



PR/095792 | Regional Regulatory Affairs Specialist

Job Information

Recruiter

JAC Recruitment Singapore

Job ID

1575392

Industry

Healthcare, Nursing

Job Type

Permanent Full-time

Location

Singapore

Salary

Negotiable, based on experience

Refreshed

February 3rd, 2026 10:42

General Requirements

Minimum Experience Level

Over 1 year

Career Level

Mid Career

Minimum English Level

Fluent

Minimum Japanese Level

None

Minimum Education Level

Associate Degree/Diploma

Visa Status

No permission to work in Japan required

Job Description

Job Title: Regional RA Specialist

Location: Singapore

About Us:

We are partnered with a leading pharmaceutical company committed to innovative healthcare solutions across Asia. They are dedicated to enhancing patient well-being through rigorous regulatory compliance and strategic pharmaceutical development.

They are seeking an experienced Regulatory Specialist based in Singapore to lead and support regulatory activities across the Asia & Arab region (excluding China & Japan). This role provides strategic regulatory advice, manages complex regional submissions and lifecycle activities, supports interactions with health authorities, and strengthens regulatory capability across local affiliates and cross-functional teams.

Key Responsibilities:

- Develop and execute regional regulatory strategies for new product registrations, renewals and major post-approval changes; anticipate risks and recommend mitigation to optimize approval timelines.
- Lead coordination with local affiliates and Responsible Persons for submissions, product registrations, renewals and lifecycle management; review and validate ASEAN CTD / eCTD documentation.
- Support interactions with Health Authorities by preparing scientific justifications, briefing materials and high-quality responses to queries.
- Perform regulatory risk assessments for submissions and variations; track risk and propose actionable mitigation plans.

- Monitor regulatory intelligence across APAC/ASEAN and the Arab region; analyze impacts and share insights to inform business decisions.

Qualifications:

- 2–5 years' regulatory affairs experience in the pharmaceutical industry, with regional exposure in APAC preferred.
- Strong understanding of APAC/ASEAN/ICH regulatory frameworks and country-specific requirements.
- Proven ability to manage complex regulatory tasks independently and provide strategic regulatory input.
- Experience preparing or reviewing ASEAN CTD / eCTD modules and supporting Health Authority interactions.

Interested applicants, please click **APPLY NOW**.

Please note, only shortlisted candidates will be contacted.

JAC Recruitment Pte Ltd
EA License Number: 90C3026
EA Personnel: R2197665

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#countrysingapore

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Company Description