AHWPTC Leaders Meeting Report

AHWP Secretariat Meeting 6-7 May 2013 Kuala Lumpur, Malaysia

AHWPTC





Group Photo



The Last AHWPTC Leaders Meeting was held on 26-27 Feb 2013, Bangkok, Thailand



AHWPTC Leaders Meeting 26-27 Feb 2013, Bangkok, Thailand

Agenda for 26 Feb (day-1)

No.	Item	Name
1	Opening remarks	Joanna Koh
2	Adoption of the Agenda	Joanna Koh
3	Roll Call	Bryan So
4	AHWP Strategic Framework	Bryan So
5	AHWP Member Economies current status	Bryan So
	Tea Break	
6	Objective of AHWP TC Leaders meeting in Bangkok	Joanna Koh
7	SWOT analysis for AHWP TC	Ali M Al-Dalaan
8	Discussion on the role of Consultants in AHWP	Bryan So All participants
	Lunch	
9	AHWP TC's current status Individual WG work plan discussion (i)WG team members; (ii)Work items (previous and new) (iii)WG update since the Chinese Taipei meeting.	All Participants
10	AHWP/RAPS Conference update	Tran Quan



AHWPTC Leaders Meeting 26-27 Feb 2013, Bangkok, Thailand

No.	Item	Name
11	Closed Sessions for TC Leaders and Advisors in own Groups	TC Leaders
		Advisors
12	Discussion on Roles	All Participants
	1. Role of TC and WGs	Lead by Joanna
	-Where are we, where are we going? and what can we achieve	Miang ,Ali &
	2. What we would hope Advisors role to encompass	Bryan
13	Advisors sharing of GHTF experience	Advisors
	1. What are the concrete steps AHWPTC can take to achieve its goals &	
	objectives	
	2. What role do Advisors see themselves playing in the AHWP TC	
	a. Support to WGs work plans	
	b. Support to WGs to move towards alignment with Strategic Framework	
14	Upcoming AHWP meetings:	Bryan So
	(i)AHWP TC meeting (Malaysia)	
	(ii) AHWP main meeting (Malaysia)	
	(iii) AHWP Conference (Malaysia	



AHWP Technical Committee

WG1: Pre-Market Submission and **CSDT**

- BAI Huifen **TAN Ming Hao**
- Alfred KWEK

WG1a: IVDD

- Li-Ling LIU
- Jeffrey CHEN

WG2: Post-Market Surveillance and Vigilance

- Yorkie CHOW Jennifer MAK
- Saini KULWANT

WG3: Quality Management System

- Ali M AL-DALAAN
- Fe Bin LIFW

WG4: Quality System Audit

WG5: Clinical Safety/ Performance

WG6: Capacity **Building** and **Regulatory Training**

STG: Medical Device Nomenclature

YANG Lian Chun

- Abdulah AL Regulator Chair Rama New չկան Chair & WG5 Chair & Getted on 28 Feb 24 հետո
- Negrin MG2 Chair: effected POAT TON MON 2006.}
 - Sumati RANDEO

Jack WONG

Carol YAN



AHWP Technical Committee Advisory Panel

No.	Title / Given Name / Surname	Company
1	Dr Philippe Auclair	Abbott Laboratories
2	Mr Michael Gropp	Medtronic, Inc.
3	Mr Leighton Hansel	Abbott
4	Dr Eamonn Hoxey	1&1
5	Dr Petra Kaars-Wiele	Abbott
6	Mr Greg Leblanc	Cook Canada
7	Dr Peter Linders	Philips Healthcare
8	Mr Benny Ons	BD Europe
9	Mr Grant Ramley	Aseptico Inc
10	Mr Scott Sardeson	3M Health Care

^{*} Surnames in alphabetic order



Summary of discussion 26 Feb 2013

- AHWP strategic framework elements
 - Need for continued education & capacity building
 - Identification & training of future leaders
- AHWP member economies current status
 - More granularity & qualification in regards to status summary for different member economies
- SWOT analysis
 - Challenges presented mirror those faced by GHTF & other harmonization forums
 - Potential issue of lack of industrial commitment for harmonisation
 - Opportunity for AHWP to develop a guideline that member economy can adopt
 - Active participation and commitment to develop guidelines.
 - Different requirements may be explored for lower risk v.s. higher risk MDs
- Discussion on the rule of Consultants in AHWP
 - Consideration for Liaisons to include persons apart from regulator and industry background. (e.g. academic associations, standardisation bodies, patient groups)
 - To define roles and objectives of observers or liasion participation



