GEETANJALI UNIVERSITY

GEETANJALI INSTITUTE OF PHARMACY

SKILL DEVELOPEMENT PROGRAM

CliMed

SIX-DAY PROFESSIONAL DEVELOPMENT PROGRAM (PDP) ON

ICH GCP E6 (R3), BIOMEDICAL ETHICS, AND ICMR CLINICAL TRIAL GUIDELINES

ONLINE PDP

*EXCLUSIVE COVERAGE ON ANIMAL STUDIES IN DRUG DEVELOPMENT, ADME, PHARMACODYNAMICS & PHARMACOKINETICS

BY GEETANJALI INSTITUTE OF PHARMACY, GEETANJALI UNIVERSITY UDAIPUR & CLIMED RESEARCH SOLUTIONS, INDIA

- GET YOURSELF CERTIFIED IN GCP & BIOMEDICAL ETHICS
- 24-MONTHS CERTIFICATE VALIDATION WITH VERIFICATION

WHO MUST ATTEND:

- Pharmacy and Health professionals
- Researchers/Ph.D. Students
- Industry Delegates
- PG Students/Pharm D Interns
- Ethics Committee Members

LIMITED 50 SEATS

8-13 November 2024

6.30 PM - 8.00 PM FRI TO WED

Virtual on Zoom Platform

ETHICAL COMMITTEE EX.

Dr. Ashok Shenoy

Professor of Pharmacology
Kasturba Medical College
Mangalore

INDUSTRY EXPERT

Ms. Sarabjit Sidhu

Manager Clinical Trial Transparency, GSK Pharma

CLINICAL EXPERT

Dr. Walli Mohammed

Pharm. D, Pg. Dip (Clinical Research), Ph.D.
Clinical Research Specialist
Hyderabad

ACADEMIC PARTNER

Dr. Mahendra Singh Rathore

Principal, GIP, Geetanjali University, Udaipur, RJ

INDUSTRY PARTNER

Dr. Ajit Singh

Founder, CliMed & Curio, CliMed Research Solutions, India

FEE: 499/- INR ONLY

Register now at :

https://rzp.io/rzp/ichgcpnov

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www.climedacademy.com

PDP SCHEDULE

Resource Person	Topics	Date & Timing
Dr. Ashok Shenoy Professor of Pharmacology, Kasturba Medical College, Mangalore	Importance of ICH GCP & Biomedical Ethics Indian Bioethics Guidelines	8.11.2024 Friday 6.30 PM – 8.00 PM
Ms. Sarabjit Sidhu Manager Clinical Trial Transparency, GSK Pharma	History and Origin of Ethical Principles in Clinical Research & ICH: history, purpose, principles, guidelines	9.11.2024 Saturday 6.30 PM – 8.00 PM
Ms. Sarabjit Sidhu Manager Clinical Trial Transparency, GSK Pharma	Good Clinical Practice (GCP)- ICH E6: Responsibilities of Sponsors, Investigators, Monitor Ethics committees: Roles & responsibility of IEC and IRB	10.11.2024 Sunday 6.30 PM – 8.00 PM
Ms. Sarabjit Sidhu Manager Clinical Trial Transparency, GSK Pharma	Protocol contents and Types of Clinical Study Design Informed consent form and consenting process	11.11.2024 Monday 6.30 PM – 8.00 PM
Dr. Ajit Singh Founder & CEO CliMed & Curio Research Scientist for ICMR, KMC, MAHE	New Drug Rules 2019 & ICMR Bioethics Guidelines	12.11.2024 Tuesday 6.30 PM – 8.00 PM
Dr. Walli Mohammed Pharm. D, Pg. Dip (Clinical Research), Ph.D. Clinical Research Specialist	Animal studies in drug development, ADME, Pharmacodynamics & Pharmacokinetics	13.11.2024 Wednesday 6.30 PM – 8.00 PM

WHY YOU SHOULD ATTEND:

- **Upgrade yourself with the latest ICH GCP E6 R3 Guidelines**
- Get certified in Good Clinical Practice Guidelines, ICMR Clinical Trial Guidelines
- Update yourself on updates after the COVID-19 era
- Learn from the industry and academic experts for both aspects
- Special coverage on - Ethical Committee Formation, Role of Responsibilities of various stakeholders, Informed consent form and compensation in clinical trials
- Keep yourself ready to participate in clinical trials and grant proposals
- Perfect Networking opportunities