SHODH CLINICALS
Deep Dive to Deliver Quality

FUNDAMENTALS OF CLINICAL RESEARCH: GCP & BIOETHICS



SATURDAY
21st Nov, 2026

PROGRAM OVERVIEW

This program is a comprehensive and practical introduction to global ethical and regulatory standards essential for conducting clinical trials. It covers ICH-GCP E6 (R3) guideline, ICMR guidelines, NDCT rules 2019 and key ethical considerations to ensure participant safety, data integrity, and regulatory compliance.

PROGRAM HIGHLIGHTS

- ICH-GCP E6(R3)
- ICMR Guidelines
- NDCT Rules 2019
- Pre & Post Assessment
- Certificate of Completion

WHY GCP MATTERS?

For Researchers
Encourages innovative designs and streamlined operations.

For IRB/IEC Members
Ensures ethical oversight & participant safety in clinical trial.

For Participants
Ensures safety, dignity, and transparency.

For the Industry
Promotes efficient, high quality, and globally compliant trials.

REGISTER NOW !

Fee: ₹2000/-
Per Participant

"Clinical research is the key to unlocking medical breakthroughs that transform patient care and improve lives."



– Dr. Francis Collins

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