

The 22nd AHWP Annual Meeting Program

The Grand Hotel, Nelson Mandela Road, Vasant Kunj, New Delhi

4 - 8 December 2017

Day 1 Agenda: 4 December 2017 (Monday)

08:00-09:10 Hrs	Registration of the Delegates	
TIME (Hrs)	INAGURAL SESSION (09:00 – 10:40)	
09:10-09:20	Opening Address	Dr. RK Vats , Additional Secretary, Ministry of Health and Family Welfare, Government of India
09:20-09:30	Address	Mr. Zamane Abdul Rahman , AHWP Vice Chair, Chief Executive, Medical Device Authority, Ministry of Health Malaysia, Malaysia
09:30-09:40	Address	Dr. Henk Bekedam , WHO Representative to India
09:40-09:50	Address	Dr. Jagdish Prasad , Director General Health Services, Ministry of Health and Family Welfare, Government of India
09:50-10:00	Address	Ms. Preeti Sudan , Secretary, Ministry of Health and Family Welfare, Government of India
10:00-10:10	Address	HE Ms. Anupriya Patel , Honourable Minister of State, Health and Family Welfare, Government of India
10:10-10:20	Address	HE Mr. Ashwini Kumar Chaubey , Honourable Minister of State, Health and Family Welfare, Government of India
10:20 -10:30	Address	HE Mr JP Nadda , Honourable Union Minister, Health and Family Welfare, Government of India
10:30-10:35	Vote of Thanks	Dr G. N Singh , Drugs Controller General of India, DGHS, Ministry of Health and Family Welfare, Government of India
10:35-11:00	Photo Session & High Tea	

Day 1 Agenda: 4 December 2017 (Monday)

TIME (Hrs)	TECHNICAL SESSION I: Overview of Medical Devices Regulation (11:00 –13:10)
	<p>Chairs : Mr. Zamane Abdul Rahman, AHWP Vice Chair, Chief Executive, Medical Device Authority, Ministry of Health Malaysia, Malaysia</p> <p>Sh. Sudhir Kumar, Joint Secretary (Regulation), Ministry of Health and Family Welfare, Government of India</p> <p>Co-Chairs : Dr. G.N. Singh, Drugs Controller General (I), CDSCO Dr. S.E Reddy, Joint Drugs Controller (I), CDSCO</p>
	Speakers:
11.00-11.10	<p>Introductory Remarks</p> <p>Mr. Ali M. Al-Dalaan, TC Chair, AHWP, Executive Director, Medical Devices Sector, Saudi FDA, Kingdom of Saudi Arabia</p>
11:10-11:50	<p>An overview of regulations of Medical Device and IVDs in India (India Regulatory update)</p> <p>Dr. V.G Somani, Joint Drugs Controller (I), CDSCO</p>
11.50-12:10	<p>An overview of regulations of medical devices and IVDs in Singapore (Singapore Regulatory Update)</p> <p>Dr. Rama Sethuraman , Deputy Director, Health Science Authority, Singapore</p>
12:10-12:30	<p>An overview of regulations of medical devices and IVDs in Japan (Japan Regulatory Update)</p> <p>Mr. Hiroshi YAGINUMA , Director, Office of Regenerative Medicine Product Evaluation Medical Device Evaluation and Licensing Division Pharmaceutical Safety and Environmental Health Bureau, MHLW</p>
12:30-12:50	<p>An overview of regulations of medical devices and IVDs in South Korea (South Korea Regulatory Update)</p> <p>Mr. Seil Park, Assistant Director , Division of High-tech devices, Department of MD evaluation Korea MFDS</p> <p>Mr. Young Wook Ahn, Assistant Director Division of In-vitro Diagnostic Devices Department of MD evaluation</p>
12:50-13:10	<p>An overview of regulations of medical devices and IVDs in Australia (Australia Regulatory Update)</p> <p>Mr. Michael Flood , Ex-TGA, Locus Consulting Pty Ltd Australia</p>

