EDQM and European Pharmacopoeia: State-of-the-art Science for Tomorrow’s Medicines

International Conference organised by the European Directorate for the Quality of Medicines & HealthCare (EDQM), Council of Europe,

on the occasion of


19-20 June 2019, Strasbourg, France

PROGRAMME
Wednesday, 19 June 2019

08:00-08:45 Registration & Welcome Coffee (Hemicycle - Palais de l’Europe)

OPENING SESSION

08:45-09:00 Welcome Addresses
Ms Snežana Samardžić-Marković, Director General of Democracy, Council of Europe
Jean-Baptiste Mattéi, Ambassador, Permanent Representative of France to the Council of Europe
Dr Susanne Keitel, Director, EDQM, Council of Europe

PLENARY SESSION

Moderator: Dr Tobias Gosdschan
Outgoing Chair of the European Pharmacopoeia Commission

09:00-09:30 Building Synergies to strengthen and support healthcare in Europe
Dr Andrzej Rys, Director Health Systems, Medical Products and Innovation, European Commission

09:30-10:00 Globalisation and cross-border cooperation in Europe: the perspective of a national competent authority
Dr Dominique Martin, Director General, Agence nationale de sécurité du médicament et des produits de santé (ANSM)

10:00-10:30 Contributions of the European OMCL Network to the protection of Public Health
Ms Patricia Courselle, Former Chair of the OMCL Advisory Group

10:30-11:00 Coffee break and Poster session

11:00-11:30 The Certification of Suitability Procedure - 25 years of a success story
Dr Jean-Louis Robert, Chair of the Certification Steering Committee

11:30-12:00 Combatting falsified medicines - the EDQM’s holistic approach in support of the MEDICRIME Convention
Dr Karl-Heinz Buchheit, EDQM, Council of Europe

12:00-12:30 European Directorate for the Quality of Medicines & HealthCare (EDQM) & the European Pharmacopoeia (Ph. Eur.)
Dr Susanne Keitel, Director, EDQM, Council of Europe

12:30-14:30 Lunch break (EDQM premises)
Wednesday, 19 June 2019

WORKSHOP SESSIONS

Five workshops will run in parallel with each session being repeated once, except for Biotherapeutics, Certification, 3Rs and ATMPs and OMCL Network. Interpretation into French will not be provided during the workshop sessions.

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<td>14:30-17:30</td>
<td>Impurities</td>
<td>Finished Product Monographs</td>
<td>Biotherapeutics</td>
<td>General Methods</td>
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WORKSHOP: IMPURITIES
19 June 2019, Room 100

Moderator: Prof. Torbjörn Arvidsson, Chair of the Ph. Eur. Group of Experts on Organic Chemistry (10A)

14:30-15:00 Impurity Control in the European Pharmacopoeia (Ph. Eur.)
Dr Gabriella Török, Chair of the Ph. Eur. Group of Experts on Organic Chemistry (10B)

15:00-16:00 Challenges linked to the Control of Antibiotics: the EU Antibiotics Guideline and its impact on dossiers and assessments.
Dr Jan Smeets, Centrient Pharmaceuticals Netherlands B.V, the Netherlands
Dr Uwe Lipke, Federal Institute for Drugs and Medical Devices (BfArM), Germany

16:00-16:30 Coffee break

16:30-16:50 Experience of implementation of ICH Q3D within the Certification Procedure (CEPs)
Dr Cristian Sampaolesi, EDQM, Council of Europe

16:50-17:10 Control of Impurities: Challenges linked to the Establishment of Reference Standards
Dr Jochen Pauwels, EDQM, Council of Europe

17:10-17:30 Feedback from Users
Panel discussion with the moderator and the speakers

17:30 Close of the first day – Reception (Palais de l’Europe)
WORKSHOP: FINISHED PRODUCT MONOGRAPHS
19 June 2019, Room 550
Moderator: Dr Tobias Gosdschan, Outgoing Chair of the European Pharmacopoeia Commission

14:30-14:50 Experiences of The U.S. Pharmacopeial Convention (USP)
Mr Bruk Alemayehu, U.S. Pharmacopeial Convention, United States of America

14:50-15:10 Perspective of a Regulatory Authority
Ms Andrea Cseh-Palos, National Institute of Pharmacy and Nutrition, Hungary

