veeda clinical research®

Empowering Your Clinical Trials from Start to Finish:

Discover Our Comprehensive Services

Overview

As a trusted partner to pharma and biopharma companies around the world, we are committed to bringing life-saving medicines to market through expertly managed clinical trials. With a team of experienced professionals and in-depth capabilities, we offer comprehensive end-to-end support throughout the clinical trial process. From protocol development and site selection to regulatory compliance and data analysis, our proven expertise enables us to deliver high-quality and timely outcomes for your clinical studies. With our tailored approach and commitment to excellence, you can trust us to help you achieve your drug development goals and make a meaningful impact on patient health.

Clinical Trial Services

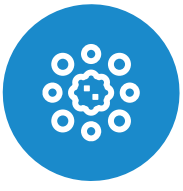
- Medical Writing - Protocol, ICF, IB, Study Report etc
- Conducting Feasibility & Site Set up activity
- Regulatory Services Application processing Technical presentation -Liasioning
- Site Monitoring, Project Management & Safety Monitoring
- Pharmacy and Laboratory services including PK and Immunogenicity analysis capabilities
- Data management, Biostatistics including eCRF capabilities
- Safety Database and Pharmacovigilance

Our Patient Trials Capabilities

40+ Completed Patient Studies	**4000+** Patients Enrolled	**250+** Sites	**900+** Investigators Database	**16+** Ongoing Studies (Including Phase I / II / IIa studies, Patient PK & Clinical Endpoint)	Successfully completed **21** USFDA & **2** EMA inspections across sites without 483 observations

Expertise That Extends Across Multiple Therapy Areas

Oncology



Psychiatry

Gastroenterology

Cardiology

Rheumatology

ENT

Dermatology

Respiratory

Endocrinology

Ophthalmology

Gynaecology

Infectious Diseases

Veeda's Experience Across Various Phases of Drug Development Lifecycle

POC Study / Phase I Study

- HIV
- Covid -19 Vaccine (Phase I / II)
- Colon or Pancreatic Cancer

Phase II / III

- Relapsed Advanced Tumors and classical Hodgkin Lymphoma (cHL)
- Asthma
- COPD
- Human Head Lice Infestation

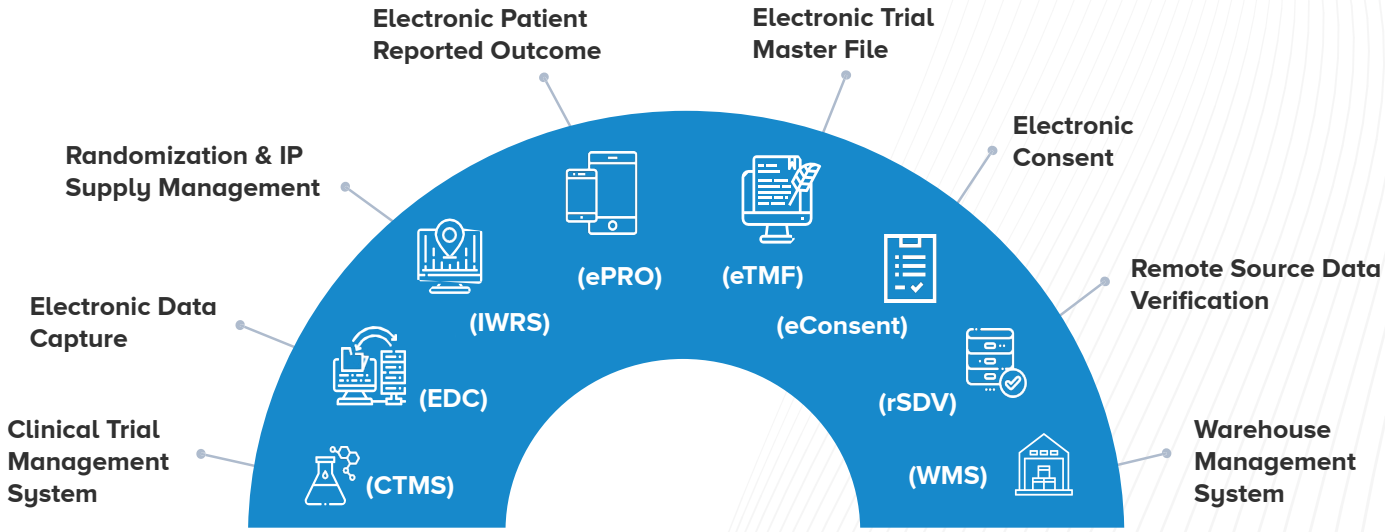
Clinical End Point

- Open Angle Glaucoma
- Chronic Idiopathic Constipation

Patient PK

- Multiple Myeloma
- Postmenopausal Osteoporosis
- Schizophrenia
- Advanced Prostate Cancer
- Ovarian Cancer
- Breast Cancer
- Iron Deficiency Anemia
- Renal Carcinoma
- Small Cell Lung Cancer
- Rheumatoid Arthiritis

Access to e-Clinical Platforms - Designed to help Research Sponsors get studies up & running quickly



Robust & Industry Leading Data Management Support

- Database & eCRF Design, Edit Checks, Programmed Dynamics, & UAT
- EDC Platform development: Medrio, Main EDC and Octalsoft
- Medical Coding using MedDRA and WHODrug Global
- Data Quality, Data Cleaning, & Query Resolution
- SAE Reconciliation/External Data Reconciliation
- SAS-certified Statistical Programmers for validation of the results
- External Vendor & Project Team Management
- Document Creation including DVS, DMP & Data Entry Guidelines
- Audit Readiness & Study Documentation Management

Our Approach for Patient Recruitment, Retention & Site Selection

We can help you identify the best strategy to mitigate costs, risks, and identify the best sites providing meaningful insights into your recruitment challenges, evaluating the impact of competitive trials, and optimizing patient access and retention.

- **Sound understanding of operational nuance in clinical studies** including site and patient-level considerations
- **Rapid study start-up** with our insights-driven site selection and feasibility services together with our site budgeting and contracting services solve common trouble spots in the start-up of clinical trials
- **Patient recruitment and retention** by combining a dedicated study team with a customized site recruitment strategy, enrolment becomes focused, effective, and streamlined so that clinical studies begin and stay on schedule
- **Relationship with leading hospitals, key opinion leaders, and investigators,** which can support recruitment strategy

Why Choose Veeda for your Next Clinical Trial Program?

- Establish site network, enabling timely recruitment
- Experienced study team with a deep understanding of scientific and operational considerations to mitigate study challenges
- Monitoring strategy to ensure high quality and compliance
- Proven history of completing studies within the agreed timeline and cost

To know more about

our expertise in Clinical Trials, mail us at

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Partners in creating a healthier tomorrow

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