CENTER FOR PHARMACEUTICAL PRODUCTS SAFETY

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Customer Event Tashkent

25 April 2024

Agenda

08:00–09:00 . . . Sign-in and welcome

09:00–09:30 . . . Keynotes

- Venkat Surendra Nath Koduru, Senior Vice President, Regions & Program Operations USP (10')
- Alessandro Slama, Vice President, Regions & Program Operations USP (10')
- Alisher Temirov, Interim Director, Center for Pharmaceutical Products Safety (10')

09:30–14:30 . . . Session 1 - Pharmacopoeia quality standards and current regulatory issues

Time	Topic	Speaker
09:30 10:00	Perspectives of Localization of Pharmaceutical Manufacturing	Abdulla Abdisalamovich Azizov, Director of the Pharmaceutical Industry Development Agency
10:00 10:30	Medicines We Can Trust: USP Impact	Stefano D'Amico, USP Strategic Customer Development Manager EMEA
10:30 11:00	USP Comprehensive Solutions Helps To Mitigate Risks To Pharmaceutical Quality and Accelerate Product Development	Amanda Guiraldelli, USP Scientific Affairs Manager EMEA
11:00 11:15	Coffee Break	
11:15 11:45	Current issues of development and implementation in practice of the State Pharmacopoeia of the Republic of Uzbekistan	Khabibulla Jalilov, Head of Pharmacopeia Development Dept, Center for Pharmaceutical Products Safety, MoH Uzbekistan
11:45 12:00	Post-Marketing Surveillance: the shared responsibility of the manufacturer with public health	Gulnora Zufarova, Advisor to the Director, Center for Pharmaceutical Products Safety, MoH Uzbekistan
12:00 12:15	Pharmacovigilance	Elvira Svechnikova, Head ofPharmacovigilance Dept., Center for Pharmaceutical Products Safety, MoH Uzbekistan
12:15 12:30	Registration Process for Medicines and Medical Devices	Mukaramoy Ergashova, Head of Laboratory of pharmaco-toxicological analysis, Center for Pharmaceutical Products Safety, MoH Uzbekistan
12:30 13:00	Q&A with speakers: Oybek Khayrullaev, Head of Registration Dept., Center for Pharmaceutical Products Safety, MoH Uzbekistan Akmaral Kabdenova, The Director of the Laboratory Center of the NCEM branch in Almaty, Kazakhstan Ardak Tulegenova, Head of the Department for the Improvement of the SPh of Kazakhstan Mehmet Kursat Derici, VP of Economical Assessment and Laboratory Services - Turkey Yerken Dautbaev, General Director, National Center for Medicines and Medical Devices Evaluation, MOH Kazakhstan -	
13:00 14:00	LUNCH	
14:00 14:30	Flexibility approach and the possibility of its application in the development of pharmacopeial standards	Ardak Tulegenova, Head of the Department for the Improvement of the SPh of Kazakhstan
Moderated by: Foreign Pharma Association of Uzbekistan		

14:30–17:00 . . . Session 2 - Determination of Impurities in Pharmaceuticals: Why and How?

Time	Topic	Speaker
14:30 15:00	USP's Perspective on Impurities Control in Pharmaceuticals	Amanda Guiraldelli, USP Scientific Affairs Manager EMEA
15:00 15:30	Impurities in excipients: USP Tools and Regulatory Focus	Christian Zeine, USP Senior Scientific Affairs Manager EMEA
15:30 16:00	Measures to prevent adverse events related to contamination of medicines with EG and DEG in Uzbekistan	Khurshid Tursunov, Laboratory of Medicines Quality Control and Standardization, Centerfor Pharmaceutical Products Safety, MoH Uzbekistan
16:00 16:15	Coffee Break	
16:15 16:45	Validation, Transfer and Verification of Analytical Procedures: Enable Accurate, Reproducible Results While Streamlining Your Analyses	Amanda Guiraldelli, USP Scientific Affairs Manager EMEA
16:45 17:00	Q&A	
Moderated by: Stefano D'Amico, USP Strategic Customer Development Manager EMEA		

17:00–17:30 . . . Wrap Up and Closure

- Venkat Surendra Nath Koduru, Senior Vice President, Regions & Program Operations, USP (10')
- Alessandro Slama, Vice President, Regions & Program Operations USP (10')
- Alisher Temirov, Interim Director, Center for Pharmaceutical Products Safety (10')

Venue

Hyatt Regency, Navoi Street 1 A, Tashkent, Uzbekistan