Job Title: Senior Clinical Research Associate (CRA)

Company: ACE Research – African Contract Research Organization (CRO)

Location: Must be in an African Country

Job Summary: ACE Research, a leading African full-service Contract Research Organization (CRO), is seeking experienced Senior Clinical Research Associate (CRA) contractors to join our dynamic team. As a Senior CRA at ACE Research, you will play a pivotal role in the planning, execution, and monitoring of clinical trials. Your expertise will ensure the successful completion of projects while maintaining the highest ethical and regulatory standards.

Key Responsibilities:

1. **Site Management:** Oversee and manage clinical trial sites, ensuring adherence to protocols, GCP, and regulatory requirements.
2. **Monitoring:** Perform site visits, including pre-study, initiation, interim, and close-out visits, to ensure data integrity and protocol compliance.
3. **Site Training:** Provide training and guidance to site staff on study protocols, procedures, and compliance with regulatory requirements.
4. **Documentation:** Maintain accurate and organized trial documentation, including essential documents, case report forms, and source documents.
5. **Quality Assurance:** Ensure data quality and integrity through regular data review and verification.
6. **Risk Management:** Identify potential risks and develop risk mitigation strategies.
7. **Reporting:** Prepare and submit monitoring reports, addressing findings and actions taken.
8. **Regulatory Compliance:** Stay up-to-date with local and international regulations, ensuring studies are conducted in compliance with these standards.
9. **Team Collaboration:** Collaborate with cross-functional teams, including project managers, data managers, and sponsors, to ensure successful trial execution.
10. **Relationship Building:** Foster positive relationships with investigational sites, clients, and regulatory authorities.

Qualifications:

- Bachelor's degree in life sciences or related field (advanced degree preferred).
- Minimum of 5 years of experience as a CRA in clinical research.

- In-depth knowledge of ICH-GCP guidelines and regulatory requirements.
- Strong communication and interpersonal skills.
- Ability to work independently and as part of a team.
- Detail-oriented with excellent organizational and time-management abilities.
- Proficiency in using clinical trial management systems (CTMS) and electronic data capture (EDC) tools.
- Willingness to travel to clinical trial sites as needed.

Why ACE Research:

- Join a dynamic and growing CRO committed to excellence in clinical research.
- Work with a collaborative and supportive team of professionals.
- Competitive compensation and benefits package.
- Opportunities for career growth and development.

If you are a dedicated and experienced CRA looking to take the next step in your career, apply today to be part of our mission to advance healthcare through clinical research at ACE Research.

How To Apply:

Submit your CV through our website.