Asia-Pacific Economic Cooperation

FDA

2025 APEC
Good Registration Management (GRM) Center of Excellence (CoE) Workshop

Accelerated approval for drugs of unmet medical needs
Special Lecture: Combination Products

Save the Date

Scheduled Date : August 26–28, 2025
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. In-person training.
2. 2.5 days of plenary sessions designed with lectures, group discussions, and applied case studies for all attendees.

Travel & Accommodations

Funding for travel eligible economies may be available for regulators. Regulator representatives willing to share recent GRM implementation status in their member economies may be prioritized.

CoE Hosting Institution : Taiwan Food and Drug Administration (TFDA)
Contact Information : GRMCOE2025@gmail.com

QR code for more details