Senior/Advanced Project QA Professional

> **ノボ ノルディスク ファーマ株式会社での募集です。 メディカルGQP・GMP・...**

Job Information

Recruiter
JAC Recruitment Co., Ltd.

Hiring Company
ノボ ノルディスク ファーマ株式会社

Job ID
1578853

Industry
Pharmaceutical

Company Type
International Company

Job Type
Permanent Full-time

Location
Fukushima Prefecture

Salary
7 million yen ~ 15 million yen

Work Hours
08:50 ~ 17:10

Holidays
【有給休暇】有給休暇は入社時から付与されます 初年度 12日 1ヶ月目に付与（最高20日付与）【休日】完全週休二日制 年末...

Refreshed
June 14th, 2026 02:00

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 NJB2361383】
As a QA Professional you will play a key role in ensuring compliance with GxP regulations and maintaining the highest quality standards. Your responsibilities will include:

・Providing QA oversight and support for projects in Koriyama factory in Japan to ensure timely delivery with the right quality.

・Participating in project and validation activities to resolve quality related issues.
・Performing quality spot checks and approving validation documentation to ensure the highest quality level.
・Reviewing and approving complex change control requests SOPs trend reports and validation related documents.
・Preparing annual product reviews and facilitating validation and other complex internal trainings.
・Acting as a coach during audits and inspections and ensuring compliance with regulations and corporate/local SOPs.
・Providing proactive guidance to departments on quality related activities and ensuring that current Novo Nordisk procedures and health authority requirements are reflected in quality decisions.

Required Skills

【Your skills qualifications】
We are looking for a dedicated and experienced QA professional who can bring the following qualifications to the role:

・Minimum 5 years' QA experience or minimum 7 years' experience in QC/production or related roles （healthcare/pharmaceutical industry preferred） with a strong quality mindset and solid understanding of QA responsibilities.
・Expert knowledge of quality management systems and cGMP production processes with deep understanding of validation in pharmaceutical manufacturing preferred.
・Ability to make autonomous high・level decisions on complex quality issues and to perform effectively with minimal supervision.
・Strong cross・functional leadership skills to drive process and team activities and to support others' ideas and actions.
・Native level Japanese and business level English communication skills both written and oral.

Company Description

医療用医薬品、医療機器の開発、輸入・製造、販売