CareerCross by JAC Recruitment

【1000～1500万円】【R D】Associate Director Study Alliance Management …

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
1524906

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
1000万円 ~ 1500万円

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 【有給休暇】 初年度 4～16 日 （ 1 か月目～ ） 入社月により付与日数が…

更新日
2025年03月06日 16:24

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 NJB2283616】
■職務内容 / Job Description
Associate Director Study Alliance Management （ADSAM） is accountable/responsible for oversight of related full outsourcing studies to achieve milestones as scheduled with maintenance of quality and mediator among Japan CRO study team Global CRO representative Japan Associate Director Clinical Development （J ADCD） and AZ Global clinical team.

The ADSAM will contribute to optimize an alliance on full outsourcing studies from the planning to the regulatory inspections （e.g. create/improve oversight platform the agreement and Japan specific requirements through influencing global alliance team members and optimize upcoming study operation planning/execution by J ADCD）.

The ADSAM will take capability and skill development for Japan Study Manager （J SM） and contribute to improve related activities in the Japan Development Operations （J DO）.

スキル・資格

■応募資格（経験、資格等）/ Qualification （Experience Skill etc.）
【経験 / Experience】
＜必須 / Mandatory＞
At least 5 years' experience in pharmaceutical industries or multinational healthcare organization.
Extensive knowledge of clinical operations project management tools and processes
Proven experience of clinical development / drug development process in various phases of development and therapy areas including health authority GCP inspection.
Proven ability to learn by working in multiple phases TAs and/or different development situations.
Experience from leading clinical projects and deliverables or similar expertise from other areas of drug development （such as pharmaceutical development）.

＜歓迎 / Nice to have＞
Working with external bodies such as co development companies and key opinion leaders as a leading person.
Leadership of significant cross functional change programmes/initiatives

【資格 / License】
＜必須 / Mandatory＞
Bachelor's Degree

＜歓迎 / Nice to have＞
Medical or biological science or discipline associated with clinical research

【能力 / Skill set】
＜必須 / Mandatory＞
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
Excellent communication relationship building and negotiation skills
Proactively identifies risks and issues and possible solutions
Basic knowledge and experience of quality management

【語学 / Language】
＜必須 / Mandatory＞
日本語 Japanese：ネイティブ
英語 English：英語 English： Business English （Achieve common understanding at the context level with customers）

会社説明

医療用医薬品の開発、製造及び販売