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Impurities in Drug Products and Drug Substances - A USP Approach (Live Webcast March 29-31, 2022)

Art. ID	USP-CM-476-01
Unit	Each

Description

This course integrates ICH Guidance and FDA policy for impurities, relevant USP General Chapters about impurities, and USP's approach to impurities in monographs. It also provides insights to USP-NF General Chapters <476> and <1086> and include case studies for impurities in the development and revision of USP monographs. /// Upon completion of this course, you will be able to: Discuss the origin and classification of impurities in pharmaceuticals. / Explain global guidance's for impurities. / Describe the USP approach to impurities in drug substances and drug products. / Discuss the USP general chapters on impurities. /Explain the USP approach to harmonization across pharmacopeia. / Demonstrate knowledge of USP's approach for impurities in documentary standards via case studies. /// Duration: 3/29/2022, 3:00 AM EDT - 3/31/2022, 5:30 AM EDT