

【1200 ~ 1500万円】【Medical】Medical Evidence Lead（Med.EL），メディカル本部 エビデ…

アストラゼネカ株式会社での募集です。 メディカルアフェアーズのご経験のある方は…

Job Information

Recruiter
JAC Recruitment Co., Ltd.

Hiring Company
アストラゼネカ株式会社

Job ID
1622056

Industry
Pharmaceutical

Company Type
International Company

Job Type
Permanent Full-time

Location
Tokyo - 23 Wards

Salary
12 million yen ~ 15 million yen

Work Hours
09:00 ~ 17:15

Holidays
【有給休暇】初年度 10日 1か月目から 【休日】完全週休二日制 年末年始 【有給休暇】※入社月により付与日数が異なります。詳…

Refreshed
November 12th, 2026 14:56

General Requirements

Career Level
Mid Career

Minimum English Level
Business Level

Minimum Japanese Level
Native Level

Minimum Education Level
Post Grad Degree (PHD/MBA etc)

Visa Status
Permission to work in Japan required

Job Description

【求人No NJB2423114】
■職務内容 / Job Description

Medical Evidence Lead（Med. EL）is a project leadership role that generates valuable real world evidence and accelerates the use of real world data for medical evidence generation in line with brand strategy. Med. EL applies clinical research

methods methodological principles and a strong understanding of the strategic business context in the relevant therapeutic area to enable high quality evidence generation across study planning execution and reporting.

Depending on study needs and organizational assignment Med. EL may serve in either one or both of the following leadership capacities:

・ Study Lead（Operational Lead） driving cross functional study planning project management

operational execution external service provider selection and oversight and delivery throughout the

study lifecycle

・ Methodological Lead driving methodological reliability and scientific robustness of study design

protocol development analytical requirements interpretation and reporting

Med. EL works cross functionally with internal and external experts including Data Scientists Statisticians Medical Affairs colleagues External Service Providers（ESPs） and/or research collaborators as appropriate to ensure that study activities and outputs are aligned with study objectives timelines quality standards budget relevant regulations and internal procedures.

Required Skills

■応募資格（経験、資格等）/ Qualification（Experience Skill etc.）
【経験 / Experience】
＜必須 / Mandatory＞
Clinical research experience（ 5 years） meeting any of the following:
Leading protocol development data management analysis result interpretation reporting and publication
Working in R D and/or Medical function in pharmaceutical industry
Working in epidemiology public health research clinical research or related evidence generation fields

＜歓迎 / Nice to have＞
・ Working experience in CVRM or R I area
・ Clinical project leader/study leader experience
・ Research using statistics and epidemiology
・ Planning and execution of database research using personal health data
・ Experience in a data science role
・ Oversight of CROs/ESPs for clinical studies

【資格 / License】
＜必須 / Mandatory＞
・Master's degree in a scientific discipline・

＜歓迎 / Nice to have＞
・PhD in a scientific discipline or MD

【能力 / Skill set】
＜必須 / Mandatory＞
・ Clinical research methods study design and applied methodological principles
・ Project Management
・ Leadership and Communication
・ Medical writing
・ Cross functional stakeholder management

＜歓迎 / Nice to have＞
・ Scientific Publication
・ Ability to review analysis dataset and tables figures and listings specifications

【語学 / Language】
＜必須 / Mandatory＞
日本語 Japanese：Native or equivalent（日本語検定1級レベル）
英語 English：Practical English communication skill / TOEIC 800

＜歓迎 / Nice to have＞
・Working experience outside Japan and/or in global organizations/teams

Company Description

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