



SHODH CLINICALS
Deep Dive to Deliver Quality

FUNDAMENTALS OF CLINICAL RESEARCH: GCP & BIOETHICS

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.



SATURDAY
5th Sep, 2026

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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