

CORPORATE PROFILE



ABOUT US

01

Fervor is the first customer-centric CRO

02

Founded by expert dedicated to advancing medical science.

03

Fervarriors strategically balance

- Science
- Patient
- Clinical Development
- Data Analytics



ABOUT US



Global CRO

Global full service CRO providing integrated services



Experience

Highly Experienced Team with Extensive Knowledge



Outreach

4 Continents, 6 Countries, 8 Offices across the world



Regulatory Expertise

Impeccable Regulatory track records



Quality

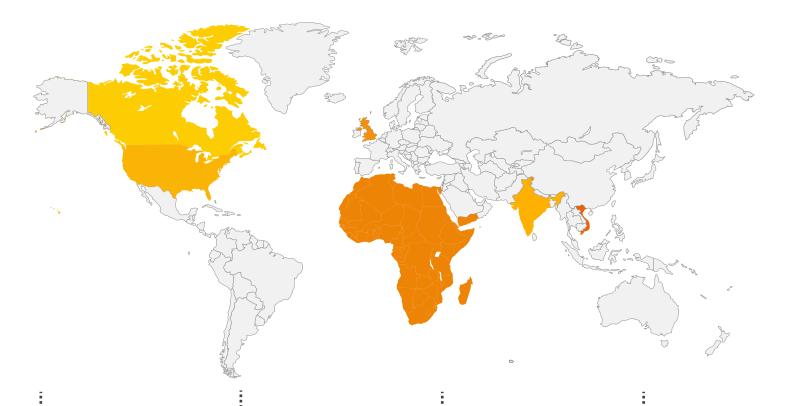
Always Audit and Inspection Ready





GLOBAL PRESENCE





Ontario 3388 sandwich street, appt. 213 Windsor, Ontario -N9C1B1 Canada

CANADA

US

New York 575 Lexington Avenue, New York, NY 10022

UNITED KINGDOM

56 Pennington, Orton goldhay, Peterborough Pe2 5rb

AFRICA

Sierra Leone9 Jakema Street, Bo,South Sierra Leone

VIETNAM

Hanoi 27 Hang Bai, Hanoi Vietnam

INDIA

Himachal Pradesh Aashirbad Building, I & II Floor, Kamal Marg, Nehran Pukhar, Dehra - 177104

INDIA

INDIA

Delhi

H-1/603, DDA

Multi-Storey,

Sector-18B, Dwarka,

New-Delhi-110087

Hyderabad B, Plot no. 20, Avanthi Enclave, Madinaguda, Hyderabad Telangana-500049



Global Comprehensive Service Portal -

Drugs & Device Development

Early
Development
Service

Clinical Research Units

PK/PD Modelling

Data Visualization & Analysis

Pharmacodynamic Models & Simulations

Commercialization & Outcomes

Real World Intelligence

Real World Evidence Strategy & Analytics

Real World & Late Phase Research

Direct to patient contact solutions

Strategic Regulatory
Services

Language Services

Clinical & Medical Translations

Linguistic Validation

Functional Services Provider

FSP Resources for Clinical Development & Post Marketing Program MedTech
Development
Services

