SUNDAY, MAY 24, 2015 | PRECONFERENCE WORKSHOPS

	Workshop 1 [Half Day]	Workshop 2 [Full Day]	Workshop 3 [Full Day]
08:30 -12:00		Non-CRF Data (eData) is Equivalent Important as CRF Data in Clinical Trials	How to Complete a CDISC-Compliant CRF Annotation
13:30 - 17:30	Essential Biostatistics Concepts Made Easy		

MONDAY, MAY 25, 2015 | CONFERENCE DAY 1

13:30 - 17:30	Opening Plenary Session + Special Forum
17:30 - 19:00	Networking Reception

TUESDAY, MAY 26, 2015 | CONFERENCE DAY 2

	Theme 1	Theme 2 / Theme 12	Theme 3
	Understanding China Regulatory Landscape	**Theme 2: CFDA Town Hall Theme 12: Hot Topics and Late Breaker**	**Clinical Science and Clinical Trial Operation**
Session 1 08:30 - 10:00	**Session 0101** Successful Stories of Bringing Innovative Product into China Market	**Session 1201** How to Create Efficient Communications for Stakeholders of Drug R&D Chain	**Session 0301** The Responsbilities to Speed Up Clinical Trials
10:00 - 10:30	TEA BREAK		
Session 2 10:30 - 12:00	**Session 0102** ICH E17 MRCT Guidance Development Update	**Session 1202** Expedited Development, Access and Approval Pathways	**Session 0302** The Feasibilty of Initiating Clinical Trials by Single IRB Approving in China
12:00 - 13:30	LUNCH		
Session 3 13:30 - 15:00	**Session 0203** China Food and Drug Administration (CFDA) Town Hall (Part I)		
15:00 - 15:30	TEA BREAK		
Session 4 15:30 - 17:30	**Session 0204** China Food and Drug Administration (CFDA) Town Hall (Part II)		

WEDNESDAY, MAY 27, 2015 | CONFERENCE DAY 3

	Theme 1	Theme 12	Theme 3
	Understanding China Regulatory Landscape	**Hot Topics and Late Breaker**	**Clinical Science and Clinical Trial Operation**
Session 5 08:30 - 10:00	**Session 0105** How Transparency Evolve in Key Authority and How It Help Industry R&D Capability Build Up	**Session 1205** Medical Affairs - How to Deliver the Latest Medical Knowledge to the Health Professional	**Session 0305** Cinical Science (Part I) - Development Strategy
10:00 - 10:30	TEA BREAK / ANNOUNCEMENT OF POSTER PRIZE WINNERS		
Session 6 10:30 - 12:00	**Session 0106** How Multiple-Disciplinary Department in Authority Works Together Collaboratively to Manage/Regulate Product Quality	**Session 1206** On the Way - China Innovative Pharmaceutical Company	**Session 0306** Clinical Operation - Risk Based Monitoring
12:00 - 13:30	LUNCH		
Session 7 13:30 - 15:00	**Session 0107** How Far Away is the Innovative Medicine to China Patients	**Session 1207** Drug-Diagnostic Co-Development and Regulatory Requirement and Challenges for Companion Diagnostic Test	**Session 0307** Cinical Science (Part II) - IMCT Latest Guidance Population PK Value in Development
15:00 - 15:30	TEA BREAK		
Session 8 15:30 - 17:30			**Session 0308** Project Management - Study Wide

SUNDAY, MAY 24, 2015 | PRECONFERENCE WORKSHOPS

Workshop 4 [Full Day]	Workshop 5 [Half Day]		
Exploring Medical Communications Services to Support Patient Care Decisions	Lean: Innovative Approaches for Authoring Clinical Regulatory Documents		

THE 7TH DIA CHINA ANNUAL MEETING AT-A-GLANCE

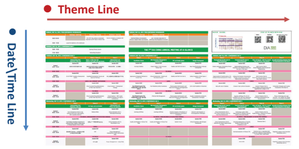
TUESDAY, MAY 26, 2015 | CONFERENCE DAY 2

	Theme 4	Theme 5	Theme 6	Theme 7
	Quantitative Science	**Risk Assessment, Management & Communication**	**CMC and Quality System**	**Biologic Development**
Session 1 08:30 - 10:00	**Session 0401** Oncology Statistics	**Session 0501** Patient Safety in First-In-Human Clinical Trial	**Session 0601** Managing Post-Approval CMC Changes in a Global Market	**Session 0701** New Trend/Technology in Biologic Development
10:00 - 10:30	TEA BREAK			
Session 2 10:30 - 12:00	**Session 0402** Application of Bayesian Methods for Clinical Trials	**Session 0502** Clinical Development - Benefit and Risk Based Decision	**Session 0602** CHP Townhall	**Session 0702** Safety Evaluation of New Types of Antibodies
12:00 - 13:30	LUNCH			
Session 3 13:30 - 15:00		**Session 0503** Post Marketing Safety Surveillance	**Session 0603** Multiple Sites for Biologicals in China	**Session 0703** Development of Biosimilars in China (Part I): Interpretation of Biosimilar Guidance
15:00 - 15:30	TEA BREAK			
Session 4 15:30 - 17:30		**Session 0504** Effective Risk Management	**Session 0604** Effective Management of Contract Manufacturing Organization	**Session 0704** Development of Biosimilars in China (Part II): Opportunities and Challenges