13:10-14:10	Lunch
	TECHNICAL SESSION II (14:10 –15:20)
14:10-14:30	<p>An overview of regulations of medical devices and IVDs in South Africa (South Africa Regulatory Update)</p> <p>Ms. Andrea Julsing Keyter, Deputy Director : Medical Devices, National Department of Health Inspectorate & Law Enforcement Unit, South Africa</p>
14:30-15:00	<p>Medical Device Single Audit Program (MDSAP) – The MDASP Experience</p> <p>Mr. Dominik Reterski, VP Quality & Regulatory, Asia Pacific, Medtronic</p>
15:00-15:20	<p>Design controls, Risk Management, Verification and Validation and product release</p> <p>Mr. Fred Viaud, VP Quality & Regulatory, Philips Healthcare</p>
	TECHNICAL SESSION III (15:20 –16:50)
15:20-15:40	<p>Adapting your ISO 13485 to the new Requirements</p> <p>Mr. Grant Ramaley, Convener, Medical Device Working Group for ISO 13485 International Accreditation Forum, Aseptico, Director of Regulatory Affairs</p>
15:40-16:00	Tea Break
	TECHNICAL SESSION III (Continue)
16:00-16:25	<p>Role of Technical Standards and updates on international technical standards</p> <p>Dr. Peter Linders, Director, Standards & Regulations, Regulatory Standards, Philips Healthcare</p>
16:25-16:50	<p>Clinical Trial Environment - Australia</p> <p>Dr. Catherine Bourgeois, Vice President, Field Clinical Affairs, Emerging Markets & ANZ, Abbott</p>
	PANEL DISCUSSION I (16:50-17:40)
16:50-17:40	<p>Accessibility, Affordability of Medical Devices and Safeguards of Consumer Rights</p> <p>Moderator: Mr. Pavan Choudhary, Director General and Chairman of MTaI</p> <p>Speakers: Ms. Jaishree Gupta, Consumer Online Foundation, New Delhi Dr. V. G. Somani, Joint Drugs Controller, CDSCO Dr. Madhur Gupta, Technical Officer –Pharmaceuticals, WHO, Country office for India Ms. Manisha Sridhar, Regional Advisor, WHO SEAR office India Dr. Y. K Gupta, Prof. & Head Dept. of Pharmacology, AIIMS New Delhi Ms. Leena Menghaney, Head South Asia, Access Campaign, Medecins Sans</p>



	Frontieres, India Mr. Himanshu Baid , MD, Poly Medicure Limited, India
17:40-17:50	Closing Remarks Dr. Jeong-Rim Lee , TC Co-Chair, Director, Cardiovascular Devices Division ,Department of MD evaluation , Ministry of Food and Drug Safety (MFDS)

CULTURAL EVENING FOLLOWED BY GALA DINNER BY INVITATION (18:30 ONWARDS):

END OF DAY 1

Day 2 Agenda: 5 December 2017 (Tuesday)

