Hybrid Event

PQRI
Product Quality Research Institute

SAVE THE DATES!

REGISTRATION TO OPEN IN FALL 2026.

PQRI Workshop:

Co-processed Excipients to Enhance Medicinal Product Development and Continuous Manufacturing

TUESDAY - WEDNESDAY, APRIL 14-15 2027

USP Meeting Center
12601 Twinbrook Pkwy
Rockville, MD

WORKSHOP OBJECTIVES

This 1-1.5-day PQRI Workshop will bring together leaders and subject matter experts (SMEs) from regulatory agencies, industry, and academia to discuss the various issues and views on Co-processed Excipients (CPEs) and what is needed to encourage the use of CPEs to enhance innovation in pharmaceutical development.

This workshop aims to facilitate interaction among stakeholders and will address the following topics

- What are CPEs and how are they made and controlled?
- Technology, Manufacturing, and Formulation Benefits of CPEs
- Safety Assessment of CPEs
- Regulatory Considerations and Challenges
- Marketing and Supply Chain Concerns
- Small group discussions ("breakout sessions") moderated by SMEs will further explore the above topics.

PQRI encourages anyone who is interested in CPEs use in product development to register for this workshop and join the in-depth discussions surrounding the technical benefits and regulatory hurdles that exist for CPEs and possible concepts for reducing the regulatory burden related to CPEs use in medicinal product development and continuous manufacturing.

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www.pqri.org