International Workshop: The Chinese and the European Pharmacopoeias - The New Editions -
18 March 2016

Location: EDQM premises, Strasbourg, France
Working languages: English/Chinese (simultaneous interpretation provided)

FINAL PROGRAMME

Morning session:

Moderators:
Dr Susanne Keitel, Director, EDQM and Mr Wei Zhang, Secretary General, Chinese Pharmacopoeia Commission

Opening Remarks & Welcome Address
9h00-9h15 Dr Susanne Keitel, Director, EDQM, Council of Europe
Mr Wei Zhang, Secretary General, Chinese Pharmacopoeia Commission

The Chinese Pharmacopoeia Commission (ChP)
- Mission, scope and organisation of the Chinese Pharmacopoeia Commission
9h15-9h35 Mr Wei Zhang, Secretary General, Chinese Pharmacopoeia Commission

- The 2015 edition of the Chinese Pharmacopoeia: what’s new?
Brief introduction about 2015 Edition Chinese Pharmacopoeia
9h35-9h55 Mr Wei Zhang, Secretary General, Chinese Pharmacopoeia Commission

Chinese traditional medicines and materials
9h55-10h25 Prof. Dean Guo, Expert of Chinese Pharmacopoeia Commission

Chemicals section
10h25-10h55 Ms Xiaohong Zhang, Deputy Director of Chemicals Department, Chinese Pharmacopoeia Commission

10h55-11h10 Coffee Break
Biologicals, Excipients & packaging materials sections

11h10-12h00 Dr Xiaoxu Hong, Deputy Director of Comprehensive Department, China Pharmacopeia Commission

12h00-12h30 Discussion

12h30-13h45 Lunch break

EDQM and the European Pharmacopoeia (Ph. Eur.)
- The 9th edition of the European Pharmacopoeia & current hot topics
13h45-14h25 Mrs Cathie Vielle, Secretary to the Ph. Eur. Commission & Head of the European Pharmacopoeia Department, EDQM, Council of Europe
- Ph. Eur. reference standards
14h25-15h05 Dr Andrea Lodi, Head of the Laboratory, EDQM, Council of Europe
- The Certification of suitability procedure & the EDQM inspection programme
15h05-15h45 Mrs Hélène Bruguera, Head of the Certification Division, EDQM, Council of Europe

15h45-16h05 Discussion

The Importance of Pharmacopoeial Standards and Their Added Value for Stakeholders
- Round table discussion with Chinese and European industry associations
  Speakers and panellists:
  - Mr. Koen Nauwelaerts, Quality and Regulatory Manager EGA, (European Generic and Biosimilar Medicines Association)
  - Mr. Lionel Randon, Biotech Regulatory CMC Merck, EBE (European Biopharmaceutical Enterprises)
  - BEng Merieke Van Dalen, Aspen Oss, CEFIC (European Chemical Industry Council)
  - Dr Mark Wiggins, MSD, EFPIA (European Federation of Pharmaceutical Industries and Associations)
  - Dr Frank Milek, Aug. Hedinger, IPEC (Internal Pharmaceutical Excipients Council Europe)
16h05-17h05 All speakers, moderators of the day, and industry representatives

CLOSING REMARKS

17h05 – 17h15 Dr Susanne Keitel, Director, EDQM, Council of Europe and
Mr Wei Zhang, Secretary General, Chinese Pharmacopoeia

Ca. 17h15 Networking reception