TIME (Hrs)	TECHNICAL SESSION IV: AHWP Playbook Training Workshop (9:00-10:40)
09:00-09:10	<p>Welcome speech – Technical Committee Capacity Building Program</p> <p>Ms. Tran Quan, Capacity Building Team Leader, VP, Regulatory Affairs, Asia Pacific Medical Technology Association (APACMed), Laos PDR</p>
09.10-09.20	<p>Opening Speech Play Book Training Session</p> <p>Mr. Ali M. Al-Dalaan, (TC Chair, AHWP), Executive Director, Medical Devices Sector Saudi Food & Drug Authority, Kingdom of Saudi Arabia</p>
	<p>Introduction of the 2 Day Training Program</p> <p>Ms. Joanna Koh, AHWP PB Program Co-ordinator & Lead Trainer, Principal Consultant MDnet Regulatory Consultants, Singapore</p>
09:20-10:00	<p>Preparatory steps to MD Controls –ASEAN experience / Centering on the AMDD elements</p> <p>Mr. Zamane Abdul Rahman Chief Executive, Medical Device Authority, Ministry of Health Malaysia</p>
10:00-10:40	<p>“Takes 2 to Tango” – CSDT/EP and standards</p> <p>Mr. Seet Wing Gang, Head Regulatory Intelligence, Greater Asia, Becton Dickinson</p>
10:40-11:00	Tea Break
	TECHNICAL SESSION V (11:00-13:00)
11:00-11:40	<p>Classification Rules GMD/IVD D –Why The Rules? Is there an alternative?</p> <p>Mr. Greg LeBlanc, M.Sc., RAC, Director, Regulatory Affairs and Quality Systems, Cook (Canada) Inc.</p>
11:40-12:20	<p>CAB Role and Grouping – A balance between Regulatory Controls and Processes with the Economics of MD Industry</p> <p>Dr. Vincent Lam , MHS Manager and Senior Product Specialist, TUV SUD Product Service</p>
12:20-13:00	<p>Objectives and limitations: In-country Lab Testing</p> <p>Dr. G. Kalyan Varma, VP, Global Business Field-Electrical, Business Stream Products-IMEA, TUV Rheinland</p> <p>Mr. Petra Kaars-Wiele, Senior Director Regulatory, Quality & Labelling, Abbott Diagnostics</p>

13:00-14:00	Lunch
	TECHNICAL SESSION VI (14:00- 14:50)
14:00-14:25	Clinical Investigation and evaluation for Medical Devices: Regulator perspective Dr. Mijung Son, Regulator (MFDS) Korea
14:25-14:50	Clinical Investigation and evaluation for Medical Devices: Regulator perspective Mr. Arthur Brandwood, Brandwood Biomedical, Founder and Principal Consultant
	PANEL DISCUSSION II (14:50- 15:40)
14:50-15:40	Clinical Investigations and Real World Evidence Moderator: Ms. Sumati Randeo Director Global Strategy Regulatory Affairs & Advocacy Abbott Laboratories Panellists: Ms. Katy Peterson , Director, Global RA Boston Scientific Ms. Kate Hyeong Joo Kim Director, Regulatory Strategy & Innovation, ASPAC Director Regulatory Affairs, North Asia, Johnson & Johnson Medical, Republic of Korea Mr. Kwan Han Ong , Regional Associate Director, Clinical Affairs, Asia Pacific Regional Associate Director, Regulatory Affairs, APAC Dr. VG Somani , Joint Drugs Controller (I), CDSO
15:40-16:00	Tea Break
	PANEL DISCUSSION III (16:00-16:40)
16:00-16:40	Regional Agreements , Challenges (TRIPS Agreement) and access to Medical Device Provisions In TRIPS Moderator: Mr. Rajiv Nath , Forum Coordinator, Association of Indian Medical Device Industry (AiMeD) Panellists: Dr. VG Somani , Joint Drugs Controller (I), CDSO Dr. S.B. Sinha , Advisor, Healthcare Technology, National Health Systems Resource Centre, New Delhi Dr. Anand Grover , Senior Advocate, Supreme Court of India, the Director of the HIV/AIDS Unit of Lawyer's Collective (India) Ms. Aradhana Johri , Former Secretary Department of Pharmaceuticals, Ministry of Chemicals and Fertilizers; Secretary of Finance (Disinvestment); Former Additional Secretary, Ministry of

	Health and Family Welfare, Government of India
	TECHNICAL SESSION VII (16:40-17:20)
16:40-17:00	<p>Performance evaluation of IVD regulation</p> <p>Dr. Benny Ons, Director of Regulatory Affairs at BD Diagnostics and BD Biosciences Europe, Becton Dickinson International Becton Dickinson B.V.</p>
17:20-17:30	<p>Closing Remarks</p> <p>Ms. Joanna Koh, AHWP PB Program Co-ordinator & Lead Trainer, Principal Consultant MDnet Regulatory Consultants, Singapore</p>
17:00-17:20	<p>Liaison Member Updates:</p> <p>(i) GS1 Updates</p> <p>(ii) UDI Databases: Useful Tools for Regulatory Controls. How to utilize them in the MD Life Cycle especially in referencing for Pre Market Assessments</p> <p>Ms. Ulrike Kreysa, VP, Healthcare, GS1 Global Office</p>

