



【800～2000万円】 Deputy Director Pharmacovigilance

安全性情報（臨床開発・製販後GVP）のご経験のある方は歓迎です。

## Job Information

### Recruiter

JAC Recruitment Co., Ltd.

### Hiring Company

非公開

### Job ID

1563534

### Industry

Pharmaceutical

### Company Type

International Company

### Job Type

Permanent Full-time

### Location

Tokyo - 23 Wards

### Salary

8 million yen ~ 20 million yen

### Work Hours

09:00 ~ 18:00

### Holidays

【有給休暇】有給休暇は試用期間満了後から付与されます 年間20日付与 【休日】完全週休二日制 土 日 祝日 年末年始 年末年始...

### Refreshed

February 5th, 2026 00:00

## General Requirements

### Career Level

Mid Career

### Minimum English Level

Fluent

### Minimum Japanese Level

Native

### Minimum Education Level

Bachelor's Degree

### Visa Status

Permission to work in Japan required

## Job Description

【求人No NJB2334734】

### ■POSITION SUMMARY

This position leads the pharmacovigilance (PV) function in Japan ensuring the effective delivery of PV consulting services to pharmaceutical companies. The role requires a balance of hands on project delivery team leadership and business development responsibilities. The individual will build and develop a local PV team while collaborating closely with global colleagues to align with the overall practice strategy.

The role covers a broad scope of client engagements from early development consulting to post marketing PV strategies. It involves frequent client interaction internal coordination and participation in external networking and industry discussions.

#### ■RESPONSIBILITIES

##### Delivery

- Accountable for successful delivery of all PV consulting projects in Japan.
- Serve as a project leader or subject matter expert (SME) in PV where necessary.
- Provide guidance and problem solving support to Japan based consultants facing project challenges.
- Collaborate with the global team on staffing for Japan based PV projects.

##### Business Development

- Identify and pursue new business opportunities based on client needs and market trends.
- Foster strong client relationships through regular communication and visits.
- Support proposal development and represent the Japan PV function at client meetings and industry events.

##### Team Development

- Collaborate with global and local leadership to define hiring needs and recruit local talent including independent consultants.
- Lead onboarding coaching and performance management of PV team members in Japan.
- Serve as the go to person and leader for the Japan PV team ensuring team cohesion and alignment with company goals.

##### Thought Leadership

- Contribute to the strategic direction of the global PV practice identifying opportunities for service expansion and innovation.
- Represent the company at industry forums conferences and networking events sharing insights and thought leadership.

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## Required Skills

#### ■MINIMUM QUALIFICATIONS

- Practical experience in one or more of the following domains: pharmacovigilance (preferred) risk management medical information regulatory operations or clinical operations.
- Proven leadership experience managing teams or cross functional groups.
- Demonstrated ability to manage client relationships in a consulting or service based environment.
- Experience working in or with global teams.
- Business level fluency in both English and Japanese.

#### ■PREFERRED QUALIFICATIONS

- Established professional network in the pharmaceutical or CRO industry.
- Excellent interpersonal skills and the ability to collaborate with international teams.
- Willingness and ability to travel domestically and internationally for client engagements.

#### 必須条件

- ・ 以下いずれかの分野での実務経験：ファーマコビランス（歓迎）、リスクマネジメント、メディカルインフォメーション、薬事オペレーション、臨床開発オペレーション。
- ・ チームまたはクロスファンクショングループのマネジメント経験。
- ・ コンサルティングまたはサービス提供型環境におけるクライアント対応経験。
- ・ グローバルチームでの業務経験。
- ・ ビジネスレベルの日本語および英語力。

#### 歓迎条件

- ・ 製薬業界またはCRO 業界での確立されたネットワーク。
- ・ 高い対人スキルと国際チームとの協働能力。
- ・ 国内外への出張に対応可能な柔軟性。

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## Company Description

ご紹介時にご案内いたします