



# PR/116978 | QA Assistant Manager

# 募集職種

# 人材紹介会社

ジェイ エイ シー リクルートメント タイランド

#### 求人ID

1537146

#### 業種

その他 (メーカー)

### 雇用形態

正社員

#### 勤務地

タイ

#### 給与

経験考慮の上、応相談

#### 更新日

2025年04月30日 16:21

# 応募必要条件

# 職務経験

3年以上

# 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

ビジネス会話レベル

### 最終学歴

短大卒: 準学士号

# 現在のビザ

日本での就労許可は必要ありません

# 募集要項

The company specializes in delivering Electronics Manufacturing Services (EMS) and regional manufacturing solutions for global customers. Renowned for its commitment to continuous improvement, the company has earned numerous awards and recognitions. The ideal candidate for this position should possess strong leadership skills and expertise in quality systems, particularly within the electronics, medical, or food industries. Demonstrate a dedication to personal and professional growth, evolving alongside the company while adapting effectively to change.

Position: QA Assistant Manager

Subordinates: 34 persons (including QA operator)

Salary: 80,000 – 100,000 THB / month

<u>Location:</u> Laem Chabang Industrial Estate\_

### Responsibilities:

- · Coordinate with plant organizations to review policies, procedures, regulations, and work instructions.
- · Review and monitor new or updated QMS and relevant regulation to ensure the compliance of company system
- Manage audit programs, management reviews, and reporting.
- Oversee complaints, CAPA system, and risk management (ISO14971).
- Maintain and supervise for Complaint file, Corrective action & Preventive action system and Risk Management (ISO14971)
- Maintain audit programs and Management review and provide reporting results to management
- Facilitate communication between functions and multi-facility groups.

# **Qualifications:**

- Bachelor's degree or higher in Engineering field or equivalent industry experience .
- A minimum of 10 years of experience in the Quality System Management
- Experienced in the position of QMR is advantage.
- Knowledge of FDA Quality System Requirements 21CFR820, 21CRF PART11, MDD/MDR, ISO 13485, ISO 14971, DMR, DHR and Process Validation IQOQPQ, GMP and GDP is advantage.
- Leadership skills including ability to maintain confidentiality.
- Good communication, teamwork, and organizational skills to carry out projects to meet timelines.
- · Good command in English.

会社説明