Appendix 1

AGENDA

Date	Tiem	Agenda	Speaker (TBD)
11/23/2024	All day	Registration	
11/24/2024	8:00-8:30	Sign in	
Topic 1: GHWP Promotion of Global Medical Device Regulatory Convergence, Coordination, and Trust			
11/24/2024	8:30-9:00	Opening Remarks	GHWP Leadership
	9:00-10:00	Harmonization, Coordination, and Trust in Global Medical Device Regulation—GHWP Strategic Framework Towards 2026	Mr. Guobiao Gao GHWP SAB Member
	10:00-10:30	Promoting Industrial Innovation through Medical Devices Regulatory Science	Prof. Yingjun Wang GHWP-GZA President
	10:30-11:00	Group Photo, Tea break	
	11:00-11:30	Trends in Innovation for High-end Medical Equipment Industry (industry/tech)	Dr. Milind Sabnis Head of Advisory, Healthcare, Frost & Sullivan
	11:30-12:00	Hot Topics and Development Trends in Global Regulation of High-end Medical Equipment	Miang Tanakasemsub Head of Regulatory Affairs, Asia Pacific, Johnson & Johnson Vision
	12:00-14:00	Lunch	
Topic 2: Regulation Requirements and Common Issues for Active Medical Devices in China			
11/24/2024	14:00-14:40	Inovative Research on Clinical Trial	Center For Medical

		Pathways for Medical Devices	Device Evaluation, NMPA
	14:40-15:20	Encouraging the Innovation and Development of Medical Devices — Progress in the Technical Review of Medical Devices	Jun Li； Deputy Director； Center For Medical Device Evaluation, NMPA ； Vice President, Technical Committee, GHWP
	15:20-15:50	Tea break	
	15:50-16:30	Pre-marke Regulation of Active Medical Devices	Liang Peng, Deputy Director, The 1st Evaluate division, Center For Medical Device Evaluation, NMPA
	16:30-18:00	Seminar—Regulatory Innovation for Innovative Medical Devices and Clinical Trial Risk Identification	Department of Medical Device Registration, NMPA; Department of Medical Device Regulation, NMPA; Center For Medical Device Evaluation, NMPA
Topic 3: Innovation and Regulation in the Medical Robotics Industry			
11/25/2024	9:00-9:30	Current Status and Trends of the Medical	Prof. Lining Sun,

		Robotics Industry in China	School of Mechanical and Electrical Engineering, Univ. of Suzhou
	9:30-10:00	Current Status and Trends of Medical Robotics Regulation in China	Pengfei Yang, Director, The 2nd Evaluate division, Center For Medical Device Evaluation, NMPA
	10:00-10:40	Enriching Research and Development in Medical Robotics for Clinical Translation	Prof. Bryan So, GHWP Secretary-General, Chinese University of Hong Kong
	10:40-11:20	Current Status and Trends of the Medical Robotics Industry in Global	Emily Li, Sr Regulatory Affairs Manager, Medtronic Greater China
	11:20-12:00	Introduction to New Registration Regulations for Active Medical Devices in Saudi Arabia	Mr. Hanadi Alousaimi, Senior Scientific Evaluation Expert, SFDA
	12:00-13:30	Lunch	
Topic 4: Innovation and Regulation in the High-end Medical Imaging and Therapeutic Equipment Industry			
11/25/2024	14:00-14:30	High-end Medical Imaging Diagnosis and Treatment Equipment and Technology	Mr. Yee Huang Vice CEO, Siemens China

	14:30-15:00	Medical Imaging Equipment Registration in APAC Countries	Alfred Kwek GHWP SAB Member
	15:00-15:40	Medical Gas System (MGS) – Essential Principles of Safety and Performance (EPSP) – Standards for Demonstrating Compliance	Al-Khairi Mohd Daud GHWP TC WG8
	15:40-16:20	Introduction to New Regulatory Guidelines for Active Medical Devices Registration in Vietnam	Doan Quang Minh DAV
	16:20-17:00	The Road to Innovation and Internationalization of Surgical Robots	Dr. He Chao, the founder, executive director, and CEO of Shanghai MicroPort Medical (Group) Co., Ltd.
	17:00-18:00	Panel Discussion	Speakers from Topic 3 and 4, Representatives from Regulatory Authorities of Various Countries/Regions
Topic 5: Innovation and Regulation in the Artificial Intelligence Medical Device Industry			
11/26/2024	8:30-9:00	Current Status and Trends of the Artificial Intelligence Medical Device Industry	Jyh-Ching Yaur Vice CEO, Philips Greater China
	9:00-9:30	Frontline Developments in AI Medical Devices and Digital Healthcare	Prof. Xiangmin Xu, Vice President of South China University of Technology
	9:30-10:10	Standardization and Regulatory Issues of AI/ML Based Medical Devices in Korea	Prof. Hwiyoung Kim

			Yonsei University , Korea
	10:10-10:50	Introduction to Active Medical Device Regulation in Thailand	Sirinmas Katchamart FDA THAI
	10:50-11:50	Panel Discussion	Speakers from Topic 5, Representatives from Regulatory Authorities of Various Countries/Regions
	11:50-12:30	Graduation Ceremony	
On-site Training			
11/26/2024	14:00-18:00	On-site Training	
2024/11/27	All Day	Participants Depart	