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Senior Clinical Research Associate Fsp

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Company: Hays

Location: Australia

Category: other-general

Senior CRA | FSP experience | Early phase oncology trials | Chinese language skills desired | Anywhere in AU Location: this vacancy is open to candidates based either in Sydney, Melbourne, Brisbane or Adelaide.

Your new company

A leading global CRO is expanding its clinical team! This mid-sized CRO is seeking a CRA to work on a brand new FSP project, the first of its kind in the organisation. Headquartered locally in Sydney CBD, this global CRO boasts having 30+ locations worldwide. Working directly on a Biotech's trials you will play a pivotal role in ensuring studies are conducted in accordance with ICH-GCP, protocol and sponsor requirements. This CRO prioritises reinvesting some of it's profits into the innovation of solutions for improving the health of developing patient populations and challenging healthcare issues. This CRO is proud of its ethical motivations and in-house network of highly experienced clinicians.

Your new role As a Clinical Research Associate/Senior Clinical Research Associate you will be responsible for ensuring assigned clinical projects are conducted in accordance with ICH-GCP, protocol and sponsor expectations. You will be involved in the site set up, routine monitoring and close out of mainly Phase I oncology studies. You will liaise directly with study sites partnering with clinical research nurses and clinicians to ensure study compliance.

Ensure ethics submissions are made by investigator sites in a timely manner. You will contribute to improving the health of patients globally and establishing treatment solutions for some of the more challenging diseases. What you'll need to succeed You will be an experienced CRA with 3+ years of independent monitoring experience in the commercial clinical research industry, ideally within an FSP model. You will enjoy monitoring, travelling to study sites and liaising directly with site staff. You will have availability to travel interstate. Previous experience working in early phase Oncology trials. Chinese language skills are advantageous. Excellent communication, interpersonal and influencing skills. You will be a self-motivated research professional who thrives on working in a fast-paced environment. Moreover, you will have professional experience of study start up straight through to close out. What you'll get in return Flexible working arrangements - you will have the benefit of working onsite, hybrid or WFH (depending on location). Competitive salary package, a supportive and friendly team environment and flexible working. You will be joining an organisation focused on employee wellbeing that prides itself on providing long term opportunities for career development. You will have the opportunity to work on complex and cutting-edge clinical trials. What you need to do now

If you're interested in this role, click 'apply now' to forward an up-to-date copy of your CV to , or call us now on . If this job isn't quite right for you but you are looking for a new position, please contact us for a confidential discussion on your career.

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