

ALL INDIA INSTITUTE OF AYURVEDA (AIIA), NEW DELHI

&

INSTITUTE OF APPLIED STATISTICS, KANPUR

In collaboration with

CENTRAL COUNCIL FOR RESEARCH IN  
AYURVEDIC SCIENCES (CCRAS), NEW DELHI

Jointly Presents

One-Month Online  
Certificate Programme on

# GOOD CLINICAL PRACTICE FOR AYURVEDIC MEDICINE

**Programme Duration:**

One-Month (1 to 30 April 2025)

**Programme Mode:**

Online & Recorded Session

**Live Session Time:**

06:00 PM - 07:00 PM

**Platform:**

**Live Session:** Cisco WebEx Meeting

**Recorded Session:** YouTube Private Link



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# CERTIFICATE PROGRAMME ON GOOD CLINICAL PRACTICE (GCP)



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# CERTIFICATE PROGRAMME ON GOOD CLINICAL PRACTICE (GCP)



## ABOUT THE CERTIFICATE PROGRAMME ON GCP

The Good Clinical Practice (GCP) certificate programme is designed to provide thorough training in the skills and knowledge necessary to conduct clinical research according to the International Committee for Harmonization of GCP guidelines (ICH GCP). This program aims to foster the development of multidisciplinary and multi-sectorial teams to advance clinical research. It also seeks to promote connections among trainees and other scientists from universities, research institutes, and similar institutions. By encouraging these collaborations, the program aims to create a supportive and empowering environment for participants, helping them become proficient professionals in clinical research. The one-month certificate program will be delivered online, featuring live and recorded sessions by the Institute of Applied Statistics, Kanpur, and the All India Institute of Ayurveda, New Delhi.

## OBJECTIVE

- Develop multidisciplinary and multi-sectorial teams dedicated to advancing clinical research.
- Foster and promote connections among trainees, scientists, and professionals from universities, research institutes, and similar establishments.
- Create a supportive and empowering environment for participants.
- Enhance participants' proficiency and capabilities in the realm of clinical research.
- Facilitate collaboration and growth among clinical research professionals.
- Ensure participants are well-versed in ethical, regulatory, and methodological aspects of clinical research.

## WHO SHOULD ATTEND ?

Faculty members or students with degrees in MD/MS, life sciences, B.Tech, B.Sc. in microbiology, life sciences, botany, zoology, food science, food technology, BE, B.Pharm, M.Pharm, Pharm D, MBBS, BDS, BHS, BUMS, BAMS, or BSMS, and who have consistently demonstrated strong academic and research backgrounds.

## SESSIONS OFFERED ON PLATFORMS



**REGISTRATION FEE:**  
**1500/- INR Per Participant**

## REGISTRATION:

For registration please visit our official website: [www.iasdcs.com](http://www.iasdcs.com)

- E-Certificates will be provided to all the participants.
- Access of Recorded Session and PDF Notes will be provided.

## PROGRAMME DURATION

01 April to 30 April, 2025



## LIVE SESSION TIME

06:00 PM - 07:00 PM

# CERTIFICATE PROGRAMME ON GOOD CLINICAL PRACTICE (GCP) PROGRAMME SCHEDULE



## 01 APRIL 2025 - INAUGURAL OF THE PROGRAMME

### 02 April Introduction to Good Clinical Practice (GCP)

- Overview of GCP: Definition, history, and significance
- Importance of GCP in clinical research
- Introduction to Ayurveda and its integration with modern clinical practices

### 03 April Ethical Principles and Guidelines

- Ethical principles in clinical research
- Types of Ethical Review and Requirement
- International and national guidelines (ICH-GCP, WHO-GCP)
- Relevance of ethics in Ayurveda research

### 04 April Regulatory Requirements

- Regulatory bodies (FDA, EMA, DCGI)
- Key regulations and guidelines for Ayurveda research
- Compliance with national and international regulations

