

## FSP CRA II (Nanchang/Haikou/Shenzhen)

[Apply Now](#)

Company: Fortrea

Location: Guangzhou

Category: business-and-financial-operations

As a leading global contract research organization (CRO) with a passion for scientific rigor and decades of clinical development experience, Fortrea provides pharmaceutical, biotechnology, and medical device customers a wide range of clinical development, patient access and technology solutions across more than 20 therapeutic areas. With over 19,000 staff conducting operations in more than 90 countries, Fortrea is transforming drug and device development for partners and patients across the globe.

### **Essential Job duties**

The below statement are intended to describe the general nature of the job and are not intended to be an exhaustive list of all responsibilities, skills and duties.

Responsible for all aspects of study site monitoring including routine monitoring and close-out of clinical sites, maintenance of study files, conduct of pre-study and initiation visits; liaise with vendors; and other duties, as assigned

Responsible for all aspects of site management as prescribed in the projects plans General on-site monitoring responsibilities

Ensure the study staff who will conduct the protocol have received the proper material and instructions to safely enter patients into the study

Ensure the protection of study patients by verifying that informed consent procedures and protocol requirements are adhered to according to the applicable regulatory requirements

Ensure the integrity of the data submitted on Case Report Forms(CRFs) or other data

collection tools by careful source document review Monitor data for missing or implausible data

Ensure the resources of the Sponsor and Fortrea are spent wisely by performing the required monitoring tasks in an efficient manner, according to SOPs and established guidelines, including managing travel expenses in an economical fashion according to Fortrea travel policy

Ensure audit readiness at the site level Travel, including air travel, may be required and is an essential function of the job

Prepare accurate and timely trip reports

Responsible for all aspects of registry management as prescribed in the project plans

Undertake feasibility work when requested

Participate in and follow up on Quality Control Visit(QC) when requested

Recruitment of potential investigators, preparation of EC submissions, notification to regulatory authorities, translation of study related documentation, organization of meetings and other tasks as instructed by supervisor

Might be requested to work in a client facing environment

Track and follow up on Serious Adverse Event(SAE) reporting, process production of reports, narratives and follow up of SAE Independently perform CRF review; query generation and resolution against established data review guidelines on Fortrea or client data management systems as assigned by management

Assist with training, mentoring and development of new employees, e.g. co-monitoring

Coordinate designated clinical projects as a Local Project Coordinator (with supervision, if applicable), and may act as a local client contact as assigned Perform other duties as assigned by management

## **Experience**

### **Minimum required:**

At minimum of 1(One) year of clinical monitoring experience is preferred, or an equivalent combination of education and experience to successfully perform the key responsibilities

of the job can be substituted

Ability to monitor study sites independently according to protocol monitoring guidelines, SOP and local regulatory Guidelines

Have a full understanding of the Serious Adverse Event (SAE) reporting, process production of reports, narratives and follow up of SAEs

Good planning, organization and problem solving skills

Ability to work with minimal supervision

Good communication and interpersonal skills

Good analytical and negotiation skills

Computer competency

Fluent in local office language and in English, both written and verbal

Works efficiently and effectively in a matrix environment

### **Education/Qualifications/Certifications and Licenses**

#### **Minimum Required:**

University or college degree, or certification in a related allied health profession from an appropriately accredited institution (e.g., nursing licensure). An equivalent amount of experience can be substituted as appropriate.

Thorough knowledge of ICH Guidelines and understanding of local regulatory requirement

Thorough knowledge of monitoring procedures

Basic understanding of the clinical trial process

Valid Driver's License

Fortrea is actively seeking motivated problem-solvers and creative thinkers who share our passion for overcoming barriers in clinical trials. Our unwavering commitment is to revolutionize the development process, ensuring the swift delivery of life-changing ideas and therapies to patients in need. Join our exceptional team and embrace a collaborative

workspace where personal growth is nurtured, enabling you to make a meaningful global impact.

