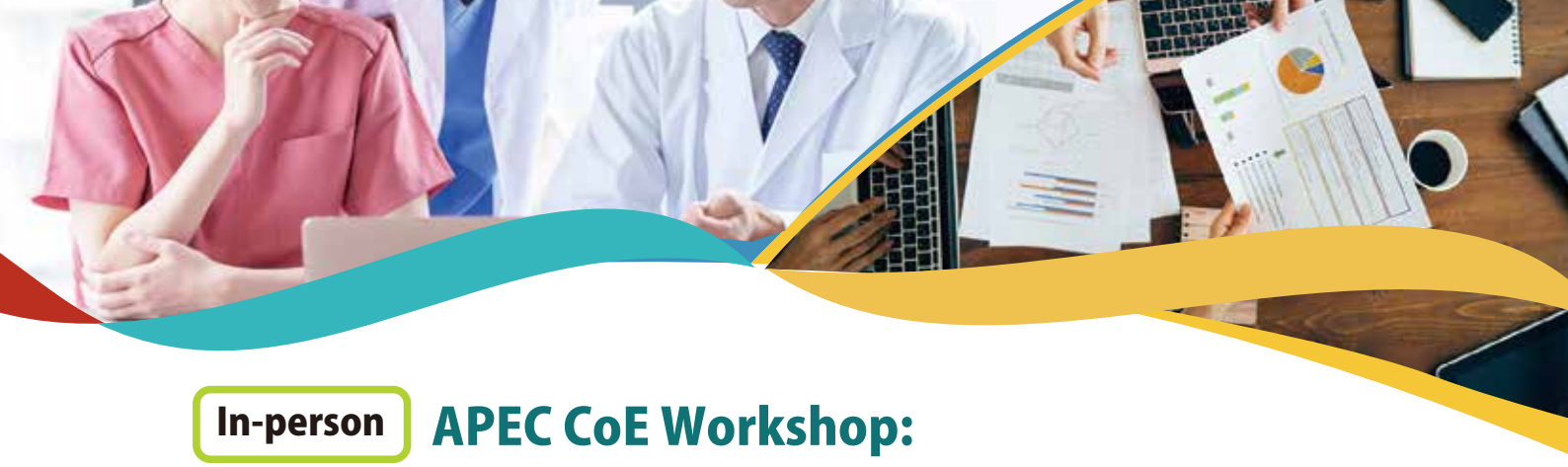


In-person **APEC CoE Workshop:**

PMDA-ATC Pharmacovigilance Seminar 2027



Dates:

February 17-19, 2027

Location:

Tokyo

Audience:

Beginner to Intermediate level

Regulators only

Applications due:

December 11, 2026

https://www.pmda.go.jp/english/symposia/0366.html

Program overview:

- Overview of pharmacovigilance in Japan
- Development and implementation of Risk Management Plan (RMP)
- Adverse Drug Reaction (ADR) case evaluation
- Identification of safety specifications
- Risk communication using labeling and risk minimization materials
- Utilization of Real-World Data for post-marketing drug safety

PMDA
独立行政法人 医薬品医療機器総合機構
Pharmaceuticals and Medical Devices Agency

APEC
Asia-Pacific Economic Cooperation

PMDA-ATC
PMDA Asia Training Center for pharmaceuticals and medical devices regulatory affairs