



【1100～万円】西神エンジニアリングSeishin Engineering/課長 Senior Manager/M1

日本イーライリリー株式会社での募集です。管理職（工場・技術マネジメント）のご...

#### 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

日本イーライリリー株式会社

#### 求人ID

1582823

#### 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

兵庫県

#### 給与

1100万円～経験考慮の上、応相談

#### 勤務時間

08:45～17:30

#### 休日・休暇

【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 完全週休2日制 (土...

#### 更新日

2026年03月19日 15:17

#### 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 NJB2366933】

【職務概要】

Engineering Manager is responsible for the management of Engineering・Group.

In order to keep the continual supply of high quality finished products to the Japanese market which meet the respective regulatory requirement this position is accountable for supervising the control over maintenance validation and・

environmental control utilizing sufficient knowledge and experience for GMP issues. They're also responsible for talent development.

【職務内容】

Responsible and owns of engineering documents training and activities.

Plans for technical GMP improvement to be conducted at Engineering and control and check the implementation.

In case of technical issues such as deviations and change proposals in packaging area technically control the process to assure the quality.

Develop local processes and procedures in accordance with Engineering management system.

Provides technical oversight and stewardship for the production operations

Ensures staff are qualified and that the required initial and continuing training of technical support personnel is carried out and adapted according to need to ensure appropriate operations support.

Ensures the process and product are well characterized and that the process description and control strategy including the measures acceptable ranges and specifications (and the justification of these measures acceptable ranges and specifications) have been documented in a process flow document (PFD) or its equivalent.

Ensures that the processes facilities equipment or systems for which they are responsible have been validated or qualified and

Ensures that the processes facilities equipment or systems for which they are responsible are compliant capable in control and maintained in a validated or qualified state.

---

## スキル・資格

### 応募資格\*\*

Bachelor grade or equivalent in engineering scientific background. ·

- Minimum 5 years experience in manufacturing environment ·
- Knowledge of GMP Pharmaceutical Science Pharmaceutical Law Finance · Strong Communication Skills ·
- Leadership capability in management of group members and execution of improvement work. ·
- Ability to influence at all levels of organization ·
- Ability to maintain rapport with internal and external partners ·
- Cross functional communication skill

---

## 会社説明

医療用医薬品の輸入・製造・販売