



## Pharmacovigilance Reporting Associate

ICONクリニカルリサーチ合同会社での募集です。安全性情報（臨床開発・製販後...

### 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

ICONクリニカルリサーチ合同会社

#### 求人ID

1521185

#### 業種

CRO

#### 会社の種類

外資系企業

#### 雇用形態

正社員

#### 勤務地

東京都 23区

#### 給与

450万円 ~ 600万円

#### 勤務時間

09:00 ~ 17:30

#### 休日・休暇

【有給休暇】有給休暇は入社時から付与されます 入社7ヶ月目には最低10日以上 【休日】完全週休二日制 【休日】：土曜、日曜、祝...

#### 更新日

2025年02月06日 17:14

### 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

専門学校卒

#### 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 NJB2204577】

#### Overview

- ・ Serve as safety reporting processor or lead for multiple safety reporting providing management support as designated.
- ・ Recognize exemplify and adhere to ICON's values which center around our commitment to People Clients and Performance.
- ・ As a member of staff the employee is expected to embrace and contribute to our culture of process improvement with a

focus on streamlining our

processes adding value to our business and meeting client needs.

- Complete all departmental project activities accurately in accordance with ICON SOPs Study Specific Procedures regulatory requirements and client processes.
- Responsible for safety reporting or safety reporting intelligence activities on assigned projects working in a customer focused approach and an audit and inspection ready mindset.
- Demonstrate skills pertaining to client management safety reporting project scope submission compliance quality and budget.

Detail

- The following safety information case processing tasks related to clinical trials/post marketing of pharmaceutical products
- Receipt of information on Adverse event triage numbering confirmation of details entry into database/QC
- Creation of explanatory text for case course (Japanese and English) /QC
- Primary evaluation of the necessity of reporting to the PMDA / QC of the evaluation details
- Preparation of reports to PMDA/QC
- Escalation coordination etc. to customers
- Operations incidental to the above

\*Our Safety Reporting team will allow you to experience the ICCC study start up not just safety reporting. At first senior members will support you. You could expand your experience.

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## スキル・資格

- Experience required for any of the following
- PV experience especially PMDA submission experience required.

Experience with ICCC is better.

- 2+ years of CRA experience
- Fluency in Japanese business level English

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## 会社説明

1. 医薬品、医療機器、再生医療等製品、ワクチン等にかかる臨床開発、市販直後調査、製造販売後調査、臨床研究等の受託事業  
2. 労働者派遣事業