



2008 Clinical Trial Symposium on International Standard for Operation of Clinical Research Site &
The First China Clinical Research Site Summit

临床研究基地国际化管理研讨会 暨：中国首届临床研究基地管理峰会



Date: 15-16 December 2008

Venue: G/F, Kai Chong Tong, School of Public Health, The Chinese University of Hong Kong (CUHK) Prince of Wales Hospital, Shatin, NT

15 December 2008 (Monday) – Day 1	
08:30-09:00	Registration
09:00-09:45	<p>Welcome and Opening Address</p> <p>Prof FOK Tai Fai, Dean of Faculty of Medicine, CUHK 霍泰辉院长 香港中文大学医学院院长</p> <p>Prof ZHU, Yizhun, Dean of School of Pharmacy, Fudan University, China 朱依淳教授 复旦大学药学院院长兼职教授</p> <p>Prof Benny Zee, Director of Centre for Clinical Trials (CCT), School of Public Health, CUHK 徐仲镛教授 香港中文大学医学院助理院长(研究)公共卫生学院临床试验中心教授/中心主任</p>
09:45-10:30	<p>Plenary Session</p> <p>Towards the international standard in conducting clinical research in China: the Implementation of Good Clinical Practice "深入规范GCP的执行 促使中国临床研究国际化"</p> <p>Ms CAO Cai, Vice-Director, CCD of the SFDA of China 曹彩主任药师 国家食品药品监督管理局药品认证管理中心副主任</p>
10:30-11:00	Break (coffee/tea)
11:00-11:30	<p>Clinical research site management in China: current status and forecast "内地临床研究机构管理"</p> <p>Prof LEE Xuening, Director, Organization for Drug Clinical Trial, Zhongshan Hospital, Fudan University, China 李雪宁教授 上海中山医院药物临床实验机构主任</p>
11:30-12:00	<p>Clinical research site management in China: case studies (1) 中国疫苗临床研究中GCP的执行情况</p> <p>Prof LI Rongcheng, Director of the Division of Virology Research, Guangxi Zhuang Autonomous Region Center for Disease Prevention and Control 李荣成副主任医师 广西壮族自治区疾病预防控制中心病毒科科长</p>
12:00-12:30	<p>Clinical research site management in China: case studies (2) 在中国实施大型疫苗临床研究的经验分享</p> <p>Prof ZHU Fengcai, Jiangsu Centers of Disease Prevention and Control, China, Chief 朱凤才主任医师 硕士生导师 (江苏省疾病预防控制中心)</p>
12:30-14:00	Lunch
14:00-14:30	<p>The importance of industry-independent academic research in clinical practice</p> <p>Prof Francis K L Chan, Professor, Department of Medicine & Therapeutics, CUHK 陈家亮教授 香港中文大学医学院内科及药物治疗学讲座教授</p>
14:30-15:00	<p>Balance between Science & Ethics - Roles of clinical research ethics committee</p> <p>Prof Benny C Y Zee, Director of CCT, School of Public Health, CUHK 徐仲镛教授 香港中文大学医学院助理院长(研究)公共卫生学院临床试验中心教授/中心主任</p>
15:00-15:30	Break (coffee/tea)
15:30-16:00	<p>Lessons learnt from the FDA inspection/audit</p> <p>Prof Juliana C N Chan, Professor, Department of Medicine & Therapeutics, CUHK 陈重娥教授 香港中文大学医学院内科及药物治疗学讲座教授</p>
16:00-16:30	<p>Clinical trial site operation and management: perspectives from international pharmaceutical companies</p> <p>Dr TANG Haiwen, Director R&D, Medical Affairs Vaccines, North Asia, GSK 唐海文博士 葛兰素史克生物制品北亚地区医学部总监</p>
16:30-17:30	Panel Discussion (1): What is the expected role of clinical trial centre?

	16 December 2008 (Tuesday) – Day 2
09:00-09:30	US FDA GCP regulated clinical trials: case studies <i>Ms Anne POHLMAN, Clinical Research Coordinator, Department of Medicine, University of Chicago, USA</i>
09:30-10:00	Good Clinical Practice – a pre-requisite: Is it demanding? <i>Dr Christopher HUANG, the Managing Director of PHARMACONS (CRO) – China/Australia)</i> 黄晓博士 法码康 (CRO,中国/澳大利亚)总裁
10:00-10:30	Clinical research from non-industry sponsored perspective <i>Dr LI Chi Kong, COS, Department of Paediatrics, Prince of Wales Hospital, Hospital Authority, Hong Kong</i> 李志光醫生 香港医院管理局威尔斯亲王医院儿科部门主管
10:30-11:00	Break (coffee/tea)
11:00-12:00	Panel Discussion (2) Is there a bigger role for clinical trials centre in conducting clinical research?
12:00-12:30	Conference Closing <i>Prof Benny C Y Zee, Director of CCT, School of Public Health, CUHK</i> 徐仲镓教授 香港中文大学医学院助理院长(研究)公共卫生学院临床试验中心教授/中心主任
12:30-14:00	Lunch
14:00-14:30	Visit The School of Public Health, CUHK

Organizers:



Centre for Clinical Trials, School of Public Health
The Chinese University of Hong Kong



School of Pharmacy
Fudan University

Co-Organizer:



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