HKAPI Regulatory Affairs Course, Seminar 6

TOPIC: Be a Competitive Regulatory Expert of Clinical Trial in Hong Kong

Learning Outcomes:

- To gain an understanding of the clinical research development of the Asian region
- To understand the regulations and policy trend pertaining to the clinical research development
- To apply the concepts and learning gained in the context of the HK regulatory landscape

Date: 23 Nov 2017 (Thu)

Venue: Lecture Theatre 2, 1/F, William MW Mong Block, 21 Sassoon Road, Pokfulam

Language: Cantonese

Agenda:

Time (pm)	Topic	Speaker
2:00 – 2:05	Opening Remarks	
2:05- 2:45	Clinical Research Development and Policy Trends - Hong Kong and Asia region	Henry Yau *Managing Director & Honorary Assistant Professor* *Clinical Trial Centre, The University of Hong Kong*
2:45- 3:15	Impact of Regulatory Affairs in Understanding Clinical Data Submission	Kim Lee *HKAPI Clinical Trial Taskforce Lead*
3:15- 3:30	**Break**	
3:30- 4:00	Clinical Trials: Law and Regulations in Hong Kong	Vincent Chiang *Senior Pharmacist (Drug Import and Export)* *Drug Office, Department of Health*
4:00- 4:30	Update and Overview of China Regulations on Clinical Trials	Clement Ngai *Partner, Baker & McKenzie LLP*
4:30-5:20	Panel Discussion: Competitive Advantages of Hong Kong in Doing Clinical Trials – APAC perspective	Henry Yau Kim Lee Vincent Chiang Clement Ngai
5:20-5:30	**Closing Remarks**	