

Study Manager

アストラゼネカ株式会社での募集です。 臨床開発リーダー・臨床開発プロジェクトマ...

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
アストラゼネカ株式会社

求人ID
1610547

業種
医薬品

会社の種類
外資系企業

雇用形態
正社員

勤務地
東京都 23区

給与
700万円 ~ 1000万円

勤務時間
09:00 ~ 17:15

休日・休暇
【有給休暇】有給休暇は入社時から付与されます 入社7ヶ月目には最低10日以上 【休日】完全週休二日制 年末年始 【有給休暇】※...

更新日
2026年10月30日 20:00

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
ネイティブレベル

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 NJB2341402】
'Responsible for leading and managing clinical delivery team of contributing experts to operationally deliver defined clinical studies or clinical project work package / activities to time cost and appropriate quality in Japan R D.Clinical Operations is the organisational home for the operational delivery and programme and project management of clinical projects drug and non drug.

スキル・資格

【経験 / Experience】
< 必須 / Mandatory >
・At least 5 years' experience from within the pharmaceutical industry or similar organization or academic experience.
・Extensive knowledge of clinical operations project management tools and processes
・Understanding of the skills and knowledge required for the successful delivery of a clinical study e.g. ICH GCP/J GCP local regulations study management
・Good experience of clinical development / drug development process in various phases of development and therapy areas・

< 歓迎 / Nice to have >
・Working experiences in Clinical Development/ Quality Assurance especially Oncology experience
・Knowledge of process improvement methodology such as Lean Sigma/Change Management

【資格 / License】
< 必須 / Mandatory >
・Bachelor's Degree in Science or related discipline

< 歓迎 / Nice to have >
・Medical or biological sciences or discipline associated with clinical research.・
・Advanced degree is preferred.・

【能力 / Skill set】
< 必須 / Mandatory >
・Excellent Communication and relationship building skills
・Proven ability to prioritize and manage multiple tasks with conflicting deadlines
・Excellent interpersonal skills and proven ability to facilitate team building and team work
・Proactively identifies risks and issues and possible solutions
・Demonstrates ability to prepare and deliver study related training materials
・Demonstrates ability to plan coordinate and facilitate internal and external meetings
・Demonstrates professionalism and mutual respect
・Displays excellent organization and time management skills excellent attention to detail and ability to multi task in a high volume environment with shifting priorities
・Proven learning agility
・Basic knowledge and experience of quality management

< 歓迎 / Nice to have >
・Project management certification・
・Ability to interact widely and effectively within the company at all levels. Good networking skills
・In depth knowledge of clinical and drug development process
・Fiscal and financial awareness

【語学 / Language】
< 必須 / Mandatory >
・英語 English：Business English（Achieve common understanding at the context level）
・日本語 : Native

会社説明

医療用医薬品の開発、製造及び販売