राजकीय आयुर्विज्ञान संस्थान
Government Institute of Medical Sciences
GIMS, GR. Noida

AN AUTONOMOUS INSTITUTE UNDER GOVT. OF UTTAR PRADESH

6-MONTHS HANDS -ON
CLINICAL TRIALS TRAINING PROGRAM

Bridge the gap between classroom learning and practical exposure. Join India's emerging clinical research ecosystem at a premier Government Medical Institute.

PROGRAM HIGHLIGHTS

DURATION
6 Months

BATCH SIZE
5-10 Participants

MODE
Classroom Training, Practical Learning, Hands-On Experience on Live Projects .

VENUE
GIMS CR Unit

WHO CAN APPLY

- MBBS/BDS
- Pharmacy
- Life Sciences
- B.Sc/M.Sc Nursing
- Biotech
- BAMS/BHMS
- Allied Health

*Final year students eligible

THE INDUSTRY

Why Clinical Research?

India is rapidly becoming a global hub for clinical trials. There is a massive demand for professionals strictly trained in GCP, ethics, and Indian regulatory mandates (NDCT Rules).

- ✓ High growth career trajectory
- ✓ Contribute to life-saving drug development
- ✓ Global opportunities and recognition

THE INSTITUTE

Why Choose GIMS?

- Government Medical Institute
- Active Clinical Research Unit
- Ongoing National/Intl. Studies
- Training by Expert Investigators

COMPREHENSIVE CURRICULUM

8 CORE MODULES

MODULE 1
Intro to Clinical Research
- Drug Development Phases
- ICH-GCP Guidelines
- Key Stakeholders

MODULE 2
Ethics & Regulations
- IEC & Informed Consent
- NDCT Rules & CDSCO
- Participant Rights

MODULE 3
Trial Start-Up
- Site Feasibility
- Budgeting & Contracts
- Study Initiation

MODULE 4
Clinical Operations
- Recruitment & Enrollment
- CRF/eCRF Completion
- IP Accountability

MODULE 5
Safety Reporting
- AE / SAE Definitions
- SUSAR Reporting
- Regulatory Timelines

MODULE 6
Data Management
- Source Data Verification
- Query Resolution
- Quality Assurance

MODULE 7
Essential Docs
- Investigator Site File (ISF)
- Delegation Logs
- CAPA Preparation

MODULE 8
Live Trial Exposure
- **EC Submission Mocks**
- **Real Monitoring Visits**
- **Final Practical Project**

PRACTICAL EXPOSURE

Our program combines academic learning with real-world clinical trial experience, giving participants practical exposure under the mentorship of leading investigators and industry professionals.

REGULATORY & START-UP

Experience site feasibility assessments, essential document preparation, and Ethics Committee submissions.

PATIENT MANAGEMENT

Learn practical participant screening, administering informed consent, enrollment, and retention strategies.

DATA & SAFETY

Hands-on CRF/eCRF completion, source document verification, query resolution, and live SAE reporting.

TMF MANAGEMENT

Maintain the Trial Master File (TMF), organize investigator meetings, and coordinate with stakeholders.

MONITORING & AUDITS

Support monitoring visits, ensure audit/inspection readiness, and learn CAPA implementation.

LIVE OBSERVERSHIP

Observe and learn from live ongoing clinical research projects directly at the GIMS clinical unit.

CAREER OPPORTUNITIES

- Clinical Research Coordinator (CRC)
- Clinical Trial Associate (CTA)
- Clinical Research Associate (CRA)
- Clinical Data Coordinator
- Regulatory Affairs Executive
- Pharmacovigilance Associate
- Quality Assurance Executive

CERTIFICATE OF COMPLETION

Issued by GIMS, recognized in the industry. Awarded post successful completion of modules and attendance.

YOUR INVESTMENT YIELDS

INDUSTRY READY

STRONG NETWORK

PRACTICAL SKILLS

CONTACT & REGISTRATION

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SCAN TO REGISTER

Limited seats available.
Register today.