

AGENDA

Friday, 17 th of November 2023	
09:00	Welcome and Introduction
09:05	ICH-E6(R2): Good Clinical Practice Principles and Guideline Content Q&A
10:15	The Informed Consent Process Q&A
11:00	Break
11:20	Set-up of a Clinical Trial at the Site Q&A
12:00	Compliance in Document Management Q&A
13:00	End of Day 1
Saturday, 18 th of November 2023	
09:00	Critical Elements in Conducting Clinical Trials Q&A
10:00	PI's responsibilities in Safety Reporting in Clinical Trials Q&A
11:15	Risk-adapted Management of Clinical Trials Q&A
12:00	Multiple Choice Test
13:00	End of Day 2