CAPRISA

CENTRE FOR THE AIDS PROGRAMME OF RESEARCH IN SOUTH AFRICA

Clinical Research Associates (Freelance Consultants)

CAPRISA is inviting applications for suitably qualified and experienced Clinical Research Associates to form part of a panel of freelance CRA's, to coordinate all aspects of the clinical monitoring process in accordance with all relevant regulations and legislation, study processes and procedures.

These positions will involve conducting site visits (eThekwini and Vulindlela) to assess protocol, regulatory and good clinical practice (GCP) compliance and manage required documentation to meet quality assurance audits for clinical trials.

Essential functions: Evaluate the quality and integrity of study site practices related to the protocol and adherence to applicable regulations.
Create and maintain appropriate documentation regarding site management, monitoring visit findings by compiling visit reports and other required study documentation.
Review the progress of studies by monitoring regulatory submissions and approvals, case report form (CRF) completion and submission, and data query generation and resolution.
May support start-up and close-out phase.

Qualifications and experience:
Degree in health sciences, or a certified health care professional.
2-5 years of experience as a clinical research monitor, conducting independent on-site monitoring in all types of monitoring visits from Phase I - IV studies.
Excellent understanding and demonstrated application of ICH and SA GCP and applicable Standard Operating Procedures.

Personal qualities and competencies: Ability to organize and manage multiple priorities; Attention to detail/Quality orientation; Communication and collaboration; Ability to work under pressure.

Period: 1 year consulting assignment

To apply send a detailed CV and cover letter to caprisa@caprisa.org

Closing Date: 4 March 2021