15:10-15:30 Perspective of an OMCL
Dr Charlotte Brenier, National Agency for Medicines and Health Products Safety (ANSM), France
[Chair of the Ph. Eur. Group of Experts on Synthetic / Semi-synthetic Products (10D)]

15:30-16:00 Coffee break

16:00-16:40 Industry’s Perspective
From an Innovator: Dr Veronique Pinilla, UCB Pharma, Belgium
From a Generic manufacturer: Dr Manish Gangrade, Cipla Ltd., India

16:40-17:00 Experiences of the European Pharmacopoeia
Dr Ulrich Rose, EDQM, Council of Europe

17:00-17:30 Feedback from Users: Panel discussion with the moderator and the speakers

17:30 Close of the first day - Reception (Palais de l’Europe)

THE NEED FOR MONOGRAPHS ON BIOOTHERAPEUTICS
(ACTIVE PHARMACEUTICAL INGREDIENTS AND FINISHED PRODUCTS)
19 June 2019, Room 11
Moderator: Mr Peter M J Jongen, Chair of the Ph. Eur. Group of Experts on P4 Biologicals

14:30-14:50 Biotherapeutic Products in the Ph. Eur.: have all the challenges been tackled?
Dr Mihaela Buda, EDQM, Council of Europe

14:50-15:00 Discussion

15:00-15:20 Reference Standards for Biotherapeutic Products
Dr Marie-Emmanuelle Behr-Gross and Dr Sylvie Jorajuria, EDQM, Council of Europe

15:20-15:30 Discussion

15:30-16:00 Coffee break

16:00-16:20 Industry’s Perspective
Mr Lionel Randon, Merck Serono SA, Switzerland and Dr Emmanuel Rossy, Novartis, Germany

16:20-16:30 Discussion
16:30-16:50 **Perspective of a Regulatory Authority**  
Dr Martijn Van der Plas, Medicines Evaluation Board (MEB), the Netherlands

16:50-17:00 Discussion

17:00-17:30 **Feedback from Users**: Panel discussion with the moderator, the speakers and  
Dr Joseph Albanese, Janssen J&J, USA

17:30 *Close of the first day - Reception (Palais de l'Europe)*

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**WORKSHOP: GENERAL METHODS**  
**19 June 2019, Room 10**

**Moderator**: Prof. Dr Michel Ulmschneider, Chair of the General Methods Working Party

14:30-14:50 **Chromatographic Separation Techniques and Challenges related to harmonisation**  
Prof. Dr Jos Hoogmartens, Chair of the Ph. Eur. Group of Experts on Chromatographic Separation Techniques

14:50-15:10 **Feedback on the work of the Ph. Eur. General Methods Working Party**  
Prof. Dr Michel Ulmschneider, Chair of the General Methods Working Party

15:10-15:30 **Reference Standards for Ph. Eur. Chapters**  
Dr Stefan Almeling, EDQM, Council of Europe

15:30-16:00 Coffee break

16:00-16:20 **Feedback on the work of the Ph. Eur. Vibrational Spectroscopy and Analytical Data Modelling (VSADM) Working Party**  
Prof. Dr Manel Alcala Bernardez, Universitat Autonoma de Barcelona, Spain

16:20-16:40 **Analytical QbD - Industry’s Perspective**  
Dr Graham Cook, Pfizer, United Kingdom

16:40-17:00 **Continuous manufacturing: what impact on the Pharmacopoeia**  
Dr Moheb M. Nasr, Nasr Pharma Regulatory Consulting (NPRC), United States of America

17:00-17:30 **Feedback from Users**: Panel discussion with the moderator and the speakers

17:30 *Close of the first day - Reception (Palais de l'Europe)*
WORKSHOP: OMCL NETWORK
19 June 2019, Room 8
Moderator: Dr Maria João Portela, Infarmed, Portugal

14:30-15:00 Official Control Authority Batch Release (OCABR): benefits, challenges and perspectives
Dr Volker Öppling, Paul-Ehrlich-Institut (PEI), Germany

15:00-15:30 Market Surveillance Studies: OMCL contribution to quality and safety of medicines on the market
Dr Lone Olsen, Danish Medicines Agency (DKMA), Denmark