Summary of discussion 26 Feb 2013

Working Group	Summary of Plan
WG1	 Review revised GHTF definitions in guidance & their impact on AHWP Software standards review & Draft of preliminary guidance document for internal circulation for comments
WG1a	 Development of AHWP guidances on IVD medical devices Training for AHWP member economies (Roadmap)
WG2	 Set up & promote use of electronic AE reporting forms across MEs Developing AE reporting requirements & timelines for all stakeholders Develop guidance document for MD disposal - in context of complaints Formal training of SG2/WG2 guidance document
WG3	 Review QMS guidance documents New guidance document (QMS - local industry, distributor and importer)



Summary of discussion 26 Feb 2013

Working Group	Summary of Plan
WG4	 Finalising guidance documents Training of AHWP guidance documents for MEs Development for auditing guidance for importers & distributors Survey development and distribution among member economies
WG5	 Build consensus within WG to continue framing guidance document based on GHTF SG5, ISO 14155 and ICH GCP – survey to be re-issued Providing inputs to ISO 14155 team on the latest ISO 14155 version for their upcoming meeting in Apr 2013 Comparative study of regulations & related guidance on clinical trial requirements Partnering with other TC work groups' initiatives to provide expertise & input relating to clinical safety/performance Training on GCP, GHTF SG5 GN and latest version of ISO 14155 and Clinical investigation evaluation
WG6	 Prioritise needs of different WGs Explore available training materials that caters for WG needs Map training requirements with other organisations Establish and implement basic online system for different WGs



1. How WG Membership should be determined in the future/selection criteria

- 1.1 Voting among the four options:-
- A. Person moving from member economy to non-member economy
- B. Person moving from one company to another within member economies (WG1, WG1a, WG2, WG3, WG4, WG5, WG6)
- C. Person moving from MD industry to another industry
- D. Appointing a proxy/ representative who is not based in member economy
- 1.2 Conclusion Proposed change to TOR to Secretariat:

A WG member must be based in AHWP member economy

2. Frequency of TC meetings

- 2.1 Continue to have 1 TC leaders (close) meeting plus 1 TC (open) meeting back to back with AHWP main meeting in 2014
- 2.2 Location of TC Leaders 2014 will tentatively be either in India or Hong Kong
- 2.3 TC leaders meeting would only allow the attendance of TC Advisors and WG Chair & Co-Chair
- 2.4 In case of apology, WG Chair and Co-Chair can send representative, while TC Advisors cannot send representative (who is invited in personal capacity)



3. AHWP RAPS 1st Joint Conference

3.1 Suggestions from WG on training topics should be feedback to Secretariat by 31 Mar 2013

4. Updates of Current Regulatory Status of Member Economies

Current Regulatory Status of Member Economies (as at 2012)

S/N	Member Economy	Medical Device Definition	Manufacturer Definition	Classification	Nomenclature	Essential Principles
1	Abu Dhabi					
2	Brunei	No	No	No	No	No
3	Cambodia	Yes	Yes	Yes, GHTF	No	No
4	Chile	Yes	Yes	Yes	No	Yes
5	China	Yes	No	Yes	No	No
6	Chinese Taipei	Yes	Yes	3 risk-class, & GHTF guidelines	TMDN	Yes for Class III only
7	Hong Kong	Yes, GHTF	Yes, GHTF	Yes, GHTF	Yes, AMDN	Yes, GHTF
8	India	No, moving towards GHTF	National definition & moving towards GHTF	No	No	No, moving towards GHTF
9	Indonesia	Yes, GHTF	Yes, GHTF	Yes	No	Yes, GHTF



Current Regulatory Status of Member Economies (as at 2012)

S/N	Member Economy	Medical Device Definition	Manufacturer Definition	Classification	Nomenclature	Essential Principles
10	Jordan					
11	Korea	Yes, national definition based on GHTF	No	Yes	Yes	Yes, KFDA Medical Device Act requirements are equivalent to EPs; include the contents of EPs
12	Laos	Yes	No	Yes	No	No
13	Malaysia	Yes, GHTF	Yes, GHTF	Yes, GHTF	Yes, UMDNS	Yes, GHTF
14	Myanmar					
15	Pakistan					
16	Philippines	Yes	Yes	Yes	No	No



