## 第三屆台日醫藥交流會議議程

## **3rd Joint Conference of Taiwan and Japan on Medical Products Regulation (tentative)**

Open Session: 26th November 2015

**Venue: Chang Yung-Fa Foundation International Conference Center** 

08:30-08:50	Registration		
	Opening Remarks		
	<ul> <li>Mr. Jason Chou, Deputy Secretary-general, Association of East Asian Relations, Taiwan</li> </ul>		
	<ul> <li>Mr. Takashi Hamada, Secretary General, Interchange Association, Japan Taipei office, Japan</li> </ul>		
	- Ms. Yu-Mei Chiang, Director-General, Taiwan FDA		
08:50-09:20	- Dr. Toshiyoshi Tominaga, Associate Executive Director for International Programs, PMDA		
	<ul> <li>Mr. Wei-Jen Chen, President, Taiwan Pharmaceutical Manufacturers Association (TPMA)</li> </ul>		
	- Mr. Kenichi Matsumoto, Vice Chairman, JFMDA		
	- Mr. Kiyoshi Horie, Asia Subcommittee, JPMA		
09:20-09:30	Memorial Photo Taking		
	Keynote Speech Current status of medical products regulation in Taiwan and Japan		
09:30-10:10	Moderator: (TBD) - Dr. Shiow-Ing Wu, Deputy Director-General, Taiwan FDA (20mins) - Dr. Kazuhiro Shigetoh, Executive Director, PMDA (20mins)		
10:10-10:30	Coffee Break		
Cooperation	on Accomplishments and Future Prospect for Japan-Taiwan Medical Products		
	Experience sharing on new product development in both Japan and Taiwan		
10:30-11:30	Moderator: Ms. Li-Ling Liu, Director, Division of Medicinal Products, Taiwan FDA		
	<ol> <li>Mr. Kouichi Tsuji, Terumo Corporation (20mins)</li> <li>Prof. Chia-Chi Lin, International Center of Excellence in Cancer Research, Taiwan University (20mins)</li> </ol>		
	Panel discussion (20mins)		
11:30-12:30	How to exchange meaningful GCP information		
	Moderator: Dr. Junko Sato, Office Director, Office of International Cooperation, PMDA		
	<ol> <li>Mr. Ryosuke Sakai, Office of Non-Clinical and Clinical Compliance, PMDA (20mins)</li> </ol>		

	<ol> <li>Ms. Liao, Wei-Cheng, Project Manager, Division of Medical Devices, Center for Drug Evaluation (20mins)</li> <li>Panel discussion (20mins)</li> </ol>
12:30-13:30	Luncheon

Parallel Session						
Pharmaceutical		Medical device				
	Collaboration between Taiwan and Japan for developing self-medication utilizing OTC		Currently regulation amendments and future prospects			
13:30-14:30	Moderator: Mr. Fumihito Takanashi, Deputy Director, Pharmaceutical Safety and Environmental Health Bureau, MHLW 1. Collaboration between Taiwan and Japan for Developing Non- prescription medicine Mr. Arthur Su, Board Member, Taiwan Pharmaceutical Marketing & Management Association (TPMMA) (20mins) 2. Mr. Motohito Nishizawa, Senior Advisor, Japan Self-Medication Industry, JSMI (20mins)	13:30-14:20	<ol> <li>Dr. Yu-Wen, Ruby Huang, Section Chief, Division of Medical Devices and Cosmetics, Taiwan FDA (20mins)</li> <li>Mr. Hideyuki Kondo, Deputy Director, Medical Device and Regenerative Medicine Product Evaluation Division, Pharmaceutical Safety and Environmental Health Bureau, MHLW (20mins)</li> <li>Panel discussion (10mins)</li> </ol>			
	Panel discussion (20mins)	14:20-14:40	Coffee Break			
14:30-15:40	Collaboration between Taiwan and Japan for faster approval for new drugs: "Current progress and next steps"		Learning, assessments and proposal for the collaboration activities in the medical device field			
	<ul> <li>Moderator: Dr. Shih-Chih Tsai, Senior Technical Specialist, TFDA</li> <li>1. Dr. Kanako Takatsuki, Office of International Cooperation, PMDA (15mins)</li> <li>2. Regulation comparison of new drug review between Taiwan and Japan Mr. Po-Yu Wang, Section Chief,</li> </ul>	14:40-15:30	<ol> <li>Mr. Makoto Yokote, JFMDA (15mins)</li> <li>Mr. Karen Ho, Manager, Regulation &amp; Document Management Department, United Orthopedic Corporation (15mins)</li> <li>Panel discussion (20mins)</li> </ol>			
	Taiwan FDA (15mins)	15:30-15:50	Coffee Break			

	3. Mr. Isao Sasaki, Astellas/ JPMA		Progress of WG activities	
	(10mins) 4. Ms. Carol Cheng, Chief Operating Officer, Taiwan Research- based Biopharmaceutical Manufacturers Association (TRPMA) (10mins)  Panel discussion for next step (20mins)	15:50-16:20	<ol> <li>Product Registration WG:         <ul> <li>Dr. Madoka Murakami, Office of International Program,</li> <li>PMDA (10mins)</li> </ul> </li> <li>QSD/QMS WG:         <ul> <li>Ms. Yu-Hsuan Chen, Senior</li> <li>Technical Specialist, Division of Risk Management, Taiwan FDA (10mins)</li> </ul> </li> <li>Panel discussion (10mins)</li> </ol>	
15:40-16:10	Coffee Break	16:20-16:30	` '	
16:10-17:40	Collaboration between Taiwan and Japan for faster access to new drugs under NHI system  Opening remarks for NHI session  1. Ms. Ru-Liang Shih, Medical Review and Pharmaceutical Benefits Division, NHIA (5mins)  2. Mr. Mimei Takahashi, Deputy Director, Economic Affairs Division, Health Policy Bureau, MHLW (5mins)  Presentation "Pricing effectiveness to realize faster access for NHI reimbursement"  1. Mr. Mimei Takahashi, Deputy Director, Economic Affairs Division, Health Policy Bureau, MHLW (20min)  2. Mr. Shang-Ping Chen, Researcher, NHIA (20mins)  Panel discussion inviting key member of PBRS (20mins)	16:30-17:30	Closed Meeting	
	Closing Remarks			
17:40-18:00	- Dr. Shiow-Ing Wu, Deputy Director-General, Taiwan FDA (10mins)			
	- Dr. Kazuhiro Shigetoh, Executive Director, PMDA (10mins)			
18:00	Adjourn			