15:30-16:00 Coffee break

16:00-16:40 API Testing: how OMCLs can support the control of APIs
Dr Eric Deconinck, Sciensano, Belgium and Mr Yvan Grange, National Agency for Medicines and Health Products Safety (ANSM), France

16:40-17:00 The contributions of OMCLs in the fight against Falsified and Illegal Medicines
Mr Stephen Young, Medicines and Healthcare products Regulatory Agency (MHRA), United Kingdom

17:00-17:30 Feedback from Users: Panel discussion with the moderator and the speakers

17:30 Close of the first day – Reception (Palais de l’Europe)
Thursday, 20 June 2019

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12:00-14:00 Lunch break (EDQM premises)

WORKSHOP: THE 3Rs AND ATMPs
20 June 2019, Room 11
Moderator: Dr Lukas Bruckner, Chair of the Biological Standardisation Programme Steering Committee

09:00-09:20 Achievements of the Biological Standardisation Programme and the Ph. Eur. in the field of the 3Rs
Dr Catherine Milne and Dr Gwenael Cirefice, EDQM, Council of Europe

09:20-09:40 The 3Rs: perspectives for the future
Dr Lukas Bruckner, Chair of the Biological Standardisation Programme Steering Committee
Mr Arnoud Akkermans, National Institute for Public Health and Environment (RIVM), the Netherlands

09:40-10:00 Vet Vaccines perspective
Dr Elizabeth Kamphuis, Boehringer Ingelheim Vetmedica GmbH, Germany

10:00-10:15 Feedback from Users
Panel discussion with the moderator and the speakers

10:15-10:45 Coffee break
10:45-11:15 **ATMPs: how can Ph. Eur. fulfil its role for tomorrow’s medicines? Viewpoint of OMCL/Regulator**
Dr Marie-Cristina Galli, Istituto Superiore di Sanità (ISS), Italy

11:15-11:45 **Viewpoint of Industry**
Dr Mehrshid Alai-Safar, Kite Pharma, United States of America

11:45-12:00 **Feedback from Users**
Panel discussion with the moderator and the speakers.

12:00-14:00 Lunch break (EDQM premises)

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12:00-14:00 Lunch break (EDQM premises)
WORKSHOP: CERTIFICATION  
20 June 2019, Room 8  
Moderator: Dr Jean-Louis Robert, Chair of the Certification Steering Committee

09:00-09:30 **EU Active Substance Master File (ASMF) work-sharing: Viewpoint of a Regulator**  
Ms Nienke Rodenhuis, Medicines Evaluation Board (MEB), the Netherlands

09:30-10:00 **EU ASMF work-sharing and CEPs: Viewpoint of Industry**  
Dr Martijn Klop, Synthon BV, the Netherlands

10:00-10:30 Coffee break

10:30-11:00 **The International Pharmaceutical Regulators Programme (IPRP) and its Quality Working Group for Generics**  
Mr Gary Condran, Health Canada, Canada

11:00-11:20 **What's new in Certification**  
Ms Hélène Bruguera, EDQM, Council of Europe

11:20-11:40 **Update on quality guidelines for the control of active substances**  
Dr Olaf Ludek, Icelandic Medicines Agency (IMA), Iceland

11:40-12:00 **Feedback from Users**  
Panel discussion with the moderator and the speakers

12:00-14:00 **Lunch break (EDQM premises)**
Thursday, 20 June 2019
Hemicycle - Palais de l’Europe

CLOSING PLENARY SESSION
Moderator:
Prof. Torbjörn Arvidsson, Incoming Chair of the European Pharmacopoeia Commission

14:00-14:30 International Developments
Dr Petra Dörr, Swissmedic, Switzerland

14:30-15:00 The Role of the EDQM in International Harmonisation Initiatives
Ms Cathie Vielle, EDQM, Council of Europe

15:00-15:30 Coffee break and Poster session

15:30-16:30 Reports from the Workshop Sessions & Discussions

16:30-16:45 Final Conclusions & Closing Remarks
Prof. Torbjörn Arvidsson, Incoming Chair of the European Pharmacopoeia Commission
Dr Susanne Keitel, Director, EDQM, Council of Europe

16:45 Close of the conference

More information is available at:
www.edqm.eu

E-Mail: prdd@edqm.eu

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