【1200 ~ 1500万円】Director Reg Affairs

臨床開発リーダー・臨床開発プロジェクトマネージャ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
1590158

業種
CRO

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
1200万円 ~ 1500万円

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05月28日 06:00

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

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 NJB2306555】
Job Description
RADDS Japan was established to support EBP initiative for business growth in RD S. This position requires the successful candidate to support the activities in RADDS Japan which includes contributing to the scoping integrated proposal compilation and delivery oversight of multi disciplinary biopharmaceutical development advisory/clinical trial enablement projects.

【業務内容】
・Manage staff in accordance with organization’s policies and applicable regulations; Responsibilities include planning assigning and directing work; appraising performance and guiding professional development; rewarding and disciplining employees; addressing employee relations issues and resolving problems; Approve actions on human resources matters including salary administration.

・Effectively manages the team's resources and delegates tasks commensurate with skill level; Evaluates workload quality and metrics through regular review and reporting of findings;
Effectively collaborate with operational colleagues to manage project and team related challenges;
Contributes to discussions on implementation of business strategy on a regional basis and will implement regional and global specific objectives as appropriate.

・Ensures staff have a consistent understanding and positive impression of strategy for regional and global objectives;
Leads strategic initiatives and develops implementation plans;

・Will have full financial responsibility and accountability for one or more Regulatory Affairs sites; monitors growth and performance of the sites;
Undertakes risk analysis and manages the outcome as appropriate;

・May act as a Project Manager for a large and complex stand alone project or programme involving several regulatory or technical deliverables and/or region and/or operations;

・May provide strategic regulatory and/or technical consultancy on a variety of projects;

・Competently manages meetings/expectations with Regulatory Agencies and/or groups within IQVIA;

・May take leadership role in bid defense strategy and planning;

・May lead/chair a session on Regulatory Affairs or related topics at a conference; deliver effective presentations to a broad audience;

スキル・資格

（求めるスキル、経験）

CROでのRA経験必須（ビジネスを理解している）

・At least 8 years regulatory experience including 6 years management experience.

・Requires broad management knowledge to lead teams and well as the ability to influences others outside of own job area regarding policies procedures and goals.

・Possess demonstrated working knowledge of corresponding professional grade level responsibilities skills and abilities as required for guidance of staff

Line management experience required with demonstrated success in development engagement and performance of senior staff.

・Advanced negotiating and influencing skills and the ability to identify and resolve issues using flexible adaptable approach. Remains calm assertive and diplomatic in challenging interactions with customers and staff

・Possesses an acute awareness of issues outwith the business function. Exudes confidence and authority within remit and delivers a positive example to teams

・Communicates an inspiring vision of the Regulatory Strategic Plan and helps others understand and value their role in it.

・Ability to work effectively with senior management remaining motivated and enthusiastic in times of change and other pressure situations

会社説明

ご紹介時にご案内いたします