USP Update on Chemical Medicines:

Chemical Medicines Plan for 2018-2020 and Beyond

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2018 – 2020 and Beyond

- Continue to optimize standards setting processes and systems to improve quality and productivity through an Up-To-Date (UTD) framework

- Continue to implement Process and Quality improvement recommendations.

- Continue to develop new and modernized monographs to support Up-To-Date.

- Work on Omitting monographs for obsolete medicines from the USP-NF to make the compendia as current as possible for industry and FDA,

- Continue collaboration with FDA and CHPA in the development of OTC compendial pathways

- Strengthen relationships with Industry and FDA to help modernize + develop new high priority monographs
In FY 19

- Work to improve standard development process and systems
- Develop capacity to solicit and use external stakeholder feedback to increase quality and output
- We will continue working with industry to Increase monograph donations with timely availability
- Develop and begin to deliver on a roadmap of QISP priorities through end of cycle
- Operationalize QISP policy decisions including metrics to measure effectiveness/success
- Define better quality operational metrics
Expanding scope to include a more strategic and integrated approach to improving operational efficiency and strengthening quality
Systems & Processes

Quality & Policy

Documentary Standards

Reference Standards
Quality & Policy + Systems & Process Initiatives

Quality & Policy:
- Examine the effectiveness of USP’s current scientific practices and policies in order to further improve the quality and scientific integrity of documentary and reference standards
- Identify gaps, areas for improvement, and training opportunities.

Systems and Process:
- Examine and document interdependences and capabilities to seek to develop streamlined processes and increase efficiency
- Improve ‘data-driven’ informed decision making within the organization
Thank You

Empowering a healthy tomorrow