CareerCross

＜研究開発・メディカルアフェアーズ統括本部＞信頼性保証本部　品質情報マネジメント/担当・担当課長/P1 P2/神戸本社

日本イーライリリー株式会社での募集です。 メディカルGQP・GMP・品質保証・...

Job Information

Recruiter
JAC Recruitment Co., Ltd.

Hiring Company
日本イーライリリー株式会社

Job ID
1475045

Industry
Pharmaceutical

Company Type
International Company

Job Type
Permanent Full-time

Location
Hyogo Prefecture

Salary
5.5 million yen ~ Negotiable, based on experience

Work Hours
08:45 〜 17:30

Holidays
【有給休暇】初年度 10日 2か月目から付与及び使用可能 【休日】完全週休二日制 祝日 夏季休暇 年末年始 【有給休暇】有休休...

Refreshed
June 21st, 2024 14:00

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 NJB2222086】
Local device stewardship is expected to be the master of devices and accountable for the related information/communication demonstrating following duties locally to meet business needs in Asia Pacific region.

・Understand technical principle and mechanism of devices behind scientific principles.

・Transform technical information into non technical communication in the appropriate form （e.g. verbal written sentence picture drawing） that fit to needs.
・Provide technical support for going business timely.
・Ensure appropriate consistency and accuracy between technical information and non technical communication across materials media channel and parties.
・Maintain appropriate consistency and accuracy with global information/direction owned by global product stewardship and/or process/engineering expert across devices across materials/reports.
・Obtain necessary technical information timely from global product stewardship and/or related process/engineering expert.
・Develop local training material for device and provide training including call center and third party being qualified for trainer training for the device trainings.
・Review materials/reports for device related contents.
・Educate individuals in Quality Assurance organization to help them better perform in responsible area.
・Be familiar with global and local key regulatory requirements standards those relate to devices.
・Collaborate for device related activities with associated functions understanding the requirement and the system.
・Demonstrate to be trusted and appreciated from customers.
・Evaluate the completed complaint investigation and identify an appropriate customer response.

Required Skills

下記は担当・担当課長の必須条件です。＜Education/Skill Requirements＞・+ Two years of Quality Assurance Engineer experience in Pharmaceutical Medical device Industry （e.g. injector Gastroscope （Becton Nipro Termo Canon etc） ・ Strong verbal and written communication skills in Japanese（Native） and English （Preferable: TOEIC score = 750 or above） ・A bachelor's degree in Science related field （e.g. pharmacy chemistry biology or engineering）＜Minimum Requirements＞・Have basic engineering and process knowledge or interest ・Strong Lilly product knowledge relating to how products work medicinally and functionally ・Solid collaboration and interpersonal skills ・Self management/motivated with an ability to work independently within a structured process ・Excellent teamwork with an ability to multi task ・Strong critical thinking/problem solving skills with an ability to apply rationale ・Ability to mentor/train others share learning

Company Description

医療用医薬品の輸入・製造・販売