

## Full Agenda of 2024 APEC Medical Devices CoE Workshop

Day 1 – Aug. 28, 2024 (Wed.)

| Time          | Topic   | Speaker  |
|---------------|---|--|
| 09:00 – 09:30 | Registration  |  |
| 09:30 – 09:40 | Opening Remarks   | <p><b>TFDA:</b><br/> <b>Dr. Shou-Mei Wu</b><br/>           Director-General, Taiwan Food and Drug Administration (TFDA), Ministry of Health and Welfare (MOHW), Chinese Taipei</p> <p><b>MD PWA Co-Champion:</b><br/> <b>Ms. Miwa Kanematsu</b><br/>           Principal Coordinator, Office of International Programs, Pharmaceuticals and Medical Devices Agency (PMDA), Japan</p> |
| 09:40 – 09:50 | Group Photo   |  |
| 09:50 – 10:00 | Roadmap and Core Curriculum of MD PWA   | <p><b>MD PWA Co-Champion:</b><br/> <b>Ms. Kanae Ohara</b><br/>           Coordinator, Office of International Programs, PMDA, Japan</p>  |
| 10:00 – 10:10 | Introduction of TFDA CoE Training Program   | <p><b>Mr. Ching-Wei Chang</b><br/>           Section Chief, Division of Medical Devices and Cosmetics, TFDA, MOHW, Chinese Taipei</p>  |
| 10:10 – 12:00 | Lecture #1:<br>Current Harmonization Status of Pre-Market Regulation in Each Economy<br>Presentation (30 min)<br>Coffee Break (20 min)<br>Presentation (45 min)<br>Q&A (15 min) | 15 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"> <li>• Definition of the Terms ‘Medical Device’ and ‘In Vitro Diagnostic (IVD) Medical Device’ (GHTF/SG1/N071:2012)</li> <li>• Principles of Medical Device Classification (GHTF/SG1/N77:2012)</li> <li>• Principles of In Vitro Diagnostic (IVD) Medical Devices Classification (IMDRF/IVD WG/N64FINAL:2021)</li> </ul> | <p><b>Dr. Sheng-Hui Liao</b><br/>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:<br/>Definition and Classification Practice (40 min)<br/>Group Presentation (15 min)<br/>Q&amp;A (5 min)</p>  | <p><b>Dr. Sheng-Hui Liao</b><br/>Senior Engineer, Office of Medical Device Evaluation, Center for Measurement Standards, Industrial Technology Research Institute (ITRI), Chinese Taipei</p> |
| 17:30 – 19:30 | 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 &amp; Performance and (2) Principles of Conformity Assessment</p> <ul style="list-style-type: none"> <li>• Essential Principles of Safety and Performance of Medical Devices and IVD Medical Devices (IMDRF/GRRP WG/N47FINAL:2018)</li> <li>• Principles of Conformity Assessment for Medical Devices (GHTF/SG1/N78:2012)</li> <li>• Principles of Conformity Assessment for In Vitro Diagnostic (IVD) Medical Devices (GHTF/SG1/N046:2008)</li> </ul> | <p><b>Mr. Shang-Ching Lin</b><br/>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&amp;A (15 min)</p>  | <p><b>Mr. Shang-Ching Lin</b><br/>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&amp;A (15 min)</p>  | <p><b>Ms. Yu-Hui Huang</b><br/>Specialist, Division of Medical Devices and Cosmetics, TFDA, MOHW, Chinese Taipei</p> <p><b>Mr. You-Lin Lee</b><br/>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"> <li>Clinical Investigation (IMDRF/MDCE WG/N57FINAL: 2019)</li> <li>Clinical Evaluation (IMDRF/MDCE WG/N56FINAL:2019)</li> <li>Clinical Evidence (IMDRF/MDCE WG/N55FINAL:2019)</li> </ul> <p>Q&amp;A</p>                  | <p><b>Dr. Daisuke Fujisawa</b><br/>Principal Reviewer<br/>Office of Medical Device II,<br/>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"> <li>TFDA (3 mins)</li> <li>APEC RHSC MD PWA Co-Champion (3 mins)</li> <li>APEC RHSC MD PWA Sub-Champions (3 mins each)</li> <li>Members of the planning committee or participants (2 mins each)</li> </ul> | <p><b>TFDA:</b><br/><b>Dr. Chia-Hung Chien</b><br/>Director, Division of Medical<br/>Devices and Cosmetics,<br/>TFDA, MOHW, Chinese<br/>Taipei</p> <p><b>MD PWA Co-Champion:</b><br/><b>Ms. Miwa Kanematsu</b><br/>Principal Coordinator,<br/>Office of International<br/>Programs, PMDA, Japan</p> |
| 11:00 – 11:20 | Certificate Award Ceremony   | <p><b>Dr. Der-Yuan Wang</b><br/>Deputy Director-General,<br/>TFDA, MOHW, Chinese<br/>Taipei</p>   |
| 11:20 – 11:30 | Closing Remarks  | <p><b>Dr. Der-Yuan Wang</b><br/>Deputy Director-General,<br/>TFDA, MOHW, Chinese<br/>Taipei</p>   |
| 11:30 – 13:00 | Lunch  |   |
| 13:00 – 17:00 | Manufacturing Site Visit   | Regulators Only   |