Clinical Research Services

Post Marketing Compliance

MedTech Technology

Compliance Solutions





Global Comprehensive Service Portal -

Drugs & Device Development



Biometrics

Biostatistics

Data Management

Medical Writing & Publishing

Adaptive trial

Scientific Operations

Medical Affairs

Pharmacovigilance

Interactive Response Technology

Clinical Trial Supplies
Management

Clinical Research Services

Project Management

Clinical Operations

Feasibility, Start-up & Regulatory

Site & Patient Solutions

Patient Recruitment and Retention Services

Drug Development Services

Preclinical/Non-Clinic al Development

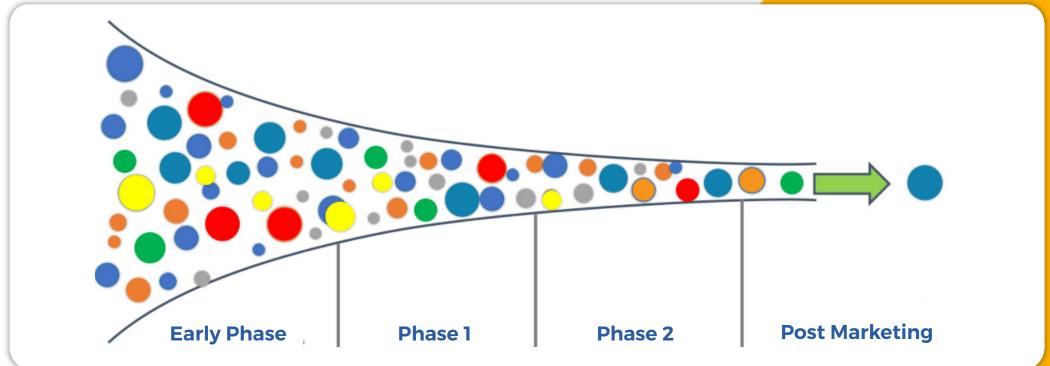
Clinical Development

Pharmacokinetics

Due Diligence

FULL-SERVICE CAPABILITIES & EXPERTISE

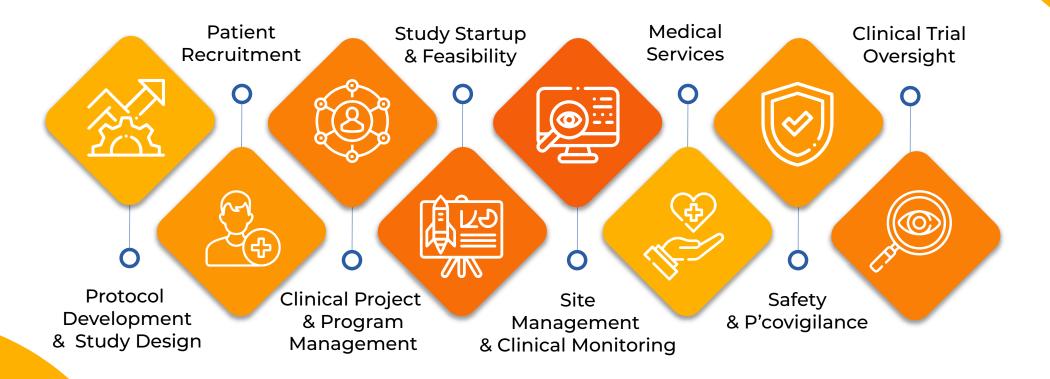
Throughout the clinical development life cycle, Fervor provides medical and regulatory leadership guidance and efficient, disciplined operational execution of studies around the world. Our comprehensive capabilities, resources and global footprint ensure quality and timely development across all phases of research.







CLINICAL TRIAL OPERATIONS



Fervor's Expertise with unmatched proactive, responsive and pragmatic approach tailor-made to match the specific needs of your projects. Fervor's global experience in effectively designing and executing studies, partner from strategy to approval in complex indications and modalities across a therapeutic areas and patient populations. Fervor offers either singly or as a full-service solution.

TECHNOLOGY UNIQUENESS

Our Clinical Development Services (CDS) Project Managers provide solution-focused project management across our single, multiple and full service and Full-Service CDS business lines, so you have one point of contact for each project within CDS. Our CDS Project Managers provide support for all non-therapeutic Business Unit specific areas, from Early Phase through the support of regulatory submissions

We offer

- ✓ Strategic of R&D
 - **Vendor Selection**
- Vendor Management
- End to End Program/Project Management
- Customized Project
- Clinical Program Rescue

Services

- Performing regulatory feasibility for a new program or a new project
- Support in planning of a Program with a multiple projects or an individual projects
- Identifying, auditing, and supporting in managing the local vendors like Central, Clinical Trial Supply management vendor, Data Management, PVG and so on
- Delivering consistency across programs and submissions
- Covering all therapeutics areas and indications

Outcomes

- Single Point of Contact
- Expertise in Clinical Development Services
- Accountability & transparency with a proven record of successful delivery
- Timelines & risks are managed proactively
- Applying knowledge from past experiences to ensure accurate financial management
- Strategic advice on registration and commercialization





REGULATORY & DEVELOPMENT SERVICES

Fervor's delivers the innovative regulatory strategies for the complex programs or borderline products or the products where the regulatory pathways are not straight forward for the Pharmaceuticals, Medical Devices, and Biotechnology companies







BIOMETRICS AND DATA ANALYTICS



FERVOR is committed to providing the highest quality services.

Our global Data Analytics group is a dedicated & experienced team of biostatistics, statistical programming, and data management professionals devoted to delivering individualized, adaptable, and dedicated services to support all your data and analytical needs.

