www.kvclinicalresearch.com

CERTIFIED ISO 9001:2015 COMPANY

K V C R Since 2015

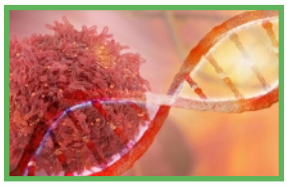
nothing beyond humanity

KV CLINICAL RESEARCH PVT. LTD.

SITE MANAGEMENT ORGANIZATION (SMO)

CIN-U73200CT2020PTC010704

THERAPEUTIC AREAS



Oncology
- Hemato Oncology
- Medical Oncology
- Surgical Oncology
- Radiation Oncology
- Immuno Therapy etc.

Cardiology
- Hypertension, IHD
- MI, Cardiac Arrest
- Cardio Surgery etc.

General Medicine
- DM Type 1&2
- Hematology
- Hypertension
- Anaemia
- Infectious diseases etc. .

Neuro-psychiatry
- Neurology
- Psychiatry Disorders etc.

HepatoGastroenterology
- IBD (Ulcerative Colitis & Chron's)
- Peptic Ulcer
- Liver disorder (Hepatitis B & C)
- GERD, Gastric Ulcer
- Constipation etc.

Uro-Nephrology
- Urology
- Nephrology
- CKD & AKD etc.

Dental
- Cavities
- Gingivitis (Gum Disease)
- Oral Cancer
- Periodontitis
- Sensitivity etc.

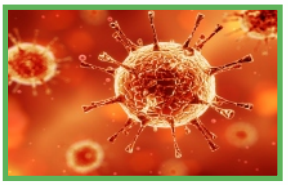
Dermatology
- Skin Cancer
- Eczema
- Acne
- Dermatitis
- Psoriasis
- Vitiligo etc.

Pediatrics & Neonatology
- Bronchitis
- Common Cold
- Immunization etc.

Orthopedic
- RA
- OA
- Osteoporosis
- Surgeries etc.

Respiratory Medicine
- Asthma
- COPD
- Intensive Care etc.

Ophthalmology
- Glaucoma
- Diabetic Retinopathy
- Cataract etc.



Immunology & Infectious Disease
- Viral Infection
- HIV
- Covid-19
- Dengue etc.

Gynecology
- PCOD
- Endometriosis
- Uterine Fibroids
- Vaginitis etc.

Note: Our research areas are not limited to above mentioned specialties...

WHO CAN CONTACT US

- CROs
- Sponsors
- Hospitals
- Clinicians
- Ethics Committees
- Educational Institutes

WHO WE ARE?

nothing beyond humanity

KV Clinical Research Services (KVCR) is a Site Management Organization (SMO) established in the year 2015. KVCR acts as a common platform between Principal Investigators/study sites and study Sponsors/CROs for the seamless execution of clinical trials. KVCR focuses on growth and development of drugs for various therapeutic areas in all phases of clinical trials in India. KVCR is built by passionate and experienced individuals working along with our clients to conduct clinical research for delivering new drugs and medical devices to the market while maintaining high quality of data.

OUR MISSION

To offer clinical trial services from Phase I - IV in key therapeutic areas, strictly compliant with ICH-GCP & NDCTR guidelines. Our multi-site investigator networks next to our well trained team, blend into a unique "tailor made solution" to match the needs and expectations of CROs and sponsors.
To collaborate with clients and to develop a better future for the community by bringing our expertise on drug development through the commitment of our expert clinical research professional.

SOLUTIONS

Our extensive database of Principal Investigators, leading hospitals, and research centers allows us to access special populations to fulfill protocol requirements and ensure rapid recruitment of eligible study subjects.

Our SOPs meet the highest standards for conducting clinical trials. We manage our sites with highly qualified personnel, experienced in conducting clinical trials while adhering to applicable regulations & guidelines. This includes ICH GCP, NDCT-2019.

OUR SERVICES

Providing services to CRO, Sponsors, Hospitals and Clinicians to facilitate the clinical trials. Also helps hospitals to set up the Clinical Research Department.

Clinical Operation Services
- Study start-up including potential sites and experienced PI suggestions.
- Study documents submission to Ethics committee.
- Feasibility of studies and initial documents preparation at site level.
- Subject recruitment & follow-up.

Clinical Data Management

We offer fast turnaround and a flexible, efficient process for any project, including :
- Data entry in paper CRF or eCRF
- Data submission on fast-track process to CRO/Sponsor
- 24/7 available for SMO services
- Document archival facility as per regulatory guidelines

Manpower Management Services

We provide following trained manpower :
- Clinical Research Coordinators
- Nurses
- Phlebotomists
- Conducts annual and proper training for Research
- Professionals including Investigators.

Ethical Committee related Services
- IEC formation and DCGI registration
- SUGAM Portal formation
- IEC Re-registration
- NAITIC Portal formation and Registration
- NABH Accreditation for IEC
- ICH-GCP training programs for IEC Members.

Note: Having experience of all the regulatory including USFDA, DCGI , EMA & TGA study and inspections

RESEARCH AREAS ON VARIOUS SYSTEM OF MEDICATIONS

Herbal & Ayurveda
Department of Ayush
Completed Several Studies
Ayurvedic Hospitals
& Clinicians

Allopathy
CDSCO (DCGI)
Completed 250+ Studies
Various superspeciality &
Multispeciality Study sites

Homeopathy
Department of Ayush
Homeopathy Hospitals

Medical Devices
CDSCO (DCGI)
Completed Several Studies
Study Sites

TYPES OF CLINICAL TRIALS

Phase I

Completed 1
Phase 1 Study

Phase II
Completed 2
Phase II studies.
Having a good
database of
experienced PIs.

Phase III & IV
Completed 200+
studies of Phase III & IV.

BA/BE Studies
Completed 45+
PK studies.

Note: Having the experience of 35 Covid19 Studies including vaccine studies

STRENGTHS

COMPLETE SOLUTION AT ONE PLACE

- **Having alliance with hospitals across India**
- **Expedited recruitment of study subjects.**
- **Expert Advisors & Clinical Research Professionals**
- **High Quality of Data**

LOCATIONS OF SITES

(KVCR is collaborated with 36 hospitals across India)

North (9 Sites)
- Delhi
- Varanasi
- Dehradun
- Ludhiana
- Chandigarh
- Patna

East (13 Sites)
- Kolkata
- Guwahati
- Bhubaneswar
- Raipur
- Ranchi

South (7 Sites)
- Bangalore
- Chennai
- Nellore
- Mysore
- Belagavi

West (7 Sites)
- Ahmedabad
- Vadodara
- Nagpur
- Nashik
- Indore
- Bhopal

HEAD OFFICE:
616-617, Sixth floor, Golden Trade Center,
New Rajendra Nagar, Raipur,
Chhattisgarh-492001, India

E-mail:
md@kvclinicalresearch.com
kpatel@kvclinicalresearch.com

Contact
94273 98575 | 90399 39374
94083 88382 | 87580 36302