



2024 APEC Good Registration Management (GRM) Center of Excellence (CoE) Workshop

 *Save the Date*

Sep. 3-5, 2024 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. 3 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)

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