### 07 April Roles and Responsibilities in Clinical Trials

- Responsibilities of sponsors, investigators, and ethics committees
- Roles specific to Ayurveda clinical trials
- Importance of delegation and documentation

### 08 April Study Protocol Development

- Components of a clinical trial protocol
- Designing a protocol for Ayurveda research
- Ensuring GCP compliance in protocol design

### 09 April Informed Consent Process

- Principles of informed consent
- Developing consent forms for Ayurveda studies
- Conducting the informed consent process ethically

### 10 April Investigator Brochure and Study Documentation

- Purpose and content of the Investigator Brochure
- Essential documents in clinical trials
- Document management in Ayurveda research

# CERTIFICATE PROGRAMME ON GOOD CLINICAL PRACTICE (GCP)



**15 April**

## **Study Design and Methodology**

- Types of clinical study designs
- Designing studies specific to Ayurveda treatments
- Ensuring methodological rigor

**16 April**

## **Patient Recruitment and Retention**

- Strategies for patient recruitment in Ayurveda trials
- Ensuring patient retention and adherence
- Ethical considerations in recruitment

**17 April**

## **Data Management and Record Keeping**

- Data management practices in clinical trials
- Electronic data capture systems
- Record keeping requirements for Ayurveda research

**19 April**

## **Monitoring and Quality Assurance**

- Principles of clinical trial monitoring
- Conducting monitoring visits
- Quality assurance processes in Ayurveda research

**21 April**

## **Adverse Event Reporting**

- Identifying and reporting adverse events
- Safety monitoring in clinical trials
- Managing adverse events in Ayurveda studies

**22 April**

## **Statistical Considerations**

- Basics of biostatistics in clinical research
- Statistical methods for Ayurveda trials
- Data analysis and interpretation

**23 April**

## **Clinical Study Reports**

- Structure and content of clinical study reports
- Writing and reviewing study reports
- Reporting outcomes for Ayurveda studies

**24 April**

## **Regulatory Submissions and Approvals**

- Preparing for regulatory submissions
- Approval processes for Ayurveda clinical trials
- Post-approval compliance



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**25 April**

## **Audits and Inspections**

- Conducting internal and external audits
- Preparing for regulatory inspections
- Responding to audit and inspection findings

**26 April**

## **Pharmacovigilance and Post-Marketing Surveillance**

- Principles of pharmacovigilance
- Post-marketing surveillance for Ayurveda products
- Integrating pharmacovigilance with GCP

**28 April**

## **Quality Management Systems**

- Developing a quality management system (QMS)
- Implementing QMS in Ayurveda research
- Continuous quality improvement

**29 April**

## **Integrating Traditional Knowledge with Modern Research**

- Challenges and opportunities in Ayurveda research
- Bridging traditional knowledge and modern clinical research
- Case studies and best practices

**30 April**

## **Future Trends and Innovations in GCP for Ayurveda**

- Emerging trends in clinical research
- Innovations in Ayurveda clinical trials
- The future of GCP in Ayurveda research

**03 MAY 2025 - ONLINE ASSESSMENT (MCQ BASED)**

**09 -10 MAY 2025 - E-CERTIFICATION**



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# CERTIFICATE PROGRAMME ON GOOD CLINICAL PRACTICE (GCP)



## CERTIFICATE PROGRAMME HIGHLIGHTS

- ✓ Comprehensive one month (Monday to Friday) online sessions on Good Clinical Practice (GCP).
- ✓ Expert-led sessions and daily query discussions at the end of each session of the programme to address doubts with experts.
- ✓ Online Assessment Tests at the end of the programme for E-Certification.
- ✓ Weekly Online Quiz for revision of the topics learned.
- ✓ Study material in the form of PDFs will be shared with all registered participants of the programme.
- ✓ All registered participants will receive an e-certificate after completion of the programme on their registered email ID.
- ✓ Access to all recorded sessions of the programme for revision purposes.
- ✓ Opportunity to network with researchers and professionals.