Current Regulatory Status of Member Economies (as at 2012)

S/N	Member Economy	Medical Device Definition	Manufacturer Definition	Classification	Nomenclature	Essential Principles
17	Saudi Arabia	Yes, GHTF	Yes, GHTF	Yes, GHTF	Yes, UMDNS & GMDN	Yes
18	Singapore	Yes, GHTF	Yes	Yes, GHTF	Yes	Yes, GHTF
19	South Africa	Yes	Yes	No	Yes, UMDNS	No
16	Thailand	Yes, GHTF	No, but definition of manufacture in the MD Act	Presently No, but being in the procedure of reclassificatio n	Yes, UMDNS& GMDN	Yes
21	State of Kuwait	Yes	No	No	No	Yes
22	Vietnam					
23	Yemen	YES	Yes	No	No	Yes





4. TC roles and responsibility

AHWPTC as the executive arm of the Working Party should perform the following roles and responsibilities to support the Working Party:

- 1) Execute the Working Party's decisions and resolutions
 - 1.1 WG leaders should report and present their recommendations to TC leaders for final endorsement after their posting of DRAFT/DRAFT PROPOSED document on AHWP website for "call for comment" from all AHWP members
- 2) Make recommendations to the AHWP Chair for decisions;
- 3) Submit resolutions to the AHWP Meetings for decisions of key issues related to the policy, direction, organization, structure and operation of the Working Party;
- 4) Provide expert opinions and advice;
- 5) <u>Develop technical documents and policy papers;</u>





- Plan and organize meetings, training, seminars, workshops and experience sharing sessions;
- 7) Work with related organizations and participate in their activities; and
- 8) Report on the progress of its activities to the AHWP Meetings.
- 9) <u>To revisit the goals and directions of the TC for in line with AHWP's objectives (and the Strategic Framework) in the light of the formation of the TC Advisory Panel in the areas of :-</u>
- 10) The work plan of each Work Group
- 11) To strengthen the TC procedures so as to facilitate TC's overall objectives.
- 12) Strengths & Weaknesses identified in last meeting





- 5. What we would hope Advisors' role to encompass
 - 5.1 To offer advice on work items which are assigned to AHWP by other international /regional org



Sharing from TC Advisors

Recommendations

1. Strategy

- a) Identification of elements of a comprehensive regulatory system needed to be addressed in the AHWP regulatory model (ARM) for IVD and MD
- b) Identification (through gap analysis) of **building blocks** that needed for the purpose in the regional context to develop the ARM (considering the differences among member economies)

2. Three levels of documents which will lead to AHWP's Best Practices

- a) Focusing more on the creation of AHWP unique documents in addition to the adoption of GHTF/IMDRF* ones [Level 1]
- b) Interpretive documents on existing documents lead to capacity building [Level 2]
- c) Collaborative decisions on borderline issues by making AHWP's decision and setting of position on standards (e.g. determine borderline definition/inclusion of certain product as MD) [Level 3]
- * GHTF documents can be regarded as level 1 or 2 doc
- # regulatory model can be regarded as level 0



Sharing from TC Advisors

Recommendations

3. Process

- a) Need more governance and oversight to verify work items fit in the approved ARM
- b) Address recruitment, qualified, balanced, consistent participation on work group
- c) Industrial participation is not limited to multi-national corporations (MNC) especially if topics are directly related to local environment
- d) Need to clarify the meaning of adoption or implementation for AHWP

4. Structure and International alignment

- a) Consider a more flexible working structure not like GHTF
- b) Roles of TC advisors
- c) Determine the AHWP liaison activities
- d) Consider to form a focus group (special task group STG) on standards



Quick Feedback from TC Leaders on recommendations

- 1. Agreed on setting priority work on sharing of "How to set up regulatory framework" for AHWP members.
- 2. AHWP makes recommendation on technical aspect instead of having the power to request jurisdiction to change their regulations to fit the endorsed AHWP GN
- 3. AHWP play book (how to set goal, plan resources, implement strategy, etc) can be useful to help members who are preparing their MD regulation within AHWP ME
- 4. Acknowledged and appreciated the recommendations from TC advisors and will determine action & follow up items