END OF DAY 2

Day 3 Agenda: 6 December 2017 (Wednesday)

TIME (Hrs)	TECHNICAL SESSION VIII: AHWP Technical Committee Meeting (9:00-10:40)
09:00-09:10	<p>Recap of Play Book session 1 and introduction of session 2</p> <p>Ms. Joanna Koh, Principal Consultant, MDnet Regulatory Consultants, Singapore</p>
09:10-09:55	<p>Post Market - Medical Device Post market for Digital Healthcare: Cybersecurity prevention for MDs</p> <p>Mr. John Ramesh, MD, TUV Rheinland LLC, Oman & Regional Field Manager, Business Solutions, IMEA & APAC</p>
09:55-10:40	<p>Post Market - A Growing Global Concern: Counterfeit Medical Devices</p> <p>Dr. Nazeeh S. Alothmany, Vice Executive President, Medical Device Sector, Saudi FDA, KSA</p>
10:40-11:00	Tea Break
	TECHNICAL SESSION IX (11:00-11:30)
11:00-11:30	<p>Advertisement and Labelling</p> <p>Mr. Er Alfred Kwek, Director, Public Affairs, Edwards Life Sciences S PL</p>
	PANEL DISCUSSION IV (11:30-12:15)
11:30-12:00	<p>Play Book initiatives and Road ahead</p> <p>Moderator: Mr. Scott Sanderson, 3M Health Care, International Regulatory Affairs and Quality Compliance Leader, Medical Division, Minnesota</p> <p>Panellists:</p> <p>Dr. Arianti Anaya, Director of Medical Devices and Household Health Pdts Evaluation, MOH, Indonesia</p> <p>Ms. Agnes Sitta Kijo, Manager, Medical Devices; Diagnostics Registration, Tanzania Food & Drugs Authority</p> <p>Mr. Grant Ramaley, Aseptico, Director of Regulatory Affairs</p>
12:00-12:15	<p>Closing of AHWP PlayBook Training Program for the Current Cycle</p> <p>Ms. Joanna Koh, AHWP PB Program Co-ordinator & Lead Trainer, Principal Consultant MDnet Regulatory Consultants, Singapore</p>
	TECHNICAL SESSION X: AHWP Technical Committee (TC) Workshop (12:15 – 13:00)

12:15-13:00	Device & Operation Risks of Tele-medicine and e-Health: Regulatory Consideration across Jurisdictions Ms. Ann Graves, VP, International Regulatory Affairs, Abbott
13:00-14:00	Lunch
	TECHNICAL SESSION XI (14:00 – 15:00)
14:00-14:30	Regulatory on Medical Devices using 3D Printing – From Manufacturing to AE Reporting (e.g. QMS, QMS audit, pre-market submissions, clinical trial, AE reporting and recall) Dr. Jang Yong Choi, Deputy Director, Division of Medical Device Safety Evaluation Medical Device Safety Bureau, Ministry of Food and Drug Safety (MFDS)
14:30-15:00	Diagnostic Devices based on AI and Big Data Analytics –Thoughts on Validation and Risk Management Ms. Nicole Taylor Smith, Senior Director, Global Regulatory Affairs Policy & Intelligence, Johnson & Johnson

	PANEL DISCUSSION V (15:00- 15:40)
15:00- 15:40	Panel discussion: Challenges and issues to regulate an innovative medical devices: Barriers and Enablers to development Moderator: Ms. Miang Tanakasemsub, Head, Regulatory Affairs (Asia & Russia), Alcon Panellists: Dr. S. Eswara Reddy, Joint Drugs Controller(I), CDSCO Dr. Manisha Sridhar, Regional Advisor, WHO SEAR office India Representative from Department of Electronics & Information Technology (DeITY) Representative from Quality Council of India Ms. Kirti Arora, Associate Director, Regulatory and Quality, Boston Scientific India Mr. Matthew Hein, Senior International Trade Specialist, Industry & Analysis, U.S. Department of Commerce International Trade Administration, Office of Health and Information Technologies
15:40-16:00	Tea Break
	PARALLEL SESSIONS (16:00- 18:00)
	PARALLEL SESSION - I : AHWP TC Workshop Session - HALL 1

