16-Sep-26				
Time	**DAY1**	**Title**	**Lecturer**	**Moderator**
9:20-9:30	Opening	Opening remarks	Mr.Eiji Watanabe（PDA Japan Chapter）	
Session A: Development of Dosage Forms and Formulations Corresponding to Modalities				
9:30-10:10	A-1	Next-generation prefilled syringes that achieve higher volume, higher viscosity, and co-administration formulations	Dr.Mitsuru Takahashi （Nippon Becton Dickinson）	
10:10-10:50	A-2	Optimizing injectability of high-viscosity biologics in next-generation syringe-based autoinjectors	Mr.Norbert Jung（Ypsomed） Mr. Alessandro Morandotti （Stevanato Group）	
10:50-11:05	**Exhibitors' presentations**		**Half of exhibitors**	
11:15-11:45	**Exhibition break**			
11:45-12:25	A-3	Designing Drug Delivery Systems for Context:Case Studies in Lyophilized Drug Delivery and Microliter-Scale Dose Delivery	Mr. Markus Bauss（SHL Medical）＆Mr. Nils Ronquist（SHL Medical）	
12:30–13:10	**Lunch （luncheon seminar）**			
13:10–13:40	**Exhibition break**			
Session B Educational Lecture （The Global Landscape of the Healthcare Environment and the Diversifying Modalities）				
13:40-14:40	B-1	（From a broad perspective on drug modalities）	Mr.Norikazu EIKI（Unknowm）	
Session C: Development of Dosage Forms and Formulations Corresponding to Modalities-2				
14:40-15:20	C-1	Design Considerations for Next Generation Delivery of Oligonucleotide Therapeutics - Focus on High Concentration SC Formulations and Direct CNS Administration （IT admin.） -	Ms.Fumiko Koizumi, Dr. Tomoya Takenaka（Takeda Pharmaceutical Co.ltd.）	
15:20-15:35	**Exhibitors' presentations**		**Half of exhibitors**	
15:35-16:05	**Exhibition break**			
16:05-16:45	C-2	Development of a Subcutaneous Drug Product for Lecanemab	Ms.Naomi Sakuramoto（Eisai Co., Ltd ）	
16:45-17:25	C-3	Formulation–Device Integration in Inhaled Drug Delivery: Optimizing Delivery for Respiratory Medicines	Dr.Mireia Puig-Sellart（Phillips Medisize）	
Session D: Container Materials and Regulatory Compliance				
17:25-18:05	D-1	Elastomer Components and the Latest Regulatory Expectations: Meeting Global Standards（Pre-	Dr.Ana Kuschel（West Pharmaceutical Services）	Ms. Brigitte Reutter-Haerle （Vetter Pharma ）
18:15-20:15	**Mixer**			

17-Sep-26				
Time	**DAY2**	**Title**	**Lecturer**	**Moderator**
Session E: Usability and Risk Management				
9:00-9:40	E-1	On the Medical Device Usability Standard （JIS T 62366-1）	Mr.Hidemasa SEKIMIZU（PMDA）	
9:40-10:20	E-2	Human Factors/Usability Engineering for Combination Products from a vendor’s perspective	Dr.Ong Weiling（INTAGE Healthcare Inc.）	
10:20-10:50	**Exhibition break**			
10:50-11:30	E-3	Risk Management in Drug–Device Combination Products in Clinical Setting	Mr.Yasunori Nagano（Sanraku Hospital）	
Session F: Utilizing IoT （Electronic Information）				
11:30-12:10	F-1	Smart Self-Injection: Enhancing Clinical Trial Efficiency and Commercial Care Coordination	Dr.Stefanie Seiler（YPSOMED）	
12:15–12:55	**Lunch （luncheon seminar）**			
12:55-13:25	**Exhibition break**			
13:25-14:05	F-2	Toward the Implementation of Safe Medication Administration	Ms.Eriko Nemoto（Terumo）	
SessionG: Process Design Methods				
14:05-14:45	G-1	From Clinical Manufacturing to Commercial Supply – Key Considerations and Lessons Learned	Dr.Armin Wilfinger（Vetter Pharma）	
14:45-15:15	**Exhibition break**			
15:15-15:55	G-2	Approaches to yield improvement in redundant filtration system	Mr.Taichi Hanada（Merck）	
SessionHG: Case Studies on Material Properties and Container Closer Integrity Testing				
15:55–16:35	H-1	Considerations for the Use of COC Cartridges in Combination Products for Emerging Drug Modalities	Mr.Christoph Zauner（Schott Pharma）	
16:35-17:15	H-2	CCIT Strategies and Solutions for Combination Products	Dr.Wenliang Chen（Packaging Technologies & Inspection）	
Session I: Q&A session				
17:20-17:50	I-1	Q&A session	Mr.Eiji Watanabe（PDA Japan Chapter）	
17:50–18:00	閉会	Closing message	Mr.Eiji Watanabe（PDA Japan Chapter）	