Purpose:
To promote GRevP and GSubP cooperatively

Long-term goals:
- Promote the concept of GRM
- Enhance mutual trust for regulatory convergence among APEC member economies by 2020

To help achieve timeliness, predictability, consistency, transparency, clarity, efficiency and high quality in the content and management of reviews

To enhance the quality and efficiency of the medical product registration process by improving the quality and management of submission
Specific Activities and Timeframes

Step 1: 2011-2012
- Gap Analysis Survey for setting the foundation for Stepwise GRevP Implementation

Step 2: 2011-2016
- Planned solutions to address Gap in GRM

Step 3: 2017-2019
- Assessing impact of GRM
  - Dissemination of GRM (continued)
  - Assessing the impact of training and implementation of GRM using KPIs

Step 4: 2020~
- Reaching the Goal for Implementing GRM

Gap Analysis Survey for setting the foundation for Stepwise GRevP Implementation
APEC GRM CoE Training Activities

Significant Activities

APEC GRM CoE Training Activities

August 2019-February 2020

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APEC Training

August 2019-February 2020

2019 APEC GRM CoE Workshop in Taipei
— September 17-19, 2019 (TFDA/RAPS Taiwan Chapter)

2019 APEC GRM Pilot CoE Workshop in Bangkok
— October 26-28, 2019 (Thai FDA)
KPI Assessment of GRM Implementation is on-going. The result will be shared with all in the 10th APAC.