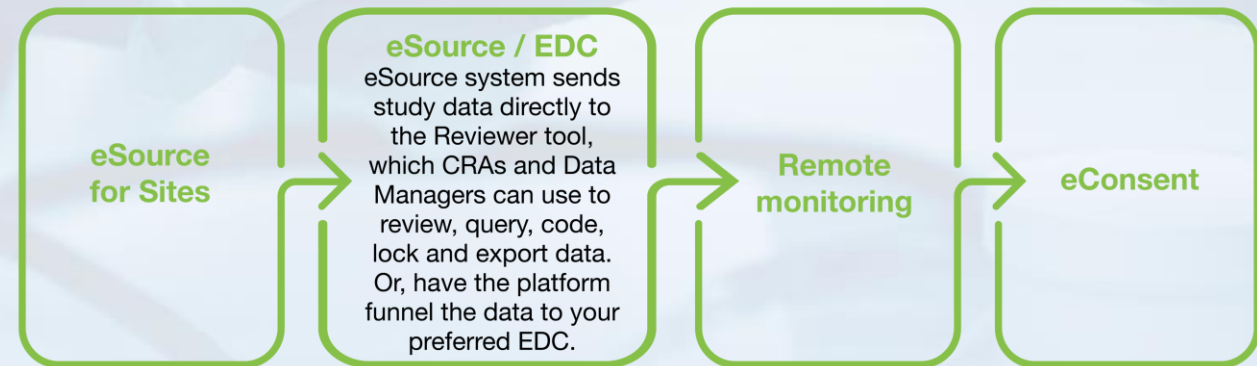
- Centralized training and oversight

- Centralized integrated eSource/EDC

For Sites:



For Sponsor / CROs:





# OUR RESEARCH NETWORK PARTNERS

- **Clinical Research Network:** 51 clinical research affiliated centres positioned in all relevant university centres, offering wide geographical coverage
- **Specialties List:** Allergology, Cardiology, Diabetes Mellitus, Nutrition & Metabolic Diseases, Dermatology, Endocrinology, Gastroenterology, Geriatrics & Gerontology, Haematology, Infectious Diseases, Internal Medicine, Medical Rehabilitation, Nephrology, Neurology, Obstetrics & Gynaecology, Oncology, Ophthalmology, Orthopaedics, Otorhinolaryngology, Paediatrics, Pneumology, Psychiatry, Psychology, Rheumatology, Urology
- **Total No. of Healthcare Professionals:** more than 900 Physicians, more than 1500 other types of Healthcare Professionals.
- **Total Equipment:** 27 CT, 30 MRI, 3 PET/CT, 2 SPECT/CT + 150 other equipment (Rx stations, DXAs, echographs, 3D mammographs etc.)
- **6 clinical laboratories** including also genetic analyses, pathology and molecular biology tests.
- **3 in-house dedicated pharmaceutical facilities that are supporting day hospitalization services**
- **We are using up-to-date** and complex automatic line processing biological laboratory tests and technologies provided by internationally renowned producers such as: GE, Abbott, Storz, Siemens Healthineers, Varian, Elekta, Hologic, Philips.
- **The sites we manage are selected after detailed analysis of the clinical trial industry.** We have taken into consideration important factors such as: projects saturation, investigators motivation, level of equipment, regional

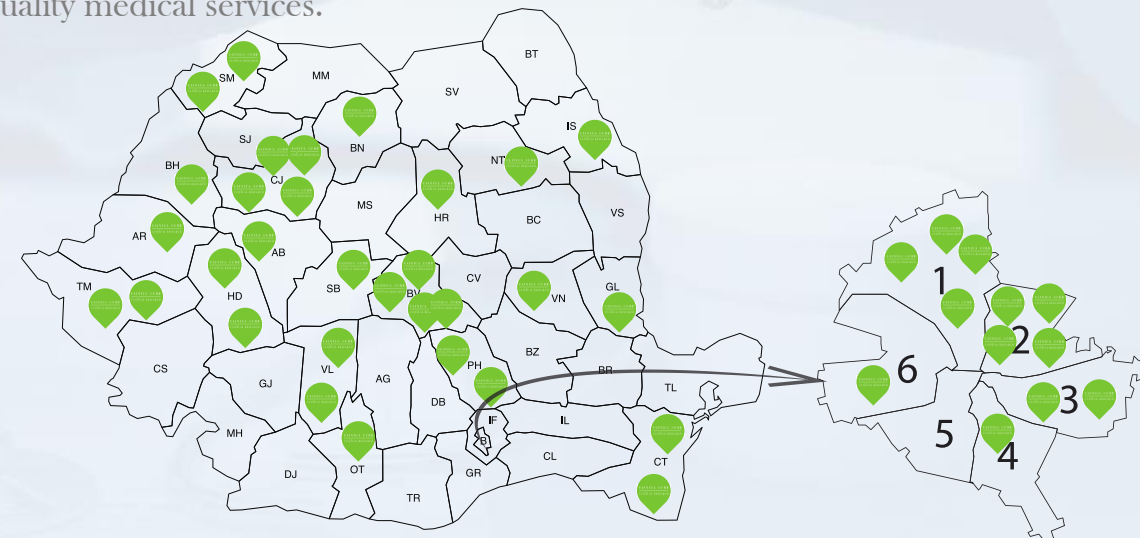
distribution of patient groups, access to medical care.

- **Recruitment capabilities** that combines:

- ❑ **Total Patient Traffic:** 1.5 million patients/year.
- ❑ **Strategic Partnership with Proprietary Digital Communication Channel** securing an **Outreach Traffic** of over 1.4 million individual users/month.

## All our Sites offer:

- Easy to find location.
- State of the art medical and lab equipment.
- Comfort and professionalism.
- Friendly staff trained in the International Standard of Good Clinical Practice (GCP).
- Top quality medical services.



# JOIN OUR RESEARCH NETWORK

Contact us!

- If you are a Sponsor: [sponsor@clinicaccbr.com](mailto:sponsor@clinicaccbr.com)
- If you are a CRO: [cro@clinicaccbr.com](mailto:cro@clinicaccbr.com)
- If you are a Clinical Research Center: [crc@clinicaccbr.com](mailto:crc@clinicaccbr.com)
- If you are a Clinical Research Professional: [crp@clinicaccbr.com](mailto:crp@clinicaccbr.com)