



# START YOUR CAREER AS A PRO with

**cliantha**  
**academy**

(An Education Initiative of Cliantha Research)

Create, Learn, Inspire with Cliantha Academy  
– A Journey of Excellence

[cliantha.com](https://cliantha.com)

# Certificate Course on Clinical Research

A Gateway of Clinical Research industry  
(Instructor-led vocational training)

Eligibility: M.Sc. (Chemistry/Microbiology/Life science)/B.Pharm/M.Pharm/Pharm D  
/BPT/BHMS/BUMS/BAMS or any other discipline

## Course Objectives



To bridge the gap between industry expectations and academic outcomes through vocational training



Aim to create skilled professionals with practical skills needed for a successful career



We aim to upskill the freshers by engraving the real-time practical abilities, enhancing their absorption towards the clinical research industry.

## Why? Cliantha Academy

Cliantha Research a full-service Clinical Research Organization (CRO), provides comprehensive and integrated offerings in Early Phase (BA-BE & Phase I), Late Phase (II-IV), Respiratory, Tobacco Research, Dermatology, Consumer Research, Research Lab, IVRT, Biometrics, Medical Services, Environmental Exposure Chambers (EECs).

Cliantha, by leveraging on the depth of experience provides ethical, value added customer centric services rooted in excellence, thereby enabling clinical development of pharmaceutical, biotech & healthcare companies.

Cliantha Research is headquartered in Ahmedabad and has expanded its domestic reach with 3 regional offices & has presence in USA (Florida & New Jersey), Canada (Mississauga, Winnipeg & Scarborough) & Europe (Portugal).

- >18 years of clinical research experience
- Trained >2000 clinical research professional (site staff, EC, sponsors etc.)
- Extensive & intense on-site training
- Industry experts as faculty
- Creative/Interactive content
- Placement Assistance & career counseling

# Modules

The program is designed by industry experts for students and young professionals. The training provides in-depth knowledge of Clinical Research including a theoretical aspect of the field and on site practical training.

- History of Clinical Research and Good Clinical Practice (ICH E6R2)
- New Drug and Clinical Trial Rules and Good Documentation Practice (GDP)
- Workflow of BE Study (Healthy Volunteer) and patient based Clinical Trial
- Clinical trial study design, Protocol, and Essential documents list
- AE/SAE Reporting and Overview of Protocol and Amendments
- Source Data Review & Verification Practical/Simulation
- Orientation on various regulatory Guidelines (e.g., DCGI, EMA, USFDA, NPRA, MHRA, etc.)
- CRO/SMO/Recruitment techniques/Retention of patients/Central lab, Local lab
- Management of Different clinical trial sites,
- Sites discrepancy management, EDC, IVRS, IWRS,
- Monitoring and Safety Reporting, Monitoring Visit Reports
- Monitoring Informed Consent documents and procedure
- Monitoring Investigational Product (IP) Accountability
- The Monitoring Role in Preventing Inspection Findings
- Regulatory Compliance and Quality Assurance: Audits and Inspections

**Duration**  
**6 months**

**Fees**  
**45,000**  
**INR +**  
**GST**

## **Additional Benefits on any Course:**

- Personality development, public speaking and communication workshops
- Email Etiquette and skill development program
- Use of social media for career development
- Interview preparation

Clinical Research Coordinator (CRC), Clinical Trial Associate (CTA), Clinical Research Associate (CRA), Quality Control (QC), Quality Assurance (QA), Clinical Data Management, Ethics Committee Coordinator, Centralized Monitor, Third Party Monitor, Medical Writer, Regulatory Affairs, Report Writer, Project Manager, Cost and Proposal, Trainer, Study Start-Up Associate and many more.

**Various  
job roles  
in  
Clinical  
Research**

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# Glimpse of Cliantha Academy



**Cliantha Academy Facility**



**Expert Sessions**



**Clinical Trials Site Visits**



**Interview preparation & Group Discussion**



**Role play activity**



**Soft Skills Training**



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