Clinical Research Associate (CRA)/ Respiratory Therapist (RT) with Clinical Trial Experience for Highly Innovative Product

Per Diem, Nort-Eastern Region, USA

About the Innovation

STIMIT 's mission is to empower patients to breathe, to achieve a breakthrough in critical care.

Patients on mechanical ventilation today do not breathe actively on their own, and their core breathing muscle – the diaphragm – deteriorates. STIMIT's innovative technology has been developed to stimulate the core breathing muscle – the diaphragm – of intensive care patients.

STIMIT is currently starting it's FDA IDE trial.

About the Position

For this novel treatment, we are looking for a per-diem/ temporary CRA with on-site CRA/respiratory therapy experience, for an FDA IDE Trial in the United States.

Applicants must have experience with medical devices relating in Critical Care, experience in FDA trials. Previous patient care experience as an RT or R.N. working in critical care, or related hospital settings are an important asset.

This position will travel up to 10 days in a row (mainly New York – Boston – Philadelphia area).

CRA role with significant responsibility and visibility within the organization and within the top scientific community. If your experience meets these requirements and you're looking for a challenging and well compensated new Sr. CRA/ RT role, please forward your CV.

Your Responsibilities

This position drives and monitors progress of clinical studies at the site level to verify that the rights and well-being of subjects are protected, that the reported study data are accurate, complete, and verifiable, and that the study is conducted in accordance with protocol, standard operating procedures, ISO/GCP/ FDA requirements.

- On-site clinical site support: accompany applications of novel pre-market medical device.
- Communication & Project Management: Ensure clear communication through monitoring reports, follow up letters, study memos, and study correspondence.
- CRA activities: Case report form review, source verification, verify informed consent forms, review regulatory documents and device accountability records. Issue, investigate and resolve data discrepancies. Proper documentation.
- Closely collaborate with cross functional team members and scientists and study sites throughout all study phases.

Qualifications

- BS/BA required
- CRA clinical trial experience
- Critical Care experience (ideally RT, ICU nursing or extensive prior ICU trial experience) is a plus

Contact

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