Agenda of 2020 APEC Medical Devices CoE Workshop

<table>
<tr><th>1st Part:
August 29 to September 7</th><th>2nd Part:
September 8 to September 11</th></tr>
<tr><td>Online Course 1: Introduction Session
• Opening Remarks
• Roadmap and Core Curriculum of Medical Device PWA
• CoE Training Program</td><td rowspan="4">2nd Videoconference
Case Study of Conformity Assessment: Medical Device Session</td></tr>
<tr><td>Online Course 2
• Basic Scheme of Conformity Assessment Procedure and Classification for Medical Devices (GHTF/SG1/N77&N78)
• Summary of Essential Principles for Medical Devices (IMDRF/GRRP WG/N47)
• Introduction of Case Study: Medical Device Session</td></tr>
<tr><td>Online Course 3
• Basic Scheme of Conformity Assessment Procedure and Classification for In Vitro Diagnostic (IVD) Medical Devices (GHTF/SG1/N045&N46)
• Summary of Essential Principles for In Vitro Diagnostic (IVD) Medical Devices (IMDRF/GRRP WG/N47)
• Introduction of Case Study: IVD Session</td><td>3rd Videoconference
Case Study of Conformity Assessment: IVD Session</td></tr>
<tr><td>1st Videoconference
• Q&A for Online Courses
• Q&A for the current harmonization status of pre-market regulation in each APEC member economy</td><td>4th Videoconference
Presentation of Case Studies, Q&A, and Closing</td></tr>
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