16:00-16:25	<p>Changing global regulatory environment – opportunity and threat in industry - focused on premarket</p> <p>Mr. Jeongpyo Hong, Manager, Regulatory Affairs, Health & Medical Equipment Business, Samsung Electronics</p>
16:25-16:50	<p>Pre- market approval – Essential Principles for safety, quality and performance</p> <p>Mr. Michael Flood, Principal, Locus Consulting Pty Ltd</p>
16:50-17:15	<p>Use of Real World Evidence in clinical and regulatory decision from Industry experience</p> <p>Dr. Justin Yoo, Government Affairs, Health Economics & Reimbursement, Manager Corporate Relations, Ambassador, St. Jude Medical Korea YH, Abbott</p>
17:15-17:40	<p>Post-market control of medical devices-How far we have gone towards harmonization</p> <p>Ms. Jennifer MAK Kit-shu, Senior Electronics Engineer (Medical Device Control Office), Department of Health, Medical Device Control Office, Department of Health, Hong Kong China</p>

17:40 -17:50	<p>TGA Reforms for Medical Devices & IVDs and Clinical trial landscape in major countries in APAC</p> <p>Ms. Mie Ohama, Principal clinical quality specialist at Medtronic Clinical Research Institute, International quality, Sydney Australia</p>
17:50-18:00	<p>Closing Remarks</p> <p>Sh. Aseem Sahu, Dy. Drugs Controller (I), CDSCO</p> <p>Dr. Jeong-Rim Lee, TC Co-Chair, Director, Cardiovascular Devices Division, Department of MD evaluation, Ministry of Food and Drug Safety (MFDS)</p>
	<p>PARALLEL SESSION -II : India Regulatory Session - HALL 2 (16:00-18:00)</p>
16:00 -16:25	<p>Medical Devices Rules, 2017- Technical Requirements of manufacture and import of medical devices</p> <p>Sh. Aseem Sahu, Deputy Drugs Controller (I), CDSCO</p> <p>Sh. Sunil Kulshrestha, Assistant Drugs Controller(I), CDSCO</p>
16:25-16:50	<p>Medical Devices Rules, 2017- Investigational Medical Devices/new IVD approval</p> <p>Sh. Ravi Kant Sharma, Assistant Drugs Controller(I), CDSCO</p> <p>Sh. Somnath Basu, Assistant Drugs Controller(I), CDSCO</p>
16:50-17:50	<p>Panel Discussion: Opportunities and Way Forward in Manufacturing of Medical Devices (MD)</p>

and IVD sector –“Make in India Program” by Invest India (DIPP)

Mr. Sanjay Arudi, Senior Director Regulatory Affairs Sustainable Healthcare Solutions, GE Healthcare

Mr. R. Asok Kumar, Vice President (QA&RA), Johnson & Johnson Medical, India

Mr. Himanshu Baid, MD, Poly Medicure Limited, India

Mr. Sudhakar Mairpadi, Director - Quality & Regulatory, Philips Electronics India Limited (Health Care sector)

Mr. Rajiv Nath, Forum Coordinator, Association of Indian Medical Device Industry (AiMeD)

Dr. V. G. Somani, Joint Drugs Controller (I), CDSCO

17:50-18:00

Q & A and Closing Remarks

Dr. V. G. Somani, Joint Drugs Controller (I), CDSCO

END OF DAY 3

Day 4 Agenda: 7 December 2017 (Thursday)

TIME (Hrs)

21st AHWP TC Meeting

09:00-10:40

AHWP TC & WG Leaders Meeting with TC Advisors (Closed Meeting)

