

US-JAPAN HBD EAST Think Tank Meeting 2023

Date: Thursday, December 14th, 9:30 AM-6:00 PM (JP Time)

Venue: Ariake Central Tower Hall and Conference, Hall A (4F)

Language: English & Japanese (simultaneous interpretation)

Moderator: **MIYASAKA Tomoyuki** (MHLW) & **OHASHI Moe** (PMDA)

Session A : Welcome Speeches

Time	Agenda items	Speakers and Panelists
9:30~9:35	A-1 From MHLW	YOSHIDA Yasunori <i>Deputy Director-General, MHLW</i>
9:35~9:40	A-2 From PMDA	FUJIWARA Yasuhiro <i>Chief Executive, PMDA</i>
9:40~9:45	A-3 From FDA	Jeffrey Shuren <i>Director -CDRH OFFICE, FDA</i>
9:45~9:50	A-4 From JFMDA	TAKAGI Toshiaki <i>Vice Chairman, JFMDA</i>
9:50~9:55	A-5 From AdvaMed	Janet Trunzo <i>Senior Advisor to the President, Senior Executive Vice President, Technology & Regulatory Affairs, AdvaMed</i>

Session B : 20th Anniversary Keynote Speeches

Chair: **Neal Fearnott** (MED Institute Incorporated), **HO Mami** (Yumino Heart Clinic)

Time	Agenda items	Speakers and Panelists
10:00~10:15	B-1 HBD history	Mitchell Krucoff (Duke Univ.)
10:15~10:30	B-2 Achievements of HBD activities and future expectations	SUZUKI Yuka (Clinical Research, Innovation and Education Center, Tohoku Univ. Hospital (CRIETO))
10:30~10:35	B-3 Q & A	



Coffee Break (15 min)

Session C : Learning from HBD activity and recent update

Chair : **Aaron Lottes** (Purdue Univ.), **SAITO Shigeru** (Shonan Kamakura General Hospital)

Time	Agenda items	Speakers and Panelists
10:50~11:00	C-1 Update on HBD activities - Focusing on the last 5 years-	MORIKAWA Hanako (PMDA)
11:00~11:10	C-2 What we can say now based on our experience in obtaining approval in Japan and the U.S. Case 1: Japanese industry's view	SENSHU Kazuhisa (Terumo Corporation)
11:10~11:20	C-3 What we can say now based on our experience in obtaining approval in Japan and the U.S. Case 2: U.S. industry's view	YASUHARA Daiki (Medtronic Japan)
11:20~11:30	C-4 Role of Academia in HBD Activities	YOKOI Hiroyoshi (Fukuoka sanno Hospital)
11:30~11:35	C-5 Q & A	

Session D : Evaluating the efficacy and safety of medical devices from pre-market through post-market using RWD

Chair : **Misti Malone** (FDA), **ISHII Kensuke** (PMDA)

Time	Agenda items	Speakers and Panelists
11:40~11:50	D-1 Basic Approach in utilizing RWD for regulatory decision-making	Misti Malone (FDA)
11:50~12:00	D-2 Challenges in establishing RWE for pre- and post-market clinical evaluation	NAKAMURA Masato (Toho Univ.)
12:10~12:20	D-3 Challenges in developing devices using RWD in Japan	KAWAHARA Kazuo (Boston Scientific Japan)
12:20~12:40	D-4 Panel Discussion <i>Theme: The efficient way of collecting RWD for regulatory decision-making in pre- and post-market to accelerate device development</i>	Speakers & Eric Chen (Abbott Medical) IWAISHI Chie (Edwards Lifesciences) Aaron Lottes (Purdue Univ.) SHIBA Takeshi (PMDA)



Lunch Break (60 min)

Session E : Approaches of HBD activity to promote the development of SaMD

Chair : **Eric Chen**(Abbott Medical), **OKAZAKI Yuzuru** (PMDA)

Time	Agenda items	Speakers and Panelists
13:40~13:50	E-1 Regulation of SaMD in the U.S.	Nicole Ibrahim (FDA)
13:50~14:00	E-2 Regulation of SaMD in Japan	KATO Kentaro (PMDA)
14:00~14:10	E-3 Learning from "CureApp" :how to develop and get an approval of SaMD	TANIGAWA Tomoyuki (CureApp)
14:10~14:20	E-4 Points to consider in the application of AI for medical devices	HAMAMOTO Ryuji (National Cancer Center Research Institute)
14:20~14:40	E-5 Panel Discussion <i>Theme: Strategies to promote the development of SaMD from the standpoints of industry, government, and academia</i>	Speakers & IKENO Fumiaki (Stanford Univ.)

Session F : Approaches of HBD activity to promote the development of pediatric devices

Chair : **Nicole Gillette** (FDA), **YASUKOCHI Satoshi** (Aizawa hospital)

Time	Agenda items	Speakers and Panelists
14:45~14:55	F-1 Progress and challenges in pediatric medical device development	FUJII Takanari (Showa Univ. Hospital)
14:55~15:05	F-2 U.S. Regulatory initiatives to promote pediatric medical device development	Nicole Gillette (FDA)
15:05~15:15	F-3 The road from development to approval of pediatric medical devices and future approaches.	NEMOTO Shintaro (Osaka Med. Pharm. Univ.)
15:15~15:25	F-4 Utilization of RWD in pediatric medical device development	INUZUKA Ryo (Tokyo Univ.)
15:25~15:55	F-5 Panel Discussion <i>Theme: Strategies to promote the development of pediatric medical devices from the standpoints of industry, government, and academia</i>	Speakers & Sung-Hae Kim (Shizuoka Children's Hospital) AIZAWA Koichi (PMDA) Eric Vang (Medtronic Cardiac Surgery)



Coffee Break (15 min)

Session G : What should be considered for global harmonization of medical device development through HBD activity ?

Chair : **Mitchell Krucoff** (Duke Univ.), **YABANA Naoyuki** (PMDA)

Time	Agenda items	Speakers and Panelists
16:10~16:20	G-1 An overview of the global situation surrounding medical devices	IKENO Fumiaki (Stanford Univ.)
16:20~16:30	G-2 Current situation of medical device regulations outside of Japan and the U.S.	Katharine Stohlman (Corvia Medical)
16:30~16:40	G-3 Comparing clinical practices or consultation processes in the US vs Japan	Robert Thatcher (Diaxamed, LLC)
16:40~16:50	G-4 Unique points of medical device development and advantages of global development.	IKEDA Koji (CRIETO)
16:50~17:00	G-5 Post-approval hurdles: Differences and strategies between Japanese and the U.S. insurance systems	TAMURA Makoto (Healthcare System Planning Institute (HSPI))
17:00~17:55	G-6 Panel discussion <i>Theme: Future direction of HBD activity</i>	Speakers & Nicole Gillette (FDA), NAKAI Kiyohito (MHLW) Janet Trunzo (AdvaMed), TANAKA Shiho (JFMDA, J&J) IWAMOTO Shin (MHLW)

Session H : Closing remarks

NAKAYAMA Tomonori (MHLW)



Anyone can participate in the reception party (18:30-20:00).

Venue: Ariake Central Tower Hall and Conference (3F)

Please tell us your impressions of participating in HBD East Think Tank 2023. This survey will take approximately 3 minutes to complete. Thank you in advance for your participation.

