International Workshop:
The Chinese and the European Pharmacopoeias
  - The New Editions -
17 October 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

1. 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 Mr Pengfei Tu, 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

11h10-12h15 Dr Xiaoxu Hong, Deputy Director of Comprehensive Department, Chinese Pharmacopeia Commission

12h15-12h30 Discussion

12h30-13h45 Lunch break

2. EDQM and the European Pharmacopoeia (Ph. Eur.)
   - Ph. Eur. reference standards
     13h45-14h20 Dr Andrea Lodi, Head of the Laboratory, EDQM, Council of Europe
   - The 9th edition of the European Pharmacopoeia & current hot topics
     14h20-15h25 Mrs Cathie Vielle, Secretary to the Ph. Eur. Commission & Head of the European Pharmacopoeia Department; Dr Ulrich Rose, Head of Division A, European Pharmacopoeia Dept., Mr Gwenael Cirefice, Scientific Officer of Division B, European Pharmacopoeia Dept., EDQM, Council of Europe
   - The Certification of suitability procedure & the EDQM inspection programme
     15h25-16h10 Ms Fiona McLeod, Scientific Programme Officer, Certification of Substances Division, EDQM, Council of Europe

16h10-16h30 Coffee Break

3. The Importance of Pharmacopoeial Standards and Their Added Value for Stakeholders
   - Round table discussion with Chinese and European industry associations
     - Mr Koen Nauwelaerts from Medicines for Europe, (European Generic and Biosimilar Medicines Association)
     - Mrs Barbara Freischem & Mr Lionel Randon from EBE (European Biopharmaceutical Enterprises)
     - Ms Hilde Vanneste from CEFIC (European Chemical Industry Council)
     - Mr Mark Wiggins from EFPIA (European Federation of Pharmaceutical Industries and Associations)
     - Dr Frank Milek from IPEC (Internal Pharmaceutical Excipients Council Europe)

16h30-17h30 All speakers, moderators of the day, and industry representatives

CLOSING REMARKS

17h30 – 17h40 Dr Susanne Keitel, Director, EDQM, Council of Europe and
Mr Wei Zhang, Secretary General, Chinese Pharmacopoeia Commission

17h40 Closure