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Customer Event Almaty

21 September 2023

Agenda

08:30–09:00 . . . Sign in and welcome

09:00–09:30 . . . Keynotes

- Dr. Venkat Surendra Nath Koduru, Senior Vice President, Regions & Program Operations (10')
- Dr. Alessandro Slama, General Manager, USP EMEA (10')
- Dr. Yerken Dautbaev, General Director - Chairman of the Board of National Center for Drug Expertise (10')

09:30–10:50 . . . Session 1 - Pharmacopoeia quality standards

Time	Topic	Speaker
09:30 – 10:00	General Overview of USP: Key Activities and Role in Assuring the Quality of Medicines	Stefano D'Amico, USP Strategic Customer Development Manager EMEA
10:00 – 10:20	The State Pharmacopoeia Kazakhstan as a fundamental document on the quality of medicines	Ardak Tulegenova, Head of the Department of the SPh, National Center for Medicines and Medical Devices MoH of Kazakhstan
10:20 – 10:50	The value of USP Reference standards (online)	Christian Zeine, USP Senior Scientific Affairs Manager EMEA
Moderated by: Dr. Zakiya Al-Kurdi		

10:50–11:10 . . . Coffee Break

11:10–12:30 . . . Session 2 - Drug manufacturing major aspects

Time	Topic	Speaker
11:10 – 11:30	Contracting manufacturing development in Kazakhstan: opportunities and barriers	Marina Durmanova, President, Pharma Industry Support and Development Association, Kazakhstan
11:30 – 11:50	Challenges of international pharmaceutical manufacturers in Kazakhstan	Svetlana Ospanova, Executive Director, Association of International Pharmaceutical Manufacturers in Kazakhstan
11:50 – 12:10	Development of pharmaceutical clusters in Uzbekistan	Ulugbek Elgamov, Director, Pharmaceutical Industry Development Agency, Uzbekistan
12:10 – 12:30	Validation, Transfer and Verification of Analytical Procedures	Amanda Guiraldelli, USP Scientific Affairs Manager EMEA
Moderated by: Dr. Marina Durmanova		

12:30–13:30 . . . Lunch

13:30–15:50 . . . Session 3 - Drug quality major aspects

Time	Topic	Speaker
13:30 – 14:00	Challenges during impurity testing and Pharmaceutical Analytical Impurities	Christian Zeine, USP Senior Scientific Affairs Manager EMEA
14:00 – 14:30 –	Nitrosamines Impurities	Amanda Guiraldelli, USP Scientific Affairs Manager EMEA
14:30 – 14:50	Control of visible particulate matter during the production of parenteral medicinal products	Bereke Tanaguzova, Quality Director & QP in Karaganda Pharmaceutical Complex
14:50 – 15:10	The experience of Kazakhstan in the development and production of the COVID-19 vaccine «QazVac»	Yergali Abduraimov, General Director, QazBioPharm | Lespek Kutumbetov, Chief Scientist, Biological safety Scientific Research Institute by the MoH of Kazakhstan
15:10 – 15:30	USP and Vaccine Quality: Standards to support public health and safety (online)	John Cipollo, Team Lead, Senior Principal Scientist Global Biologics
Moderated by: Dr. Bayan Moldakhmetova		

15:30–16:00 . . . Open discussion: How to organize an effective training on drug quality issues?

Moderated by: Dr. Alessandro Slama, Dr. Stefano D'Amico

16:00–16:30 . . . Wrap Up and Closure

- Venkat Surendra Nath Koduru, Senior Vice-President for Regional and Programme Activities, USP (10')
- Alessandro Slama, Senior Director for EMEA Regional Programs, USP (10')
- Ardak Tulegenova, Head of the Department of the SPh, National Center for Medicines and Medical Devices MoH of Kazakhstan (10')

Venue

The forum will be held at the DoubleTree, 115 Dosmukhamedov Street, Almaty 0500-00, Kazakhstan