[Apply Now](#)

#### Cross References and Citations:

1. FSP CRA II (Nanchang/Haikou/Shenzhen) [MobileappjobsJobs Guangzhou](#)  
[Mobileappjobs](#)
2. FSP CRA II (Nanchang/Haikou/Shenzhen) [PsychologistjobsJobs Guangzhou](#)  
[Psychologistjobs](#)
3. FSP CRA II (Nanchang/Haikou/Shenzhen) [Teacherjobsnearme Jobs Guangzhou](#)  
[Teacherjobsnearme](#)
4. FSP CRA II (Nanchang/Haikou/Shenzhen) [UsajobscentralJobs Guangzhou](#)  
[Usajobscentral](#)
5. FSP CRA II (Nanchang/Haikou/Shenzhen) [AdminjobsJobs GuangzhouAdminjobs](#)  
[Adminjobs](#)
6. FSP CRA II (Nanchang/Haikou/Shenzhen) [Nyjobscentral Jobs Guangzhou](#)  
[Nyjobscentral](#)
7. FSP CRA II (Nanchang/Haikou/Shenzhen) [HongkongjobsJobs Guangzhou](#)  
[Hongkongjobs](#)
8. FSP CRA II (Nanchang/Haikou/Shenzhen) [Searchcanadajobs Jobs Guangzhou](#)  
[Searchcanadajobs](#)
9. FSP CRA II (Nanchang/Haikou/Shenzhen) [RetailjobsnearmeJobs Guangzhou](#)  
[Retailjobsnearme](#)
10. FSP CRA II (Nanchang/Haikou/Shenzhen) [Searchcanadajobs Jobs Guangzhou](#)  
[Searchcanadajobs](#)
11. FSP CRA II (Nanchang/Haikou/Shenzhen) [Marketingjobs Jobs Guangzhou](#)  
[Marketingjobs](#)
12. FSP CRA II (Nanchang/Haikou/Shenzhen) [Courtjobs Jobs GuangzhouCourtjobs](#)  
[Courtjobs](#)
13. FSP CRA II (Nanchang/Haikou/Shenzhen) [Jobssearch Jobs GuangzhouJobssearch](#)  
[Jobssearch](#)
14. FSP CRA II (Nanchang/Haikou/Shenzhen) [Searchukjobs Jobs GuangzhouSearchukjob](#)  
[Searchukjob](#)



15. FSP CRA II (Nanchang/Haikou/Shenzhen) [Neurologistjobs](#) [Jobs Guangzhou](#)  
[Neurologistjobs](#) ↗
16. FSP CRA II (Nanchang/Haikou/Shenzhen) [Uzbekistanjobs](#) [Jobs Guangzhou](#)  
[Uzbekistanjobs](#) ↗
17. FSP CRA II (Nanchang/Haikou/Shenzhen) [Thechatgptjobs](#) [Jobs Guangzhou](#)  
[Thechatgptjobs](#) ↗
18. FSP CRA II (Nanchang/Haikou/Shenzhen) [Gardeningjobs](#)[Jobs Guangzhou](#)  
[Gardeningjobs](#)↗
19. Fsp cra ii (nanchang/haikou/shenzhen) [Jobs Guangzhou](#) ↗
20. AMP Version of Fsp cra ii (nanchang/haikou/shenzhen) ↗
21. Fsp cra ii (nanchang/haikou/shenzhen) [Guangzhou Jobs](#) ↗
22. Fsp cra ii (nanchang/haikou/shenzhen) [JobsGuangzhou](#) ↗
23. Fsp cra ii (nanchang/haikou/shenzhen) [Job Search](#) ↗
24. Fsp cra ii (nanchang/haikou/shenzhen) [Search](#) ↗
25. Fsp cra ii (nanchang/haikou/shenzhen) [Find Jobs](#) ↗

Source:<https://cn.expertini.com/jobs/job/fsp-cra-ii-nanchang-haikou-shenzhen--guangzhou-fortrea-f445e3115f/>

Generated on: 2024-05-03 by [Expertini.Com](#)