Labmix24 GmbH
Kesseldorfer Rott 24
46499 Hamminkeln
Germany

Tel: +49 (0) 2852 96064 00
Fax: +49 (0) 2852 96064 24
Web: www.labmix24.com
E-Mail: info@labmix24.com

Development and Validation of Dissolution Procedures (On-Demand)

Art. ID	USP-CM-1092-01
Unit	Each
Deliverydetails	No Dangerous Good

Description

This course has been entirely revised in order to reflect the content of the complete overhaul of USP General Chapter <1092> published in USP 38-NF 33 First Supplement with the official date of August 1, 2015. Building on your basic understanding of USP's approach to dissolution, this course provides a foundation for developing and validating dissolution methods used for batch release and stability testing. These tests are in vitro performance tests for most dosage forms, such as tablets, capsules, suspensions, transdermal patches, and suppositories. They are important components of the specifications that establish the strength, quality, purity, and bioavailability of a drug product.