Manager GDP Export Quality

武田薬品工業株式会社での募集です。 物流・受発注納品手配のご経験のある方は歓迎...

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
武田薬品工業株式会社

求人ID
1590586

業種
医薬品

雇用形態
正社員

勤務地
大阪府

給与
600万円 ~ 1100万円

勤務時間
08:00 ~ 16:45

休日・休暇
【有給休暇】初年度 12日 1か月目から 完全週休二日制（土・日）、祝日、メーデー、年末年始、他 特別有給休暇、リフレッシュ休...

更新日
2026年06月11日 05:00

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
流暢

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 NJB2375005】
【OBJECTIVES/PURPOSE:】

Ensure compliant GDP export execution・by providing operational Quality oversight and expert guidance across export/transportation activities maintaining inspection readiness and robust documentation.

Lead risk and quality event management・by identifying and mitigating distribution risks translating QMS requirements into practice and driving deviations/CAPA and change control for export lanes providers and processes.

Oversee external partners（especially central contracted distributors）・through qualification and performance monitoring effective Quality Agreements and strong cross functional collaboration across Supply Chain Site Quality LOC Quality and project teams.

As deputy RP responsible Eligible to perform all the duties and have all rights applicable to the Deputy Responsible Person in Singapore as per HSA requirements

【ACCOUNTABILITIES】

Provide operational GDP Quality oversight for export execution・across shipment preparation handover transportation and delivery ensuring activities meet applicable GDP regulations and Takeda Quality standards.

Serve as the Quality SME for export and transportation・（including cold chain where applicable）advising on packaging/ship to configurations monitoring strategy handling instructions security controls and lane suitability.

Lead risk identification assessment and mitigation・for export distribution（lanes carriers forwarders 3PLs distributors）ensuring risks are documented owned controlled and periodically reviewed in line with governance expectations.

Oversee deviation/event management for export distribution・（e.g. temperature excursions delays damages misroutes seal breaches）ensuring timely triage investigation support documented Quality impact assessment and effective CAPA implementation.

Drive change control for export GDP scope ensuring changes to lanes logistics partners distributors packaging solutions and monitoring processes are assessed and implemented with appropriate Quality review evidence and approvals.

Deliver GDP oversight of contracted distributors・（central Takeda contracts）including distributor qualification/onboarding Quality Agreement content and maintenance periodic performance review escalation management and ensuring continued compliance with Takeda expectations.

Ensure compliance with Good Manufacturing Practice and Good Distribution Practice Standards in Singapore

The company has given authority to the Responsible Person to perform the required activities as per local regulations which include:

ensuring that an effective quality system is implemented and maintained that meets GDP standard;

focusing on the management of authorized activities and the accuracy and quality of records;

ensuring that initial and continuous training prorammes are implemented and maintained;

coordinating and promptly performing any recall operations for therapeutic products;

ensuring that relevant customer complaints are dealt with effectively;

ensuring that suppliers and customers are legally approved or authorized to enable lawful supply of therapeutic products;

approving any subcontracted activities which may impact on GDP;

ensuring that self inspections are performed at appropriate regular intervals following a prearranged programme and necessary corrective measures are put in place;

keeping appropriate records of any delegated duties;

deciding on the final disposition of returned rejected recalled or counterfeit products;

approving any returns to saleable stock;

ensuring that any additional requirements imposed on certain products by national legislation are adhered to e.g. controlled drugs.

【CORE ELEMENTS RELATED TO THIS ROLE】

Maintain strong cross functional collaboration・with Global Supply Chain Trade Compliance Site Quality Organizations GDC/RDC related LOC Quality and Takeda project teams to resolve issues support initiatives and ensure aligned decision making.

Experience overseeing・global export lanes cold chain distribution and complex third party networks.

Experience with distributor qualification/oversight programs and supplier management.

Familiarity with global GDP frameworks（e.g. EU GDP guidelines）and practical application across regions.

Ability to interpret data trends（excursions lane performance partner KPIs）to drive risk based decisions

スキル・資格

・Technical/Functional（Line） Expertise
Demonstrated experience in・GDP Quality・within pharmaceutical distribution logistics or supply chain（typically 5+ years depending on internal leveling）.
Strong working knowledge of GDP expectations for・transportation ・outsourced distribution and・quality agreements.
Experience with deviation management investigations CAPA and audit readiness in a regulated environment.
Understanding of Commercial Quality requirements including regional and in country distribution controlled substance compliance supply chain quality and local quality surveillance（returns recalls complaints）

・Leadership
Ability to collaborate and partner well regionally cross functionally with Takeda stakeholder groups and RDCs
Strong communication skills with ability to influence cross functional stakeholders and external partners.
Ability to drive change by influence

・Decision making and Autonomy
Quality approval/endorsement for GDP controls impacting export lanes and distribution models（per Takeda governance）.
Role specific accountability for Quality oversight of selected distributors and logistics provider

・Interaction
Internal:・Supply Chain Governance Global/Regional Quality Logistics/Transportation teams Trade Compliance Planning Customer Service Site QA/QP RP
External:・3PLs freight forwarders carriers packaging and monitoring suppliers contracted distributors.

・Innovation
Understanding of quality and regulatory requirements and trends related to warehousing and distribution across the global regions.

・EDUCATION BEHAVIOURAL COMPETENCIES AND SKILLS:・
Scientific Degree（BSc MSc）
Minimum 10 years experience in in the pharmaceutical industry including Quality Assurance combined with a good knowledge of regulations pertaining to pharmaceutical manufacturing and distribution in the region（e.g. familiar with cGMP/GDP ISO and ICH requirements）・
Strong attention to details ability to review managing documentation（including transport temperature records shipping documents and related GDP documents/ records etc.）
Prior experience managing external GMP/GDP suppliers
Fluent in written and spoken English

・Core Competencies / Skills
Risk based decision making with strong Quality mindset
Pragmatic operational execution and ownership
Partner management and governance
Clear documentation and inspection readiness discipline
Continuous improvement and problem solving

・Deliver on Takeda Leadership behaviours
Think Strategically
Inspire others
Deliver・ priorities
Elevate Capabilities

会社説明

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