JOB ADVERTISEMENT

Position: Site Clinical Researcher for BREATHE Project

Job description

Sponsored by Welcome Trust, BREATHE is a multi-sites and multi-country evaluation being implemented in Rwanda, Kenya and Malawi. The project is a Type 1 implementation-effectiveness hybrid clinical trial design, with primary aim of determining the effectiveness of clinical interventions and secondary aim of determining the feasibility and utility of intervention strategies. The interventions include both protocolized standard of care Low Flow oxygen with monitoring and titration, and protocolized High Flow Nasal Canula with monitoring and titration for adults with hypoxemia of any cause. BREATHE is a 3 years project and its implementation in Rwanda is coordinated by CIICHIN.

CIICHIN is looking for qualified candidates for the position of clinical researcher for BREATHE project and to be based at CHUK and CHUB respectively.

The candidate will implement research protocol at his/her respective site. The clinical researcher is responsible for overseeing and ensuring all specifications are being followed on her/ his site and will be responsible of daily project activities

Duties

- Implement study activities to ensure compliance with protocols and with all relevant local and international regulatory and institutional policies.
- Assess the eligibility of potential subjects through methods such as screening interviews, reviews of medical records, and discussions with physicians and nurses.
- Enroll and follow up each eligible participant based on the protocol in place
- Participate in quality assurance audits conducted by study sponsors, or specially designated review groups.
- Anticipate and report any protocol problem, inform the site coordinator of problems, and assist in problem resolution efforts such as protocol revisions.
- Track enrollment status of subjects and document related information
- Ensure proper, appropriate and recommended storage of records of study activity including case report forms, device dispensation records, or regulatory forms.
- Participate in capacity building workshops/ trainings on standards of care, documentation procedures, continuing education activities etc.
- Conduct subject enrollment to ensure that informed consent is properly obtained and documented.
- Gather metrics from cross-functional study sites in relation to project progress

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Requirements – Abilities, Skills, and Knowledge

- Must possess excellent communication skills, both verbal and written are required so as to be able to effectively convey messages across to both study team members and other external persons
- Good problem-solving, organizational, and leadership skills are highly required
- Ability to work with little or no supervision
- Ability to pay keen attention to detail at all times
- Must be able to work within stipulated timeframes or deadlines even in the face of multiple responsibilities or projects
- Must be able to work as part of a team or in most cases lead a team
- A bachelor degree in general nursing and/or a bachelor degree in clinical medicine
- Sufficiently trained on Good Clinical Practice (GCP)
- A minimum of 2 years' experience in a scientific or health related field of which 1 year must have been worked in clinical setting, clinical research setting.
- Proficient user of relevant computer applications for the execution of daily project operations

How to apply

To apply, please send the following not later than Monday, June 26th, 2023 5:00 PM by e-mail to: info@ciichin.org, cc numutors@ciichin.org and management@ciichin.org

- A motivation letter addressed to CIICHIN CEO
- A copy of academic qualification,
- A CV (maximum two pages) with at least 2 references
- A copy of Rwandan National Identity Card;

Please include “Clinical Researcher / BREATHE Project” in the subject line of the application e-mail.

- Note: Incomplete and late application will not be considered.

Jeanine CONDO, MD, MSc, PHD
CEO CIICHIN

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