WEDNESDAY, MAY 27, 2015 | CONFERENCE DAY 3

	Theme 4	Theme 5		Theme 7
	Quantitative Science	**Risk Assessment, Management & Communication**		**Biologic Development**
Session 5 08:30 - 10:00	**Session 0405** Involving CDM Role	**Session 0505** Vaccine Safety		**Session 0705** Detection and Evaluation of Immunogenicity of Biologics
10:00 - 10:30	TEA BREAK / ANNOUNCEMENT OF POSTER PRIZE WINNERS			
Session 6 10:30 - 12:00	**Session 0406** Quality and Integrity of Clinical Trial Data	**Session 0506** Herbs Safety		
12:00 - 13:30	LUNCH			
Session 7 13:30 - 15:00	**Session 0407** Stat/DM Joint Session - The Collaboration of DM and Statistics (Part I)			
15:00 - 15:30	TEA BREAK			
Session 8 15:30 - 17:30	**Session 0408** Stat/DM Joint Session - The Collaboration of DM and Statistics (Part II)			

QUICK GUIDE:



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TUESDAY, MAY 26, 2015 | CONFERENCE DAY 2

	Theme 8	Theme 9	Theme 10	Theme 11
	Therapeutically Specific Issues in Drug Development	**Functional Specific Issues in Durg Development**	**China and Beyond - WHO PQ, Generic Drug and Botanic Drug Pathway**	**Innovative Partnership and Business Model for Drug Development in China Beyond**
Session 1 08:30 - 10:00	**Session 0801** Global Landscape of Innovative Clinical Development	**Session 0901** Medical Writing Evolution and Expansion: from Global to China, from the Past to the Future	**Session 1001** WHO Vaccine (Part I): Global Regulatory and Procurement Requirements	
10:00 - 10:30	TEA BREAK			
Session 2 10:30 - 12:00	**Session 0802** Opportunities & Transformation of TA Related Clinical Development in China	**Session 0902** Embrace the Dynamic Environmental Change: Innovations Lead to New Opportunities and Solutions	**Session 1002** WHO Vaccine (Part II): Newer than New-Innovations to Create New Health Solutions	
12:00 - 13:30	LUNCH			
Session 3 13:30 - 15:00	**Session 0803** High and Lows in Diabetes	**Session 0903** Why is Professional Medical Writer Needed in Boosting Scientific Publication in China	**Session 1003** Botanical (Part I): Developing New Drugs from Traditional Chinese Medicine for Unmet Public Health Needs	**Session 1103** Partnership Strategies and Creative Frameworks in the New Era
15:00 - 15:30	TEA BREAK			
Session 4 15:30 - 17:30		**Session 0904** Regulatory Writing: CTD, ICH E3 - Compliant CSR, and China Submission Dossier Preparation (Clinical Parts)	**Session 1004** Botanical (Part II): Developing New Drugs from Traditional Chinese Medicine for Unmet Public Health Needs	**Session 1104** Money, Technology and Legal Structure - Put Together the Jigsaw Puzzle for an Innovative Partnership

WEDNESDAY, MAY 27, 2015 | CONFERENCE DAY 3

	Theme 8	Theme 9	Theme 10	Theme 11
	Therapeutically Specific Issues in Drug Development	**Functional Specific Issues in Durg Development**	**China and Beyond - WHO PQ, Generic Drug and Botanic Drug Pathway**	**Innovative Partnership and Business Model for Drug Development in China Beyond**
Session 5 08:30 - 10:00	**Session 0805** Clinical Development (Part I) - The Critical Successful Factor for New Drug Development	**Session 0905** Technology Leads to New Opportunities for Meeting Customer Medical Information Needs	**Session 1005** Generic Drug Forum (Part I) - Global Requirements and Regulatory Initiatives	
10:00 - 10:30	TEA BREAK / ANNOUNCEMENT OF POSTER PRIZE WINNERS			
Session 6 10:30 - 12:00	**Session 0806** Clinical Development (Part II) - What Are the Gaps & Opportunities for China	**Session 0906** Innovative Approaches to Ensuring Quality in Clinical Trials and Compliance to GCP	**Session 1006** Generic Drug Forum (Part II) - Product Development for Quality Consistency	
12:00 - 13:30	LUNCH			
Session 7 13:30 - 15:00		**Session 0907** Internationalization of Chinese CRO	**Session 1007** Generic Drug Forum (Part III) - Lessons Learned from Quality and Compliance Inspections	**Session 1107** Regulatory Aspect of the Innovative Partnership (Part I)
15:00 - 15:30	TEA BREAK			
Session 8 15:30 - 17:30		**Session 0908** Site & SMO		**Session 1108** Regulatory Aspect of the Innovative Partnership (Part II)