



AMJCON 2024

6th ANNUAL NATIONAL MULTIDISCIPLINARY CONFERENCE OF
ASSOCIATION FOR MEDICAL UPDATES



NATIONAL WORKSHOP ON GOOD CLINICAL PRACTICE GUIDELINES

22

NOVEMBER
2024

LT-1 NEW ACADEMIC BLOCK
SMS MEDICAL COLLEGE, JAIPUR

HIGHLIGHTS OF WORKSHOP

1. Updated Guidelines on GCP as per CDSCO
2. Credit hours by RMC
3. Fulfilling the NMC requirement of PG
4. Resource faculty key persons in framing GCP guidelines



Workshop Coordinator and Resource Faculty

Dr. Monica Jain

Senior Professor, Department of Pharmacology,
SMS Medical College, Jaipur



Master of Ceremony

Dr. Jaya Dadheech

Senior Demonstrator Department of Pharmacology
SMS Medical College, Jaipur

SPEAKERS



Prof. (Dr.) Bikash Medhi

Professor, Department of Pharmacology,
PGIMER, Chandigarh



Prof. (Dr.) Anant Patil

Professor, Department of Pharmacology,
Dr DY Patil Medical College, Navi Mumbai



Dr. Anusha Vohra

PHOD, Pharmacology,
MGMCH, Jaipur



Dr. Priyanka Rathi

Professor, Department of Pharmacology,
NIMS & R, Jaipur



Dr. Shivankan Kakkar

Assistant Professor, Pharmacology,
SMS Medical College, Jaipur

Scientific Agenda



ICH GCP Training Workshop

22nd November 2024

Time: 09:00 AM - 5:00 PM

Program Schedule

TIME	SESSION	FACULTY/SPEAKER
09:00 am 09:20 am	Registration & Networking Participant Check-In and Networking Opportunity	
09:20 am 09:40 am (20 min)	Inauguration Introduction to Workshop Objectives and Goals	
09:40 am 09:55 am (15 min)	Pre-Test Assessment Initial Knowledge Assessment of Participants	
BREAKFAST : 9:55 am – 10:10 am (15 min)		
10:10 am 10:50 am (40 min)	Keynote Session: Overview of Good Clinical Practice (GCP) The Importance of GCP in Clinical Trials and Research Integrity	Dr. Monica Jain
10:50 am 11:30 am (40 min)	Stakeholder Responsibilities in Clinical Research Detailed Roles of the Sponsor, Investigator, and Ethics Committee	Dr. Bikash Medhi
11:30 am 12:10 pm (40 min)	Ethics Committee: Structure and Function Composition, Roles, and Responsibilities of Ethics Committees	Dr. Anant Patil
12:10 pm 12:40 pm (30 min)	Informed Consent Process in Clinical Trials Comprehensive Overview of Consent Documentation and Process	Dr. Anant Patil Dr. Monica Jain
LUNCH : 12:40 pm – 1:15 pm (35 min)		
1:15 pm 1:45 pm (30 min)	Essential Documents for Conducting a Clinical Trial Protocols, Case Report Forms, and Investigator Brochures	Dr. Shivankan Kakkar
1:45 pm 2:15 pm (30 min)	Investigational Product Accountability Management and Accountability of Investigational Products	Dr. Anusha Vohara
2:15 pm 2:45 pm (30 min)	Adverse Event Reporting and Safety Monitoring Serious Adverse Event Reporting Requirements and Procedures	Dr. Monica Jain
2:45 pm 3:05 pm (20 min)	Principles of Academic Clinical Trials Design and Implementation of Academic Clinical Research	Dr. Anusha Vohara Dr. Monica Jain
TEA BREAK : 3:05 pm – 3:20 pm (15 min)		
3:20 pm 3:50 pm (30 min)	Quality Assurance, Audits, and Data Management Standards for Quality Control, Audits, and Data Archiving	Dr. Priyanka Rathi
3:50 pm 4:10 pm (20 min)	Group Activity GCP Compliance: Identifying Errors in Case Report Forms (CRFs)	Dr. Monica Jain Dr. Shivankan Kakkar
4:10 pm 4:30 pm (20 min)	Interactive Q&A Session Open Forum with Faculty on GCP Practices	
4:30 pm 4:45 pm (15 min)	Post-Test Assessment Final Knowledge Evaluation	
4:45 pm 5:00 pm (15 min)	Valedictory Ceremony and Feedback Collection Closing Remarks and Participant Feedback	