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Michael Page www.michaelpage.co.jp

Director, Regulatory Affairs (upto 15M) - Fully Remote

RA Director - (Up to 15M) Fully Remote

募集職種

人材紹介会社
マイケル・ペイジ・インターナショナル・ジャパン株式会社

求人ID
1546587

業種
医薬品

雇用形態
正社員

勤務地
東京都 23区

給与
1000万円 ~ 1500万円

更新日
2025年07月02日 09:42

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

- Join a global regulatory consultancy as their first permanent regulatory strategy lead in Japan. This senior role supports development-stage projects, PMDA interactions, and submission planning across multiple therapeutic areas.

Client Details

- Our client is a leading global provider of regulatory, clinical, and compliance services, supporting pharmaceutical and biotech companies worldwide. Known for its high-quality service and fully remote work culture, the organization offers strong career growth and meaningful global impact.

Description

- Lead Japan regulatory strategy for global drug development programs
- Manage PMDA consultations, CTNs, and registration submissions
- Act as Japan project lead and coordinate with global teams
- Align Japan regulatory plans with global development strategies
- Review English-language submissions for Japanese use
- Oversee local vendors and consultants to ensure delivery
- Stay current on regulatory changes and contribute to planning
- Represent the company in client meetings with strategic input

Job Offer

- Fully remote work
- English-using environment with global teams
- Competitive salary up to ¥15M total compensation
- High-impact role with international client exposure
- Opportunity to shape regulatory function in Japan

To apply online please click the 'Apply' button below. For a confidential discussion about this role please contact Sobi Tantisakchaichan on +81357337165.

スキル・資格

A successful RA Director should have:

- Regulatory or development experience focused on Japan
- Strong PMDA interaction experience (clinical & pre-market)
- Proven project leadership from early development to NDA
- Comfortable operating independently and cross-functionally
- Fluent in Japanese and English
- Consulting/CRO/global pharma background preferred
- Strategic, collaborative, and hands-on mindset

会社説明

Our client is a leading global provider of regulatory, clinical, and compliance services, supporting pharmaceutical and biotech companies worldwide. With a strong reputation for quality and innovation, they offer long-term career growth and a collaborative international culture.