【1100〜1500万円】【R D】Associate Director ・ Cell Therapy Territory O…

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
1573304

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
1100万円 ~ 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
2026年05月02日 22:00

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 NJB2354086】
The Associate Director ・ Cell Therapy Territory Operations Manager （AD TOM） is a regional expert supporting Patient Operations and project teams in cell therapy. As a senior technical leader the role sets and implements compliant policies and standards for treatment centers partners with Regional Operations Directors （RODs） on site strategy leads research and internal reporting and maintains up to date best practices.
The role expands AZ’s cell therapy footprint in oncology by building patient centered service lines and leading complex

engagements with site leadership and frontline providers. It delivers operational and clinical guidance for trials and commercial products leads site activation and provides ongoing operational quality and clinical oversight post activation. The AD TOM is a field regional role reporting to a Regional Operations Director （ROD）.

You Will:
・・ Lead and support strategic initiatives across clinical and commercial cell therapy from development to post market.
・ Be a key leader within Internal AZ teams （Program Management QA Manufacturing Supply Chain Medical Affairs Market Access Compliance Sales/Marketing）.
・ Establish service lines with external partners （PIs Physicians Nursing Apheresis Cell Therapy Lab other HCPs/leadership）.
・ Serve as SME driving decisions and projects at multi country/local levels; influence strategy governance and cross functional execution.
・ Acts as the technical expert and authoritative source of knowledge in cell therapy workflows and brings expertise to site interactions.
・ Build and maintain relationships with stakeholders at assigned Sites.
・ Operate within regulatory/accreditation standards （e.g. FACT） ensuring compliance audit readiness and continuous improvement.
・ Guide clinical workflows; ensure standards; advance apheresis excellence; optimize product handling; inform health outcomes.
・ Act as primary operations readiness liaison; collaborate with Supply Chain to align logistics and site needs.
・ Provide exemplary customer service while building business relations with clinical/operational leaders.
・ Lead cross functional meetings to develop product/patient workflows for trials and future commercial products.
・ Maintain strong collaboration with Program Management SMM Supply Chain Manufacturing Quality Medical Commercial Cell Therapy Hub to manage customer centers and clinical care for the portfolio.
・ Oversee operational/clinical aspects of raw material collection product receipt/storage/distribution/infusion and COI/COC.
・ Design and deliver training to PIs MD Nursing Staff Apheresis and Cell Therapy Laboratory and Coordinator staff
・ Lead Site activation using risk based tools for certification training and monitoring.
・ Train Site HCPs on product needs; ensure COI/COC access for trained individuals with Cell Therapy Hub.
・ Oversee Site logistics and manage regional account changes with Cell Therapy Hub.
・ Respond to SOP deviations with expertise and collegiality.
・ Support/own quality records （Deviations CAPA Change Control） with Cell Therapy Hub and QA.
・ Travel up to 50%.

スキル・資格

You Have:
・ Bachelor’s degree （Nursing Physician Assistant） with thorough knowledge/hands on experience in Cell Therapy （preferred: MD or similar）.
・ 8+ years in cell therapy with progressive leadership.
・ Expertise in service lines （infrastructure resourcing costs quality metrics）.
・ Proven delivery to timelines cost and quality; collaboration with internal/external providers.
・ Excellent knowledge of cell therapy standards/regulations.
・ Experience selecting/overseeing external providers and developing contracts.

Nice to Have:
・ Advanced scientific degree; broad understanding across patient operations manufacturing interfaces quality regulatory market access; project management experience.
・ Program/Quality qualifications （e.g. PMP CPHQ）.
・ Experience across academic/Sponsor settings and countries.
・ Early phase delivery across product lifecycle and multiple therapeutic areas.
・ Salesforce or similar CRM experience.

会社説明

医療用医薬品の開発、製造及び販売