



2016 TRAINING SESSION: STRASBOURG
EUROPEAN PHARMACOPOEIA REFERENCE STANDARDS
19 APRIL 2016, STRASBOURG, FRANCE
Working language: English

PROGRAMME *(Subject to change)*

- 8:00-9:00** Registration
- 9:00-9:10** **Opening & Welcome address**
Susanne Keitel, Director, EDQM, Council of Europe
- 9:10-9:30** **European Pharmacopoeia Reference Standards (RS): A Global Overview**
Andrea Lodi, Head of the Laboratory Dept., EDQM, Council of Europe
- 9:30-9:55** **Chemical RS for qualitative purposes**
Jochen Pauwels, Laboratory Department, EDQM, Council of Europe
- 9:55-10:25** **Chemical RS for quantitative purposes**
Stefan Almeling, Deputy Head of the Laboratory Department, EDQM, Council of Europe
- 10:25-10:45** **Open discussion with the panel of speakers**
- 10:45-11:00** *Coffee break*
- 11:00-12:30** **Parallel Sessions**

Session 1	Session 2
<p>11:00 RS for Synthetic Peptides Frank Jung, Laboratory Department, EDQM, Council of Europe</p> <p>11:20 RS for Recombinant Proteins Sylvie Jorajuria, Laboratory Department, EDQM, Council of Europe</p> <p>11:40 RS for Biologicals established via the Biological Standardisation Programme Eriko Terao, Dept. of Biological Standardisation, OMCL Network & HealthCare, EDQM, Council of Europe</p> <p>12:10 Open discussion</p>	<p>11:00 Orthogonal RS characterisation techniques Matthias Weber, Laboratory Department, EDQM, Council of Europe</p> <p>11:20 Monitoring the quality of CRS Philippe Duret, Laboratory Department, EDQM, Council of Europe</p> <p>11:50 RS in herbal monographs Matthias Weber</p> <p>12:10 Open discussion</p>

- 12:30-14:00** *Lunch break*
- 14:00-14:30** **Production, Storage and Transport of Ph. Eur. Reference Standards**
Vincent Egloff, Head of the Reference Standards & Samples Division, EDQM, Council of Europe
- 14:30-15:00** **RS Documentation/QA & Safety Aspects/Regulations**
Jonna Sunell-Huet, Deputy Head of the Quality, Safety & Environment Division, EDQM, Council of Europe



- 15:00-15:20 Role of RS in the Ph. Eur.: Good practices, dos and don'ts**
Emmanuelle Charton, Head of Division B, European Pharmacopoeia Dept., EDQM, Council of Europe
- 15:25-15:40** *Coffee break*
- 15:40-15:50 Secondary Standards**
Andrea Lodi
- 15:50-16:00 Reference Standards: An inspector's viewpoint**
Sotirios Paraschos, Certification Division, EDQM, Council of Europe
- 16:00-16:30 Open discussion with the panel of speakers**
- 16:30** *Close of the meeting*

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