AHWP & TC & WG Leaders & TC Advisors

10:40-11:00

Tea Break

11:00-11:05

Welcome Speech

11:05-11:25

Opening speech by TC Chair

-Roll call by **Mr. Bryan So**, AHWP Secretariat

-Adoption of Agenda by TC Chair

-Announcement of the Election Arrangement for Office Bearers of AHWPTC and AHWPTC WGs

Mr. Ali M. Al-Dalaan,(TC Chair, AHWP),Executive Director, Medical Devices Sector, Saudi FDA, Kingdom of Saudi Arabia

Mr. Bryan So, AHWP Secretariat, Principal Consultant, Smart HealthCare, MedTech and Optics

	Unit, Hong Kong Productivity Council, Hong Kong SAR, China
	WORKING GROUP UPDATES (11:25- 15:25)
11:25-13:05 (20min each)	<p>Working Group 1</p> <p>Working Group 2</p> <p>Working Group 3</p> <p>Working Group 4</p> <p>Working Group 5</p> <p>Working Group Chair & Co-chairs</p>
13:05-14:05	Lunch
14:05- 15:25 (20min each)	<p>Working Group 6</p> <p>Working Group 7</p> <p>Working Group 8</p> <p>STG</p> <p>Working Group Chair & Co-chairs</p>
15:25-15:45	Tea Break
15:45-16:15	<p>Highlight of AHWP PB Training</p> <p>Ms. Joanna Koh, Principal Consultant, MDnet Regulatory Consultants, Singapore</p>
16:15-16:45	<p>Speech by TC Advisors Representative</p> <p>TC Advisor Representative</p>
16:45-16:55	<p>Closing Remarks</p> <p>Mr. Er Alfred Kwek, Director, Public Affairs, Edwards Life Sciences S PL</p> <p>Mr. Ali M. Al-Dalaan, (TC Chair, AHWP), Executive Director, Medical Devices Sector, Saudi FDA, Kingdom of Saudi Arabia</p>

GALA DINNER (19:00 ONWARDS): END OF DAY 4

Day 5 Agenda: 8 December 2017 (Friday)

TIME (Hrs)	21 st AHWP Annual Meeting
09:00-09:30	<p>Speech by officials, Ministry of Health and Family Welfare, Government of India</p> <ul style="list-style-type: none"> A. Opening Ceremony B. Congratulation Address C. Opening Speech by AHWP Vice-Chair <p>Mr. Zamane Abdul Rahman, AHWP Vice Chair, Chief Executive, Medical Device Authority, Ministry of Health Malaysia, Malaysia Group Photograph</p>
09:30-09:45	<ul style="list-style-type: none"> A. Roll Call B. Adoption of Agenda C. Adoption of 21st AHWP Annual Meeting Minutes <p>Mr. Zamane Abdul Rahman, AHWP Vice Chair, Chief Executive, Medical Device Authority, Ministry of Health Malaysia, Malaysia</p> <p>Supported by</p> <p>Mr. Bryan So, AHWP Secretariat, Principal Consultant, Smart HealthCare, MedTech and Optics Unit, Hong Kong Productivity Council, Hong Kong SAR, China</p>
09:45-10:15	<p>Updates by AHWP and AHWP TC</p> <p>Ms. Tran Quan, AHWP Vice Chair, Vice President, Regulatory Affairs, Asia Pacific Medical Technology Association (APACMed), Laos PDR</p> <p>Mr. Ali M. Al-Dalaan, (TC Chair, AHWP), Executive Director, Medical Devices Sector, Saudi FDA, Kingdom of Saudi Arabia</p>
10:15-10:30	<p>Announcement of Election Arrangement for Office Bearers of AHWP</p> <p>Mr. Bryan So, AHWP Secretariat, Principal Consultant, Smart HealthCare, MedTech and Optics Unit, Hong Kong Productivity Council, Hong Kong SAR, China</p>
10:30-11:00	Tea Break

