



【1000～1400万円】 ラインマネージャー

外資スペシャリティファーマでの募集です。臨床開発リーダー・臨床開発プロジェク...

Job Information

Recruiter

JAC Recruitment Co., Ltd.

Hiring Company

外資スペシャリティファーマ

Job ID

1587056

Industry

Pharmaceutical

Company Type

International Company

Job Type

Permanent Full-time

Location

Tokyo - 23 Wards

Salary

10 million yen ~ 14 million yen

Work Hours

09:00 ~ 18:00

Holidays

詳細は求人ご紹介時にご案内いたします。

Refreshed

April 16th, 2026 16:40

General Requirements

Career Level

Mid Career

Minimum English Level

Business Level

Minimum Japanese Level

Native

Minimum Education Level

Bachelor's Degree

Visa Status

Permission to work in Japan required

Job Description

【求人No NJB2352566】

CRAの採用・育成・配置を通じて質の高い試験実施を支援します。

治験実施施設の選定から開始、被験者登録、データ収集、問題解決までの監督を行い、GCPや規制に準拠したモニタリングの品質管理を担います。

施設スタッフとの良好な関係構築に注力し、試験目標達成をサポート。

チームのパフォーマンス評価や教育、予算管理も担当し、国内外の規制対応やPMDA査察支援も行います。

Required Skills

Qualification Required:

Computer Skills: Efficient in Microsoft Word Excel MS Project MS PowerPoint and Outlook

Education Required:

Bachelors degree or higher in a scientific or healthcare discipline preferred

A minimum of 5 6+ years of relevant clinical operations experience.

Other Qualifications: . . .

Associate CRM: 1 3 years of people management or project management in progressive clinical research within the biotech pharmaceutical sector CRO industry or relevant field

CRM: 3 years or above of people management and project management experience in progressive clinical research within the biotech pharmaceutical sector CRO industry or relevant field

Senior CRM: 5 years or above of people management and project management experience in progressive clinical research within the biotech pharmaceutical sector CRO industry or relevant field

CRM/Senior CRM: Solid understanding of career development and performance management activities

Documented training knowledge and application of current Regulations GCP and ICH guidelines in clinical trials required

Evidence of team leadership capabilities

Therapeutic or medical knowledge preferred

Exhibits a strong understanding of methodologies and approaches

Understanding of all aspects of monitoring and trial execution

Excellent site management capabilities with demonstrated capability to problem solve and mediate complex compliance issues.

Thorough understanding of the international aspects of drug development process including expert knowledge of international standards (GCP/ICH) health authorities local/National Health Authorities regulations and BeiGene standards

Strong written and verbal communicate effectively with site personnel country and global associates

Computer skills including proficiency in use of Microsoft office

Excellent organization skill and management of competing priorities

Company Description

ご紹介時にご案内いたします