Appendix 1

Agenda of The 1st GHWP (Guangzhou) Academy Training

Date	Time	Contents	Speakers	Participants	Venue
2024/5/26	All day	Registration			Hotel
2024/5/27	8:30-9:00	Sign in			Lecture Hall B13
	Topic 1: GHWP Promotes Global Medical Device Regulatory Convergence, Harmonization, and Reliance				
	9:00-9:30	Opening speech	GHWP Chair, Deputy Commissioner of NMPA Jinghe Xu Chair of GHWP (Guangzhou) Academy, Vice President of SCUT Yong Xu	All participants	Lecture Hall B13
	9:30-10:00	Group photo			
	10:00-10:30	Global Medical Device Innovation and Development Trends	President of GHWP (Guangzhou) Academy, Professor of SCUT Yingjun Wang	All participants	Lecture Hall B13
	10:30-11:00	Driving Convergence, Harmonization and Reliance of Global Medical Devices Regulatory: an overview of GHWP	GHWP SAB Member, Party Secretary of Center for Medical Device Evaluation, NMPA Guobiao Gao		

	11:00-11:30	GHWP Strategic Framework Towards 2026	GHWP SAB Member, Baxter Vice President of Quality and Regulatory Affairs Tran Quan		
	11:30-13:30	Lunch			
	13:30-14:30	Introduction to Chinese Medical Device Registration Regulations	Director of Medical Device Registration Department of NMPA Ling Lu	All participants	Lecture Hall B13
	14:30-15:30	Introduction to the Reform of Chinese Medical Device Evaluation System	Director of Center for Medical Device Evaluation, NMPA Lei Sun	All participants	Lecture Hall B13
	15:30-16:00	Tea break			
	16:00-17:00	Q&A on Chinese Medical Device Regulatory Policies and Evaluation	Speakers of Medical Device Registration Department and Center for Medical Device Evaluation, NMPA	All participants	Lecture Hall B13
	18:00-20:00	Collaborative Empowerment—Forum on Innovative Medical Devices Embracing the World and GHWP (Guangzhou) Academy 1st Council Meeting	NMPA	Council members	B12-504
2024/5/28	**Topic 2: Innovative Trends Shaping the Future of Medical Device Industry**				
	8:30-9:30	Global Regulation Reliance and Atlas of Medical Devices	Unit Head, Regulation and Safety, WHO Agnes Sitta Kijo	All participants	Lecture Hall B13

	9:30-10:00	Q&A on WHO	Unit Head, Regulation and Safety, WHO Agnes Sitta Kijo	All participants	Lecture Hall B13
	10:00-10:20	Tea Break			
	10:20-11:20	Development Status of the Asia-Pacific MedTech Industry	APACMed CEO Harjit Gill	All participants	
	11:20-11:50	Q&A on APAC Med	APACMed CEO Harjit Gill	All participants	Lecture Hall B13
	11:50-13:30				
	13:30-14:30	Development Status of the MedTech in American Region	Executive Secretary of IACRC Sandra Ligia González	All participants	Lecture Hall B13
	14:30-15:00	Q&A on IACRC	Executive Secretary of IACRC Sandra Ligia González	All participants	Lecture Hall B13
	15:00-15:20	Tea break			
	15:20-16:20	Development Status of Middle East and North Africa Medical Device Industry	Medical devices, Imaging, and Diagnostics Trade Association (MECOMED) Rana Chalhoub	All participants	Lecture Hall B13
	16:20-16:50	Q&A on MECOMED	Medical devices, Imaging, and Diagnostics Trade Association (MECOMED) Rana Chalhoub	All participants	Lecture Hall B13
	16:50-17:00	Set-up lecture hall			
	17:00-18:00	Panel Discussion 1- Medical Device Products Benefit the	Representatives of WHO, APACMed, IACRC, MECOMED, and GHWP Leadership\other GHWP	All participants	Lecture Hall B13