<p>11:00-12:00 (15min each)</p>	<p>AHWP Member Economy updates</p> <p>China - Mr. GAO Guo Biao, Deputy Director General of Medical Device, Registration Department, China Food and Drug Administration, People's Republic of China</p> <p>India - Dr. V. G. Somani, Joint Drugs Controller, CDSCO, India</p> <p>Tanzania - Mr. Mitangu Adam Fimbo, Director of Medicines and Complementary Products., Tanzania FDA</p> <p>APEC -Dr. Arianti Anaya, Director of Medical Devices and Household Health Pds Evaluation, MOH, Indonesia</p> <p>ASEAN- Mr. Zamane Abdul Rahman, Chief Executive, Medical Device Authority, Ministry of Health Malaysia</p>
<p>12:00-12:30 (10 min each)</p>	<p>AHWP Liaison Member Updates</p> <p>APACMed</p> <p>Mr. Fredrik Nyberg, CEO APACMed</p> <p>DITTA</p> <p>Dr. Peter Linders, Director, Standards & Regulations, Regulatory Standards, Philips Healthcare</p>
<p>12:30-13:00</p>	<p>Endorsement of New Member Economies</p> <p>A. Kenya (TBC) To be confirmed by AHWP</p> <p>Endorsement of New Liaison Members</p> <p>GMDN Speech by Representative</p> <p>Mr. Zamane Abdul Rahman, AHWP Vice Chair, Chief Executive, Medical Device Authority, Ministry of Health Malaysia, Malaysia</p> <p>Supported by</p> <p>Mr. Bryan So, AHWP Secretariat, Principal Consultant, Smart HealthCare, MedTech and Optics Unit, Hong Kong Productivity Council, Hong Kong SAR, China</p>

13:00-14:15	Lunch
14:15-14:35	<p>Secretariat Updates - Secretariat Report - Financial Report</p> <p>Mr. Bryan So, AHWP Secretariat, Principal Consultant, Smart HealthCare, MedTech and Optics Unit, Hong Kong Productivity Council, Hong Kong SAR, China</p>
14:35-14:45	<p>Resolutions:</p> <ul style="list-style-type: none"> A. STG change to WG 9 B. Amendment to AHWP TOR/HR C. Participation of WG activities <p>Mr. Zamane Abdul Rahman, AHWP Vice Chair, Chief Executive, Medical Device Authority, Ministry of Health Malaysia, Malaysia</p> <p>Supported by</p> <p>Mr. Bryan So, AHWP Secretariat, Principal Consultant, Smart HealthCare, MedTech and Optics Unit, Hong Kong Productivity Council, Hong Kong SAR, China</p>
14:45-16:00	<ul style="list-style-type: none"> A. Election of AHWP Leadership of 2018-2020 B. Election C. Endorsement of Newly Elected Office Bearers of AHWP D. Endorsement of Newly Elected Office Bearers of AHWP TC E. Endorsement of Newly Elected Office Bearers of AHWP TC Working Group <p>Mr. Bryan So, AHWP Secretariat, Principal Consultant, Smart HealthCare, MedTech and Optics Unit, Hong Kong Productivity Council, Hong Kong SAR, China</p>
16:00-16:30	Tea Break
16:30-16:40	<p>Announcement of 23rd AHWP Annual Meeting Host</p> <p>Mr. Zamane Abdul Rahman, AHWP Vice Chair, Chief Executive, Medical Device Authority, Ministry of Health Malaysia, Malaysia</p>
16:40 – 16:50	Speech by Newly Elected Chair of AHW
16:50-17:00	<p>Closing Remarks by Outgoing Chair/Vice-Chair of AHWP</p> <p>Mr. Zamane Abdul Rahman, AHWP Vice Chair, Chief Executive, Medical Device Authority, Ministry of Health Malaysia, Malaysia</p>
17:30-18:00	<p>The 6th AHWP ASL Annual General Meeting (AGM)</p> <p>ASL Members only</p>



(Transfer to another room)

END OF DAY 5

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