Advantages of conducting clinical trials in South Africa

- Established regulatory agencies and Ministries of Health. The South African Health Products Regulatory Authority (SAHPRA) was established in 1965 and provides for the monitoring and evaluation, regulation, investigation, inspection, registration and control of medicines, scheduled substances, clinical trials and medical devices, in vitro diagnostics and further matters related to the public interest.
- SA-GCP is based on ICH-GCP and provides added protection for vulnerable populations
- Established Research Ethics Committees
- Sites are ICH-GCP compliant
- Experienced clinical trial sites
- Multinational trial experience, that means sites are familiar with FDA requirements
- Patient recruitment is characteristically faster than in the US and Western Europe
- Diverse patient population
- Emerging and first world disease indications
- Seasonal reversal, with respect to the Northern Hemisphere, allows companies to shorten clinical trial timelines

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Your **Partner** In Africa acro african clinical research organisation transforming ideas into solutions

We are African
We know Africa
We live Africa
We love Africa



ACRO, African Clinical Research Organisation, a private company launched in August 2007, was the first South African, full-service, BEE contract research organisation.

ACRO, Offers services in Southern Africa and other African regions and is active across a broad range of clinical areas including, but not limited to:

> **Ophthalmology** Metabolic

Respiratory CNS

Vaccines Infectious Diseases Oncology **Medical Devices Endocrine Wound Care and** Other rare diseases Renal

An important aim of the organisation is to help improve the care of those affected by HIV/AIDS, tuberculosis, malaria and other diseases prevalent in other African populations.

ACRO is committed to capacity building across Southern Africa and has formed strategic development partnerships with a solid base in South Africa and an African emphasis.

ACRO employees have experience in a multinational environment and clients are guaranteed that their trials will be conducted to international standards, with competitive local South African costing.

ACRO is ideally positioned to "transform ideas into solutions".

ACRO offers services to donor-funded organisations, non-governmental organisations, research institutions, academic institutions, pharmaceutical companies, medical device companies, complimentary medicine, food product, cosmetic companies and government institutions.

ACRO works both within and outside the biotechnology sector.







Clinical Trial Site

site staff.)

of site administrative

Development (including site

identification, establishment

and physical infrastructure,

as well as GCP training of

Our Service Coverage in Africa

Regulatory Consulting and Submissions



Training:

- GCP Intro and Refresher
- CRA, CTA & PM Training
- Data Management Training
- Customized Training

Additional and updated information about these and other services on offer are provided on the ACRO website: www.gcro.co.za