大会详细日程				
日期 Date	分会场主题 Theme	时间 Time	演讲主题 Topic	演讲嘉宾 Speaker
9月4日 Sep. 4th	药品注册电子申报分会场 Electronic Submission of Drug Registration Parallel Session	13:30-14:10	全球药品电子申报现状及我国电子申报政策法规 The Current Status of Global Drug Electronic Submission and China's Electronic Submission Policies and Regulations	代丹 女士，北京益睿思信息科技有限公司咨询服务总监 Ms. Dan Dai, Director of Consulting Services, Beijing ERIS Information Technology Co., Ltd.
		14:10-15:00	中国药品电子申报实践经验分享 Sharing of Practical Experience in Drug Electronic Submission in China	王芳 女士，苏州泽璟生物制药股份有限公司药品注册事务部副总监 Ms. Fang Wang, Deputy Director of Regulatory Affairs Department, Suzhou Zelgen Biopharmaceuticals Co., Ltd.
		15:00-15:20	茶歇 Tea Break	
		15:20-16:00	药品申报信息及文档管理解决方案（RIM）介绍 Introduction to the Drug Submission Information and Document Management Solution(RIM)	刘作为 先生 ，OpenText生命科学行业总监 Mr. Leo Liu, Director for the Life Sciences Industry, OpenText
		16:00-16:40	药品注册文档生命周期管理与RIM应用经验分享 Experience Sharing on Lifecycle Management of Drug Registration Documents and RIM Application	刘志伟 先生，亿腾医药注册副总监 Mr. Zhiwei Liu, Deputy Director of Regulatory Affairs, Edding Pharm
		16:40-17:20	圆桌讨论：药品电子申报法规与实践讨论 Panel Discussion: Drug Electronic Submission Regulations and Practice Discussion	代丹 女士，北京益睿思信息科技有限公司咨询服务总监 王芳 女士，苏州泽璟生物制药股份有限公司药品注册事务部副总监 刘作为 先生 ，OpenText生命科学行业总监 刘志伟 先生，亿腾医药注册副总监 Ms. Dan Dai, Director of Consulting Services, Beijing ERIS Information Technology Co., Ltd. Ms. Fang Wang, Deputy Director of Regulatory Affairs Department, Suzhou Zelgen Biopharmaceuticals Co., Ltd. Mr. Leo Liu, Director for the Life Sciences Industry, OpenText Mr. Zhiwei Liu, Deputy Director of Regulatory Affairs, Edding Pharm
		17:20-17:30	会议总结 Summary	