



## 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・...

### 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

ノボ ノルディスク ファーマ株式会社

#### 求人ID

1578853

#### 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

福島県

#### 給与

700万円 ~ 1500万円

#### 勤務時間

08:50 ~ 17:10

#### 休日・休暇

【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...

#### 更新日

2026年04月30日 01:00

### 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 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.
- 

## スキル・資格

【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.
- 

## 会社説明

医療用医薬品、医療機器の開発、輸入・製造、販売