As of 6 February, 2020

Venue	Conference building				Exhibition building	
	Main Hall	Tachibana Conference Hall	Hagi Conference Hall	Shirakashi Conference Room	Meeting Room 1&2	Exhibition Hall
2/18AM 10:00-12:30	**Opening 10:00〜10:30** **K-01 10:30〜11:30 Keynote1** AI-driven pharmaceutical innovation in Japan **K-02 11:30〜12:30 Keynote2** Medical response to the Great East Japan Earthquake in the Ishinomaki Medical Zone					**Exhibition Poster 9:00〜17:00**
Lunch					**Poster Short Oral 12:45〜13:55**	
2/18PM 14:15-18:05	**C-01 14:15~15:30** JSQA-JMACCT Considerations for Using IT Systems in Clinical Trials Break **X-02 16:00~18:05** Compliance and quality assurance in the integrity of the utilized electrical data in Pharmaceutical industry	**X-01 14:15~15:55** Up-to-date initiatives for promoting clinical trials and Quality Assurance in Asian countries. Break **L-01 16:25~17:55** The history behind the FDA GLP	**L-02 14:15~15:35** Points to be noted when outsourcing non-clinical studies (GLP studies, non-GLP studies) to domestic facilities. Break **L-03 16:05~18:00** Points to be noted when outsourcing non-clinical studies (non-GLP, PK and Pharmacology studies) to overseas facilities.		**Oral Presentations** **O-01** 14:15~14:45 **O-02** 14:45~15:15 **O-03** 15:15~15:45 Break **Oral Presentations** **O-04** 16:15~16:45 **O-05** 16:45~17:15 **O-06** 17:15~17:45	
2/19AM 9:00-12:00	**L-04 9:00~10:15** Promoting quality in research in the academic environment Break **C-02-1 10:45~12:00** Different or Same? GCP Requirements among Regulatory Authorities -Subject Protection and Informed Consent Process -	**L-05 9:00~10:15 / 10:45~12:00** (10:15~10:45 Break) Future of Electronic Archiving	**X-03 9:10~10:15 / 10:45~11:50** (10:15~10:45 Break 30m) Current status and issues of clinical analysis-related regulations in each country	**X-04 9:00~10:15** The investigational medicinal products GMP in Japan (J-GMP for IMP) Break **X-05 10:45~12:00** Embark on the New Tide! PV QMS culture in Japan and Asia	**Oral Presentations** **O-07** 9:15~9:45 **O-08** 9:45~10:15 Break **Oral Presentations** **O-09** 10:45~11:15 **O-10** 11:15~11:45	**Exhibition Poster 9:00〜17:00** **Poster discussion 12:10〜13:50**
Lunch						
2/19PM 14:00-18:00	**C-02-2 14:00~16:00** Different or Same? GCP Requirements among Regulatory Authorities -GCP Inspections - Break **C-02-3 16:30~18:00** Different or Same? GCP Requirements among Regulatory Authorities - Data Integrity in Clinical Trials -	**L-06 14:00~15:55 / 16:25~17:25** (15:55~16:25 Break) What has changed by issuing OECD Advisory Document No.19?	**X-06 14:30~16:00/ 16:30~17:50** (16:00~16:30 Break) Current status of regenerative therapeutic products and Quality Assurance	**L-07 14:00~16:00** Role of QAU at the testing facility in the event of a disaster Break **L-08 16:30~18:00** “SAMURAI” and “Quality Assurance”		
2/20AM 9:00-12:40	**X-07 9:00~9:45** Basics in keeping Quality Break **X-08 10:10~12:10** GxP Regulatory Session					**Exhibition Poster 9:00〜10:30**
	Closing 12:10〜12:40					