



■CRA<FSP>

臨床開発モニターのご経験のある方は歓迎です。

募集職種

人材紹介会社

株式会社ジェイ エイ シー リクルートメント

採用企業名

非公開

求人ID

1588887

業種

CRO

会社の種類

外資系企業

雇用形態

正社員

勤務地

大阪府

給与

500万円 ~ 800万円

勤務時間

09:00 ~ 17:30

休日・休暇

詳細は求人ご紹介時にご案内いたします。

更新日

2026年04月30日 12:01

応募必要条件

キャリアレベル

中途経験者レベル

英語レベル

ビジネス会話レベル

日本語レベル

ネイティブ

最終学歴

大学卒：学士号

現在のビザ

日本での就労許可が必要です

募集要項

【求人No NJB2297517】

Job Description

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

Assures the implementation of project plans for the clients as assigned at the client office where you will be dispatched

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

General on Site Monitoring Responsibilities:

Ensure the study staff who will conduct the protocol have received the proper materials 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

Prepare accurate and timely trip reports

Manage small projects under direction of a Project Manager/Director as assigned

Serve as lead monitor for a protocol or project and may assist in establishing monitoring plans and trip report review as assigned

Review progress of projects and initiate appropriate actions to achieve target objectives

Organize and make presentations at Investigator Meetings

Participate in the development of protocols and Case Report Forms as assigned

Participate in writing clinical trial reports as assigned

Interact with internal work groups to evaluate needs resources and timelines

Act as contact for clinical trial supplies and other supplies (vendors) as assigned

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

Undertake feasibility work when requested

Conduct report and follow up on Quality Control Visit (CQC) when requested

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

Negotiate study budget with potential investigators and assist the Fortrea legal department with statements of agreements as assigned

Complete process of Serious Adverse Event (SAE) reporting process production of reports narratives and follow up SAEs

Independently perform CRF review; query generation and resolution against established data review guidelines on Fortrea or clinical data management system assigned by management

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

Co ordinate designated clinical projects as a Local Project Coordinator (with supervision if applicable) and may act as a local client contacts as assigned

Perform other duties as assigned by management

To be dispatched to the client for the project following the clients' working conditions (working hours/working places)

スキル・資格

- ・ 学士資格（薬学、医学、生物科学、看護などの分野尚可）
- ・ 臨床開発モニターとしての3年以上の経験（医薬品、医療機器）
- ・ GCP（Good Clinical Practice）の知識
- ・ 英語を使用することに抵抗感の無い方

会社説明

ご紹介時にご案内いたします