APEC CoE Workshop
PMDA-ATC Pharmacovigilance Seminar 2027

Agenda

Dates: February 17-19, 2027 Venue: Tokyo

(JST)

As of November 13, 2026

Day 1 Wednesday, February 17	Day 2 Thursday, February 18	Day 3 Friday, February 19
9:00-9:30 Registration	9:00-9:30 Registration	9:00-9:30 Registration
9:30-9:50 Opening Ceremony	9:30-11:40 **Session 5** (Group Work 1) ADR Case Evaluation/Identification of Safety Specification - Introduction - Group Discussion	9:30-10:20 **Session 9** Signal Detection and Use of AI in Pharmacovigilace
9:50-10:40 **Session 1** Overview of Post-marketing Drug Dafety in Japan		Break
		10:30-11:20 **Session 10** End-to-end Labeling including CCDS Labeling Principles, Electronic Labeling(e-product information) Initiatives, Patient Centric Labeling/ Product Information(PCPI) / Health Literacy
10:40-11:30 **Session 2** Outline of RMP in Japan		
11:30-12:40 Lunch	11:40-12:40 Lunch	11:20-12:20 **Session 11** (Group Work 2) Risk Management Plan - Introduction - Group Discussion
12:40-13:30 **Session 3** Development and Update of RMP	12:40-13:40 **Session 5** (Group Work 1) - Group Presentation - Q&A / Wrap up	12:20-13:20 Lunch
		13:20-14:20 **Session 11** (Group Work 2) - Group Presentation - Q&A / Wrap up
Break	Break	Break
13:40-15:50 **Session 4** Round Table Discussion Country Introduction	13:50-14:40 **Session 6** Communication of Safety Risk Information to Patients and Healthcare Professionals	14:30-15:20 **Session 12** Pharmacovigilance Methods
	14:40-15:30 **Session 7** Use of Published Safety Information at Clinical Site	15:20-16:10 **Session 13** Utilization of Real world Data for Post-marketing Drug Safety in PMDA
Break	Break	
16:00-16:20 Introduction of Case Studies	15:40-16:30 **Session 8** Overview of Internationlal Post-marketing Drug Safety	16:10-16:20 Feedback for Day 3
16:20-16:30 Feedback for Day 1	16:30-16:40 Feedback for Day 2	Break
Break		16:30-16:40 Closing Ceremony
16:45-18:00 Friendly Get-together		