CRK-CRO ®
Clinical Research Key

St. Mary's Mission Hospital,
Nairobi, Kenya
+254 740410277
www.crk-cro.com
contact@crk-cro.com

CRK

Clinical Trial Solutions

ADVANCING HEALTH THROUGH **ETHICAL RESEARCH.**

Our Solutions

Explore our comprehensive tailored approach for Biotech, Pharma and Medical Device companies in need of exceptional clinical trial management.

- Late Phase Clinical Trial
- Post Marketing Trials
- Medical Devices & Nutraceuticals
- Biosimilar & Generics

Efficient and Cost-effective Clinical Trial Solutions in Kenya

Our dedicated team excels at designing and carefully planning each phase of the medical development for exceptional therapies.

Leveraging our CRO expertise and resources, we are committed to advancing your therapeutic innovation, ensuring its success from concept to market.

At CRK, we provide efficient, flexible, and reliable clinical trial solutions for every phase of your clinical research. We offer the best value CRO and SMO solutions that significantly reduce clinical trial costs without compromising quality.

Explore our comprehensive, tailored approach for Biotech, Pharma, and Medical Device companies in need of exceptional clinical trial management.

Late Phase Clinical Trials

Partner with us to access the top recruitment and retention rates in Kenya and benefit from our efficiency and speed in the Late Phase clinical trials recruitment process.

Optimized Late Phase Clinical Trial Solutions

Our extensive experience makes us rapidly identify the right investigators and patient populations, ensuring efficient enrollment and achieving recruitment goals. We prioritize clear and continuous communication, providing regular project updates while strictly adhering to timelines and budgets.

Comprehensive Late Phase Clinical Development Services

We offer tailored late-stage clinical trial solutions for innovative investigational drugs and biologics across various therapeutic areas. Whether you require stand-alone services or a full-service solution, we are committed to delivering customized support that meets your specific needs and drives the success of your project.

Late Phase Clinical Trials

Your Trusted Partner in Late-Phase Trials

As your strategic partner, we help develop the right clinical trial strategy to:

- **Reduce costs**
- **Mitigate risks**
- **Ensure global regulatory compliance**

Recruitment and Retention

We take a patient-centric approach, leveraging our strategic location in the CEE region, known for its diverse patient population. Through targeted prioritization strategies, we help you meet enrollment goals, enhance trial awareness, and maintain participant engagement.

Managing Your Late Phase Trials

Our detailed and balanced trial management approach ensures the timely achievement of project milestones. We develop and oversee a comprehensive Project Management Plan, maintaining control of study execution and risk management with rapid issue identification and communication.

02 Post Marketing Trials

At CRK Clinical Research Key, our post-marketing trials provide essential insights into long-term patient safety and compliance, supporting your medical product's success in the market

Post-Marketing Trial Services for Drug Safety & Effectiveness

Post-market trial services are **essential for monitoring a drug's safety and effectiveness** once it has entered the market. **Phase IV studies** help identify **unexpected effects**, track **long-term efficacy**, and ensure **ongoing product quality**.

Expertise in Non-Interventional & Phase IV Studies

At **CRK Clinical Research Key**, we specialize in managing **non-interventional studies**, offering **customized Phase IV clinical trial solutions** for **evidence-based market management**.

Comprehensive Post-Marketing Monitoring

- **Pharmacovigilance & Post-Approval Studies**
  We **monitor and analyze data** on **unexpected side effects and adverse drug reactions**, ensuring **long-term safety assessments** and conducting **further clinical research** as needed.

- **Registry Studies for Long-Term Insights**

02 Post Marketing Trials

Our **registry studies** maintain **detailed records of patient outcomes**, providing **valuable real-world data** for continued drug evaluation.

Studies We Advocate:

We advocate for **comprehensive and impactful studies** that drive **scientific advancements** and **enhance patient outcomes**.

- Safety Surveillance Studies
- Efficacy Studies
- Comparative Effectiveness Research (CER)
- Observational Studies
- Registry Studies
- Pharmacoepidemiological Studies
- Label Expansion Studies
- Behavioral and Quality of Life Studies
- Compliance and Adherence Studies
- Real-World Evidence (RWE) Studies
- Post-Approval Commitments
- Biomarker and Genetic Studies
- Long-Term Follow-Up Studies

Medical Devices & Nutraceuticals

At CRK Clinical Research Key, we are committed to the design and clinical evaluation of medical devices and nutraceuticals, ensuring that your products meet the highest standards of safety, efficacy, and market readiness.

Comprehensive Medical Device & Nutraceutical Development

At **CRK Clinical Research Key**, we provide **end-to-end solutions** for the **clinical evaluation and development** of **medical devices and nutraceuticals**. From **regulatory consulting and clinical trial design** to **data management and post-market surveillance**, we offer **complete support** throughout the process.

Ensuring Quality & Compliance

We conduct **rigorous testing and validation** to ensure compliance with **global safety and efficacy standards**. Our **bioanalytical laboratory services** and **strict quality assurance protocols** guarantee **credible and reliable results**.

Your Trusted Partner in Product Commercialization

By partnering with us, you gain access to **expert guidance** in navigating **complex regulatory landscapes**, allowing you to **efficiently bring innovative products to market**.

Medical Devices & Nutraceuticals

What We Do

- Regulatory Consulting
- Clinical Trial Design & Management
- Data Management & Analysis
- Bioanalytical Laboratory Services
- Quality Assurance
- Post-market surveillance

Biosimilar & Generics

At CRK Clinical Research Key, we offer expert guidance in Biosimilar and generic development, ensuring cost-effective and efficient pathways to high-quality, safe, and effective clinical studies at every stage.

Comprehensive Biosimilar & Generics Development Support

If your goal is to develop **cost-effective alternatives** to existing **biological drugs (Biosimilars)** or **small-molecule drugs (Generics)**, **Comac Medical** provides **comprehensive support** and a wide range of **expert biosimilar and generic solutions**. Our team will **craft a tailored strategy** to **mitigate risks** and ensure a **smooth, cost-efficient clinical development process** for your product.

Expertise in Global Biosimilar Clinical Plans

We specialize in **designing and executing global biosimilar clinical strategies**, ensuring **timely delivery and high-quality results**. Our **key services** include:

Gap analysis for comparative analytical similarity assessments
Drug development consulting
Strategic advisory for biosimilar & generic development

Biosimilar & Generics

Enhancing Efficiency & Reducing Risk

We provide:

- **Clinical pharmacology expertise**
- **Bioanalytical laboratory services**
- **Operational support for product authorization**

All our solutions are designed to **streamline processes**, **enhance efficiency**, **reduce costs**, and **minimize risks** in biosimilar and generic development.

Key Services

- Create and execute global biosimilar clinical strategies
- Ensure timely delivery and high-quality outcomes
- Conduct gap analysis for analytical similarity testing
- Provide drug development consulting and strategic advice
- Offer clinical pharmacology expertise
- Support for Pharmacokinetics and Pharmacodynamics (PKPD)
- Deliver flexible services to enhance efficiency
- Bioanalytical laboratory testing
- Consulting support for product authorization