

Full Agenda of 2024 APEC Medical Devices CoE Workshop

Day 1 – Aug. 28, 2024 (Wed.)

Time	Topic	Speaker
08:50 – 09:10	Registration	
09:10 – 09:20	Opening Remarks	<p>TFDA: Dr. Shin-Hun Juang Director-General, Taiwan Food and Drug Administration (TFDA), Ministry of Health and Welfare (MOHW), Chinese Taipei</p> <p>MD PWA Co-Champion: Ms. Miwa Kanematsu Principal Coordinator, Office of International Programs, Pharmaceuticals and Medical Devices Agency (PMDA), Japan</p>
09:20 – 09:30	Group Photo	
09:30 – 09:40	Roadmap and Core Curriculum of MD PWA	<p>MD PWA Co-Champion: Ms. Kanae Ohara Coordinator, Office of International Programs, PMDA, Japan</p>
09:40 – 09:50	Introduction of TFDA CoE Training Program	<p>Mr. Ching-Wei Chang Section Chief, Division of Medical Devices and Cosmetics, TFDA, MOHW, Chinese Taipei</p>
09:50 – 12:00	Lecture #1: Current Harmonization Status of Pre-Market Regulation in Each Economy Presentation (40 min) Coffee Break (10 min) Presentation (70 min) Q&A (10 min)	10 min per economy
12:00 – 13:20	Lunch	
13:20 – 14:20	Icebreaker Activities	Moderator

14:20 – 14:50	<p>Lecture #2: Medical Device and IVD Definition and Classification</p> <ul style="list-style-type: none"> • Definition of the Terms ‘Medical Device’ and ‘In Vitro Diagnostic (IVD) Medical Device’ (GHTF/SG1/N071:2012) • Principles of Medical Device Classification (GHTF/SG1/N77:2012) • Principles of In Vitro Diagnostic (IVD) Medical Devices Classification (IMDRF/IVD WG/N64FINAL:2021) 	<p>Dr. Sheng-Hui Liao Senior Engineer, Office of Medical Device Evaluation, Center for Measurement Standards, Industrial Technology Research Institute (ITRI), Chinese Taipei</p>
14:50 – 15:00	Coffee Break	
15:00 – 16:00	<p>Group Practice #1: Definition and Classification Practice (40 min) Group Presentation (15 min) Q&A (5 min)</p>	<p>Dr. Sheng-Hui Liao Senior Engineer, Office of Medical Device Evaluation, Center for Measurement Standards, Industrial Technology Research Institute (ITRI), Chinese Taipei</p>
18:30 – 20:30	Welcome Reception	

*The morning session will be open to the public.

Day 2 – Aug. 29, 2024 (Thu.)

Time	Topic	Speaker
09:00 – 09:30	Registration	
09:30 – 10:00	<p>Lecture #3: Review of (1) Essential Principles of Medical Device Safety & Performance and (2) Principles of Conformity Assessment</p> <ul style="list-style-type: none"> Essential Principles of Safety and Performance of Medical Devices and IVD Medical Devices (IMDRF/GRRP WG/N47FINAL:2018) Principles of Conformity Assessment for Medical Devices (GHTF/SG1/N78:2012) Principles of Conformity Assessment for In Vitro Diagnostic (IVD) Medical Devices (GHTF/SG1/N046:2008) 	<p>Mr. Shang-Ching Lin Associate Researcher, Division of Medical Devices and Cosmetics, TFDA, MOHW, Chinese Taipei</p>
10:00 – 12:00	<p>Group Practice #2: IVD Case Study</p> <p>Case Study Introduction (10 min) (1 case)</p> <p>Group Discussion (50 min)</p> <p>Coffee Break (15 min)</p> <p>Group Presentation (30 min)</p> <p>Q&A (15 min)</p>	<p>Mr. Shang-Ching Lin Associate Researcher, Division of Medical Devices and Cosmetics, TFDA, MOHW, Chinese Taipei</p>
12:00 – 13:30	Lunch	
13:30 – 16:00	<p>Group Practice #3: Medical Device Case Study</p> <p>Case Study Introduction (15 min) (2 cases)</p> <p>Group Discussion (60 min)</p> <p>Coffee Break (15 min)</p> <p>Group Presentation (45 min)</p> <p>Q&A (15 min)</p>	<p>Ms. Yu-Hui Huang Specialist, Division of Medical Devices and Cosmetics, TFDA, MOHW, Chinese Taipei</p> <p>Mr. You-Lin Lee Associate Reviewer, Division of Medical Devices and Cosmetics, TFDA, MOHW, Chinese Taipei</p>

Day 3 – Aug. 30, 2024 (Fri.)

Time	Topic	Speaker
09:00 – 09:30	Registration	
09:30 – 10:10	<p>Lecture #4: Clinical Evaluation</p> <ul style="list-style-type: none"> Clinical Investigation (IMDRF/MDCE WG/N57FINAL: 2019) Clinical Evaluation (IMDRF/MDCE WG/N56FINAL:2019) Clinical Evidence (IMDRF/MDCE WG/N55FINAL:2019) <p>Q&A</p>	<p>Dr. Daisuke Fujisawa Principal Reviewer Office of Medical Device II, PMDA, Japan</p>
10:10 – 10:20	Group Photo	
10:20 – 10:40	Coffee Break	
10:40 – 11:00	<p>Expectations from the Workshop and Next Steps</p> <ul style="list-style-type: none"> TFDA (3 mins) APEC RHSC MD PWA Co-Champion (3 mins) APEC RHSC MD PWA Sub-Champions (3 mins each) Members of the planning committee or participants (2 mins each) 	<p>TFDA: Dr. Chia-Hung Chien Director, Division of Medical Devices and Cosmetics, TFDA, MOHW, Chinese Taipei</p> <p>MD PWA Co-Champion: Ms. Miwa Kanematsu Principal Coordinator, Office of International Programs, PMDA, Japan</p>
11:00 – 11:20	Certificate Award Ceremony	<p>Dr. Der-Yuan Wang Deputy Director-General, TFDA, MOHW, Chinese Taipei</p>
11:20 – 11:30	Closing Remarks	<p>Dr. Der-Yuan Wang Deputy Director-General, TFDA, MOHW, Chinese Taipei</p>
11:30 – 13:00	Lunch	
13:00 – 17:00	Manufacturing Site Visit	Regulators Only