

# 2026 APEC



Asia-Pacific  
Economic Cooperation

## Good Registration Management (GRM) Center of Excellence (CoE) Workshop

*Special Topic: Combination Products*

### Save the Date

Scheduled Date: August 26–28, 2026

Venue Location: Taipei, Chinese Taipei

### Target Audience

1. Regulatory professionals from authorities with hands-on experience in the management of regulatory reviews.
2. Regulatory professionals from industries with hands-on experience in the management of regulatory submissions.
3. Academia who are interested in learning GRevP or GSubP guidelines.
4. Professional bodies who are actively involved in training.

### Program Overview

1. Hybrid Training on Day 1 and Day 3, with limited online participation prioritized for overseas attendees.
2. In-Person Session on Day 2, featuring interactive case studies and group discussions with overseas and TFDA reviewers, as well as industry representatives.  
In-person participants will also be invited to the Welcome Reception Dinner.  
Limited travel support for eligible regulatory authority participants may be available.
3. Certificates will be awarded to participants who complete all three days of the workshop.

### Travel & Accommodations

Funding may be available for regulators from travel-eligible economies. Priority will be given to regulator representatives who are willing to share updates on drug review mechanisms in their economies during the workshop.

**CoE Hosting Institution: Taiwan Food and Drug Administration (TFDA)**

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**More Information: <https://apecgrmcoe2026.tw/>**



Agenda and  
more details