In Person

PMDA-ATC Pharmacovigilance Seminar 2025

Date:
February 26-28, 2025
9:30 to 17:00 JST (UTC+9)

Location: Tokyo (PMDA)

Target Audience: Regulators
Beginner to Intermediate level

Program Overview:

- Evaluation of cases of adverse drug reaction
- Identification of safety specifications
- Development and implementation of the RMP
- Labeling system and electronic labeling initiatives

Application Due: November 27, 2024

https://www.pmda.go.jp/english/symposia/0310.html

PMDA-ATC

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Pharmaceuticals and Medical Devices Agency