

Senior Clinical Research Associate (SCRA)

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Company: Randstad

Location: Hong Kong

Category: other-general

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Our client, a world leader CRO, offers world class services in clinical trial logistics, development and manufacturing of the drug product. Looking to expand their clinical research team to contribute to the future of drug discovery. Less experience will be tilted as Clinical Research Associate.

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Monitors investigator sites with a risk-based monitoring approach.

Assess investigational product through physical inventory and records review.

Documents observations in reports and letters in a timely manner using approved business writing standards.

Escalates observed deficiencies and issues to clinical management expeditiously and follow all issues through to resolution.

Conducts monitoring tasks in accordance with the approved monitoring plan. Participates in the investigator payment process. Ensures a shared responsibility with other project team members on issues/findings resolution. Investigates and follows-up on findings as applicable.

Participates in investigator meetings as necessary. Identifies potential investigators in collaboration with the client company to ensure the acceptability of qualified investigative sites.

Initiates clinical trial sites according to the relevant procedures to ensure compliance with the protocol and regulatory and ICH GCP obligations.

skills & experiences required.

Bachelor's degree in a life sciences related field or a Registered Nursing certification or equivalent and relevant formal academic / vocational qualification.

Previous experience that provides the knowledge, skills, and abilities to perform the job (comparable to 1 years as a clinical research monitor).

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skills

no additional skills required

qualifications

no additional qualifications required

education

Bachelor Degree

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