CareerCross

グローバルプロジェクトマネージャー　【関東窓口】

臨床開発モニターのご経験のある方は歓迎です。

Job Information

Recruiter
JAC Recruitment Co., Ltd.

Hiring Company
非公開

Job ID
1488689

Industry
Contract Research Organization

Job Type
Permanent Full-time

Location
Tokyo - 23 Wards

Salary
5.5 million yen ~ 14 million yen

Work Hours
09:00 〜 17:30

Holidays
【有給休暇】初年度 12日 1か月目から 【休日】完全週休二日制 土 日 祝日 GW 夏季休暇 年末年始 会社創立記念日（6/…

Refreshed
October 10th, 2024 02:00

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 NJB1052855】
「日本主導のグローバル試験」において、プロジェクトの進捗、品質、予算等を総合的に管理するグローバルプロジェクトマネージャとして、日本経つグローバルプロジェクト、アジアプロジェクトをマネージメントしていただきます。
■職務詳細
・スポンサーとの調整
・臨床開発プロジェクトのタイムラインの管理
・プロジェクトに関する提案、契約締結
・社内（日本を含むアジア、ヨーロッパ、アメリカ）の調整、進捗管理
・プロジェクトの予算管理、調整
・リスクマネジメントプランの作成

■メンバー構成：日本、ドイツ、韓国、台湾出身の方が在籍しております。

The Project Manager manages the project team in the delivery of quality clinical trial management servicesto achieve the successful overall project completion. The PM will be expected to maintain an in depthunderstanding of customer needs within the project group to focus on achieving the project's goals and tohave knowledge in the assigned therapeutic area. According to his/her knowledge and skills the PM maytake over any other tasks duties roles and job responsibilities at the discretion of the company andrelevant managers.The PM will provide a customer focused leadership role and may be assigned to manage multiple Phase Ithrough Phase IV clinical research trials across all functional areas. The role holder will assure theunderstanding and integration of all functions roles and responsibilities within the clinical project team andwill effectively coordinate and manage cross functional teams that deliver clinical projects （ClinicalMonitoring Data Management Biostatistics Medical writing Pharmacovigilance Regulatory） includingvendors if applicable.The PM will monitor adherence to project contract and budget. In addition the PM is expected to assist withproposal generation within a multifunctional matrix setting. The PM is expected to assist in businessdevelopment activities to achieve high level of customer satisfaction and therefore ensuring repeat businessopportunities.

Required Skills

臨床開発におけるグローバルプロジェクトマネジメント経験者
※アメリカ、カナダ、ヨーロッパ（ドイツ/フランス/イタリア/イギリス/オランダ）、韓国、台湾、中国出身の多国籍なメンバーが活躍しております。

■NecessaryEducation
University/college degree in a life science field （Master's or other advanced degree is preferred） orequivalent experience or equivalent education.At least 5 years of clinical research experience in a CRO biopharmaceutical company or relevant clinicalenvironment and a minimum of 2 years in team leader function （e.g. Monitor Lead （ML）） or equivalentstudy coordination or management experience. Experience in study management in Japan and/or an Asiancountry （e.g. Taiwan Korea） is required. Clinical operations management experience （e.g. ML） is required.In depth knowledge of the clinical trial process and CRO industry is mandatory for this function.Outstanding written and oral English and Japanese skills are required to qualify for this role. An additionallanguage is a plus.
■AdditionalQualifications
Knowledge and experience of project management tools; good knowledge of drug development process;drive for results; focus on customers and projects; global orientation /interests; proven expertise inmanagement/ financial control fundamentals; negotiation skills and awareness of regulatory requirementsand legal and contract issues; talent at presentations; extensive acumen in problem solving; experience atteam/staff mentoring; commitment to quality; strong interpersonal skills; ability to build a team; ability todirect and motivate staff; good internal

Company Description

ご紹介時にご案内いたします