＜研究開発・メディカルアフェアーズ統括本部＞Project Statistician/ Principal Statistic…

日本イーライリリー株式会社での募集です。 統計解析のご経験のある方は歓迎です。

Job Information

Recruiter
JAC Recruitment Co., Ltd.

Hiring Company
日本イーライリリー株式会社

Job ID
1585292

Industry
Pharmaceutical

Company Type
International Company

Job Type
Permanent Full-time

Location
Tokyo - 23 Wards

Salary
6 million yen ~ Negotiable, based on experience

Work Hours
08:45 〜 17:30

Holidays
【有給休暇】有給休暇は入社後2ヶ月目から付与されます （ 年途中で入社した社員に対する年次有給休暇は、次の入社月の区分に従い入…

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April 2nd, 2026 15:12

General Requirements

Career Level
Mid Career

Minimum English Level
Business Level

Minimum Japanese Level
Native

Minimum Education Level
Bachelor's Degree

Visa Status
Permission to work in Japan required

Job Description

【求人No NJB2348617】
職務内容
The Project statistician develops or assists in the development of protocol designs clinical plans and data analysis plans in collaboration with physicians veterinarians and/or medical colleagues. The Project Statistician is responsible for working with research associates and scientists to establish a reporting database and for analyzing data for these types of studies.

[Clinical Project Statistician]
Provide strong statistical leadership in the process of drug development.

[Real World Analytics Project Statistician]
Provide input and statistical leadership for post launch activities including Health technology assessment （HTA） real world evidence （RWE） research post market safety studies medical affairs publications to maximize the value of our products in collaboration with cross functional and cross regional partners.

主な職責/Primary responsibilities
Statistical Trial Design and Analysis

[Clinical Project Statistician]
・Operate in collaboration with study personnel to provide input on study protocol design studies and write protocols for the conduct of each study.
・Assist in or be accountable for selecting statistical methods for data analysis authoring the corresponding sections of the protocol and conducting the actual analysis once a reporting database is created.
・Collaborate with data management in the planning and implementation of data quality assurance plans.
・Maintain currency with respect to statistical methodology to maintain proficiency in applying new and varied methods and to be competent in justifying methods selected.
・Participate in peer review work products from other statistical colleagues.

[Real World Analytics Project Statistician]
・Lead statistical activities with the development of protocols analysis plans and analysis execution and interpretation for RWE research and post market safety studies.
・Provide technical statistical leadership for network meta analysis/indirect comparisons economic modelling and dossier development for Health technology assessment （HTA）.
・Ensures high quality statistical support for projects through the oversight of external suppliers and provides input into outsourcing strategy and processes.
・Influences cross functional team members regarding appropriate research method.

Communication of Results and Inferences
・Collaborate with team members to write reports and communicate results.
・Assist with or be responsible for communicating study results via regulatory submissions
manuscripts or oral presentations in group settings as well as for communicating one on one with key customers and presenting at scientific meetings.
・Respond to regulatory queries and to interact with regulators.

Therapeutic Area Knowledge
・Understand disease states in order to enhance the level of customer focus and collaboration and be seen as a strong scientific contributor.

Regulatory Compliance
・Perform work in full compliance with assigned curriculum（s） and will be responsible for following applicable Corporate Medical local and departmental policies procedures processes and training

Required Skills

■Basic Requirements:
・M.S. or Ph.D. in statistics or biostatistics

■Other Information/Additional Preferences:
・Rich experiences in healthcare field as a statistician with science background such as mathematics or epidemiology.
・Statistical and methodological knowledge in clinical development epidemiology or related field.
・Regulatory knowledge of clinical trial methodology and statistics.
・Ability to build relationships with individuals and teams.
・Good communication and presentation skills in both English and Japanese

Company Description

医療用医薬品の輸入・製造・販売