Our Services

We offer outsourced services:

- Study and Site Management
- Clinical Trials Monitoring and Auditing
- Clinical Trials Project Management
- Investigator Site Feasibility and Selection
- Regulatory Affairs
- Clinical Research Training
- Health Systems and Outcomes Research
- Contract Laboratory Services Management
- Pharmacovigilance services
- Strategic Consulting services

Our Experience

Our team has extensive experience in: infectious and non-communicable diseases, Projects Management in resource limited settings (Sub-Saharan Africa and Asian countries).

We offer Phase I - IV capabilities as a full service project based or Functional Service Resourcing model.

Our Team

Our team is composed of:

- Clinical Trials Monitors and Auditors
- Public Health Experts and Epidemiologists
- Projects and Data Managers
- Laboratory Scientists
- Physicians/Medical Doctors
- Contracts and Grants Managers
- Statisticians
- Social Scientists
- Qualified Persons

CLINWIN RESEARCH SERVICES (CRS)

About us

CRS is a mid-sized Contract Research Organisation (CRO), providing outsourced clinical development and Consulting services to Biopharmaceutical companies, Academic Medical Centres, Investigator Sites, not-for profit research organisations and Government.

Our Vision
To deliver quality and efficient services on time and cost to our clients.

Our Mission
To partner with Biopharmaceutical, Academia and CRO clients to support the successful outcome of their projects.

Our Values
We advance our clients' assigned projects through integrity, teamwork, quality and accountability.

Our Principles
Efficiency, Quality and Ethics

Our Contacts

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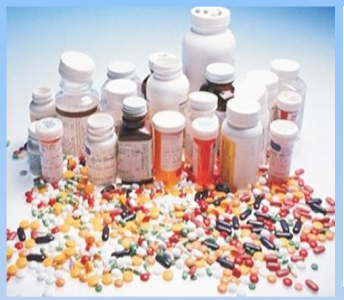
ClinWin Research Services

Efficiency. Quality. Ethics

Clinical Trials Monitoring and Consulting



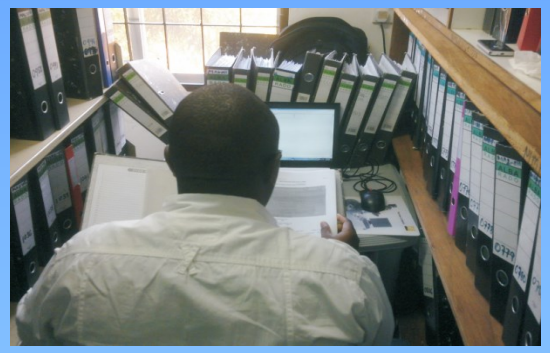
Biopharmaceutical

Technology

Data Science

Training

Clinical Trials Monitoring and Site Management



- Feasibilities and Site Selection
- Study start up and contracts negotiations
- Ethical & Regulatory submissions support
- Site Initiation, IMV & Close Out Monitoring
- Medical Monitoring
- Patient Recruitment
- Site Operations

Projects Management

- Projects feasibilities
- Risk Assessment and Management
- Regulatory and Ethical submission
- Project team management
- Resources planning and management
- Milestones tracking and reporting
- Budgets and Contracts management
- Project Communication Planning
- Quality Management
- Projects Monitoring and Evaluation

Strategic Consulting

At CRS we offer our clients innovative and cost efficient solutions to make informed business decisions. These include:

- Product development advisory services
- Protocol design and feasibility
- Operational efficiency and effectiveness
- Patient recruitment and retention services
- Risk and Compliance Operations
- Productivity and Performance Evaluations
- Contracts Negotiation and Legal Affairs
- Clinical Resourcing and Recruitment
- Document Translation service
- Market research and access

Health Systems Research

- Health Services effectiveness
- Social Science Research,
- Health Policy analysis,
- Health Services delivery,
- Health care organisational development
- Surveys and program evaluation

Clinical Research Training

We offer face to face and on-line courses to investigator site and Sponsor staff. The courses can be tailored as per client needs on all aspects of clinical research enterprise. This include:

- Entry and Advanced Level Clinical Trials Monitoring Course
- Research Project planning and Management
- Good Clinical Practice (ICH-GCP) for beginners and refresher courses
- Good Clinical Laboratory Practice (GCLP)
- Internal quality assurance
- Clinical Research Ethics
- Infectious Diseases
- Data Management
- Regulatory affairs
- Clinical Research Methodology
- Projects Management