The key to excellent data management is flexibility. We respond to your needs by nimbly adjusting to change and striving to avoid timeline modifications. This ability is the true mark of a valuable CRO partner — but FERVOR surpasses that basic data management function by applying strong cost-effective measures, quality assurances, and therapeutic and regulatory expertise.

- Statistical Analyses & SAP
- Design CRF as per CDASH Standard
- Statistical Consulting & Development Strategy
- CDSIC: SDTM & ADaM
- Double-Data Entry (Paper CRF)
- Operate Seamlessly in multiple EDC platform
- Preparation & Generation of Randomization Plan
- Interpretation of Study Result and Writing Report



THERAPEUTIC AREAS

Attain faster pathways to commercialization, achieve greater efficiencies through better designed and executed clinical research studies, and effectively navigate global regulatory requirements to optimize your product's success. Your project team will be led by medical, regulatory and operational experts with deep therapeutic experience who are fully engaged throughout your study, proving guidance and averting potential roadblocks by staying close to the project.

We treat your drug
development journey
as if it's our very own,
providing
VALUABLE INSIGHT,
UNMATCHED CLIENT

DELIVERABLES.

SERVICE, & HIGH-QUALITY

Cell & Gene Therapy

O2 Rare & Orphan Disease.

Oncolog & Hematology

O4 Central Nervous System

O5 Hepatology & Gastroenterology

Additional Areas of Focus

Infectious Disease

Endocrinology & Metabolic

Cardiology & Vascular

Dermatology

Respiratory & Pulmonology

Nephrology & Urology

Obs/Gynae/Women Health

Pediatrics

Device & Diagnostics

DRUG SAFETY AND PHARMACOVIGILANCE

FERVOR has deep expertise and experience from early phase studies to marketed products. FERVOR is a safe pair of hands to support clients in both current state whilst planning for your strategic objectives. Our PV SMEs, comprised of both Physicians and Scientists are end-to-end partners for our clients, proactively advising and collaborating in designing and maintaining PV services and systems which are fit-for-future.

Medical Affairs

- Perform a full Medical Review, including protocol deviations
- Share clinically relevant trends with sponsors
- Continuously assess therapeutic response
- Interact with CRAs to identify safety issues early

Pharmacovigilance

- Customized Safety Surveillance
- Safety Narrative Writing
- Safety Signal Management
- Aggregate Report Management
- AE/SAE Mgmt. & Reconciliation
- Individual Case Reporting
- Safety Coding (MedDRA, WHO Drug
- Blinded/Unblinded Services
- Updating of IB with new Safety Information

We know how important your product is to you and the patients who are waiting







GLOBAL REGULATORY AFFAIRS



Drive the development of your products – from preclinical proof-of-concept candidates to approved products primed for post-marketing label extension and regulatory maintenance – by partnering with an experienced partner who knows how to help you reach your long-term commercial goals.

Fervor's highly skilled team of global regulatory experts are available to provide guidance and support for your products at any stage of clinical development for drugs and medical devices.

Preparation Strategic **Borderline** Development **Products Submissions** Regulatory **FHS Combination Agency Products** Interactions Medical Regulatory **Accelerated Devices &** Compliance **Pathways Diagnostics**

Partnering with our clients in all stages of their product development programs and product lifecycle to accelerate global therapeutic development and enable access to the market



TECHNOLOGY SOLUTIONS



Leveraging Fervor's differentiated technology solutions and data collaborations

Real world Data Collaboration

EDC **Systems**

World Class Analytics & Collaborations **World Class PVG** Systems







































FERVOR SITE NETWORK

FERVOR works in partnership with the leading medical institutions across Asia to accelerate quality clinical research for its clients and has now signed 16 major strategic partnerships with hospitals and research institutions in the region.



Speed

Site Selected to Site Initiation Visit is on average 35% Faster



Patients

Average of 55% more patients per sites



Quality

12 audits & inspections over 4 years – No Major/Critical Findings



CLINICAL SUPPLIES MANAGEMENT SERVICES



FERVOR Clinical Supplies Management is an integrated biopharmaceutical supply chain solutions provider offering a full range of primary and secondary clinical packaging, temperature-controlled logistics, storage and distribution services for the global pharmaceutical, biotech and Medical Device industries.



CONTACT US

- www.fervorhs.com
- hello@fervorhs.com
- +91 8527110300 | +91 88600 09879 | + 91 78740 33093
- Pheadquarters: H-1/603, DDA Multi-story, Dwarka, New-Delhi-110078, India