		People Around the World	members\Representatives of Targeted invited		
2024/5/29	**Topic 3： A Series of Training on the Premarket Regulatory Framework of Medical Device in the World**				
	8:30-10:00	Introduction to the EU Medical Device Registration Policy - MDR and IVDR	Former member of the European Commission, Former Head of the Czech Ministry of Health Erik Hansson	All participants	Lecture Hall B13
	10:00-10:20	Q&A on Pre-Market Approval Policies in EU	Former member of the European Commission, Former Head of the Czech Ministry of Health Erik Hansson	All participants	Lecture Hall B13
	10:20-10:40	Tea break			
	10:40-12:10	Korean Medical Device Registration Review Process and Requirements	Head of the International Affairs Team of the Korean Medical Device Industry Association Sunny Woo	All participants	Lecture Hall B13
	12:10-12:30	Q&A on Pre-Market Approval Policies in Korean	Head of the International Affairs Team of the Korean Medical Device Industry Association Sunny Woo	All participants	Lecture Hall B13
	12:30-13:30	Lunch			
	13:30-15:00	Saudi Arabia Medical Device Registration Review Process and Requirements	Standards & Regulation Expert, Saudi Food and Drug Authority Fajer Alkusair	All participants	Lecture Hall B13

	15:00-15:20	Q&A on Pre-Market Approval Policies in Saudi Arabia	Standards & Regulation Expert, Saudi Food and Drug Authority Fajer Alkusair	All participants	Lecture Hall B13
	15:20-15:40	Tea break			
	15:40-17:40	Panel Discussion 2- Innovative Medical Device Policy	Representatives of EU, Korea, Saudi Arabia, and GHWP Leadership\other GHWP members\Representatives of Targeted invited	All participants	Lecture Hall B13
2024/5/30	8:30-10:00	Indonesian Medical Device Registration Review Process and Requirements	Indonesian Ministry of Health Anita Nu	All participants	Lecture Hall B13
	10:00-10:20	Q&A on Pre-Market Approval Policies in Indonesia	Indonesian Ministry of Health Anita Nur	All participants	Lecture Hall B13
	10:20-10:40	Tea break			
	10:40-12:10	Malaysian Medical Device Registration Review Process and Requirements	Assistant Director of Pre Listing Registry, Medical Devices Bureau, Ministry of Health Malaysia Vishyallenni Sathasivem	All participants	Lecture Hall B13
	12:10-12:30	Q&A on Pre-Market Approval Policies in Malaysia	Assistant Director of Pre Listing Registry, Medical Devices Bureau, Ministry of Health Malaysia Vishyallenni Sathasivem	All participants	Lecture Hall B13
	12:30-13:30	Lunch			

	13:30-15:30	Panel Discussion 3-ASEAN Countries Regulatory Reliance Path	Representatives of Indonesia, Malaysia, China, and GHWP Leadership\other GHWP Members\Representatives of Targeted invited	All participants	Lecture Hall B13
	15:30-15:50	Tea break			
	15:50-17:20	Interactive Communication 1-Introduction to Special Task Group of Common Evaluation Reliance	Acting Chair of GHWP CERP, Director of Quality Management Department of CMDE Shiqing Zhang	Members of CERP, all participants	TBD.
		Interactive Communication 2-Interaction between GHWP member representatives and trainees	Representatives of GHWP Member: Kuwait, Cuba, Jordan, Oman, Hong Kong China	All participants	TBD.
		Interactive Communication 3-Introduction to the Annual Report on the Development of Medical Device Industry in China 2023	National Medical Products Administration Institute of Medical Economics	All participants	TBD.
2024/5/31	**Topic 4: Onsite Practical Training of Innovative Medical Device Enterprises in Guangdong**				
	8:30-12:00	Onsite Practical Training of Innovative Medical Device	Wondfo Biotech (Guangzhou)	All participants	TBD.

		Enterprises in Guangdong			
	12:00-13:30	Lunch			
	Topic 5：Case Sharing: Going Global with Innovative Medical Devices				
	13:30-14:00	Showcase of Advanced Implantable and Interventional Medical Devices	Medtronic	All participants	Lecture Hall B13
	14:00-14:30	Showcase of large scale medical equipment	United-Imaging		
	14:30-15:00	Practice of Cross border Mergers and Acquisitions of Chinese Medical Device Enterprises	Blue Sail Medical		
	15:00-15:20	Tea break			
	15:20-15:50	Construction of Overseas Platforms for Chinese Medical Device Enterprises	Mindray Medical	All participants	Lecture Hall B13
	15:50-16:20	Showcase of In vitro Diagnostic Products	Abbott		
	16:20-16:40	Completion Ceremony			Lecture Hall B13
2024/6/1	**Departure**				

Note: This program will be adjusted according to the preparation of the training