Global Institute Of Health Science™

CERTIFICATE · PG DIPLOMA · FAST-TRACK PG DIPLOMA

CLINICAL TRIAL MANAGEMENT

A rare combination in one programme — and one fee.

NEW AI MODULE | LIFETIME 1-TO-1 EXPERT | FREE CAREER COACHING ₹9,999 | 2 INTERNATIONAL CERTIFICATES | 100% ONLINE

NEW · AI MODULE **NOW WITH AI IN CLINICAL TRIAL MANAGEMENT & CLINICAL OPERATIONS**

What's inside this brochure

12 PAGES

01	**Programme Overview & Why It Stands Out** — What it covers, our teaching philosophy and the headline benefits	2
02	**Revised Syllabus — Module by Module** — All 9 modules, and what each one teaches	3–4
03	**NEW: Artificial Intelligence in Clinical Trial Management & Operations** — Key topics, practice exercises and the responsible boundary	5
04	**Study Method, Certification, Skills & Career Pathways** — How the course runs, what you gain and where it takes you	6
05	**Placement, Free Career Coaching & Enterprise Value** — Free career coaching worth ₹9,999, plus enterprise value	7
06	**SAMPLE CERTIFICATES** — The two international certificates awarded	8–9
07	**Fees, Admission Procedure & Who Can Enrol** — Learning options, how to apply and who can enrol	10
08	**Frequently Asked Questions** — Scope, eligibility, AI and certificate verification	11
09	**Other GIHS Programmes & How to Reach Us** — About the institute and the full catalogue	12

Trusted by Healthcare Professionals Globally

Learn from global perspectives and prepare for impactful roles in hospitals, public health systems and healthcare organisations around the world.

INDIA | UK | USA | AUSTRALIA | CANADA | RUSSIA | NEW ZEALAND | UAE | AFRICA | EUROPE

15+ years of experience **2,000+** placed globally **Lifetime** free expert tutoring

ACCREDITED & CERTIFIED

- **ISO 21001:2018** Educational organisation management
- **UASL** Germany / UK — quality management
- **GSAAA** USA — global accreditation
- **IPR, Govt. of India** Copyright approved

ENROL ANY DAY · ADMISSIONS OPEN ALL YEAR

Scan to enrol — or apply online

www.gihsonline.com/enroll-now/

info@gihsonline.com · +91-9033768588 · www.gihsonline.com

★★★★★

300+ Google reviews

Recorded candidate satisfaction to date

PROGRAMME OVERVIEW

Build Future-Ready Skills in Clinical Trial Management

Practical | Job-Oriented | ICH-GCP | Trial Operations | Project Management | Monitoring | AI Literacy

The GIHS Clinical Trial Management programme provides a broad, practical understanding of clinical trials, Good Clinical Practice, regulatory requirements, study planning, site management, monitoring, quality and project-management activities. The syllabus also gives exposure to pharmacovigilance and drug-safety concepts, for a wider view of participant protection and clinical-development operations.

Clinical trials are not clinical practice. In practice you use established treatments; in a trial you collect the evidence that establishes one. This programme is built around that distinction and everything it demands — ICH GCP, regulatory requirements and the operational discipline of running a study to time, budget and standard.

> **"We Teach What the Job Requires."**
>
> GIHS connects trial-management theory with the operational reality of clinical development — timelines, vendors, budgets, sites, monitoring, documentation and the quality systems that hold it all together.

Why This Programme Stands Out

- Clinical trial management from study planning through site close-out
- ICH-GCP, Schedule Y, ICMR and Indian GCP concepts
- Clinical trial design, project planning and project management
- Protocol, CRF, essential documents and informed-consent concepts
- Site selection, initiation, recruitment and retention planning
- Vendor management, contracts, budgeting and stakeholder coordination
- Clinical monitoring, source-data verification and drug accountability
- QC, compliance, auditing, SOPs and 21 CFR Part 11 awareness
- Drug-safety and pharmacovigilance exposure
- NEW dedicated AI module for trial management & clinical operations
- Certificate, PG Diploma and Fast-Track learning pathways

REVISED SYLLABUS

Curriculum, Module by Module

The full GIHS syllabus, regrouped into 9 clearly sequenced modules so you can see exactly what each stage of the programme teaches and why it comes where it does. Module 9 is the new Artificial Intelligence module.

MODULE 01 Foundations of Clinical Research

What clinical research is, how it differs from clinical practice, and how a medicine travels from the laboratory to the market.

- Introduction to clinical research
- Clinical trial phases
- Pharmacological principles of clinical research
- Drug development and launch

MODULE 02 Regulations & Key Stakeholders

The rulebook and the people bound by it — who is accountable for what, and to whom.

- Indian regulation (domestic regulation)
- ICH GCP / Schedule Y / ICMR / Indian GCP
- Key stakeholders in clinical research
- Ethics Committees and Institutional Review Board
- Responsibilities of the sponsor and the investigator
- Sponsor–investigator and sponsor–vendor responsibilities

MODULE 03 Trial Design & Project Management

Designing a study that can actually answer its question, then running it to time and budget.

- Clinical trial design
- Vendor selection and management
- Project planning
- Project management

MODULE 04 GCP, Protocol & Essential Documents

The paperwork that is the trial — if it is not documented to GCP standard, it did not happen.

- Principles of Good Clinical Practice
- Protocol design and CRF design
- Essential documents and regulatory requirements
- IND application and NDA application
- Clinical study report
- Informed consent process and documentation

MODULE 05 Study Setup & Site Management

Getting a site open and recruiting — the stage where most trials lose time.

- Site selection and pre-study visits
- Site initiation
- Subject recruitment and retention planning
- Site contracts and budgeting

MODULE 06 Quality Control, Compliance & Auditing

The systems that keep the data trustworthy and the trial inspection-ready.

- QC and QA in clinical research
- Compliance, auditing and inspections
- 21 CFR Part 11 and electronic records
- Site auditing
- Sponsor compliance and auditing
- SOPs for clinical research

MODULE 07 Clinical Monitoring

The CRA's working week — verifying at the site that the data match the source.

- Roles of the lead and principal investigator
- Clinical monitoring activities
- CRF review and source data verification
- Drug safety reporting
- Drug accountability work
- Routine site monitoring and site close-out visit

MODULE 08 Pharmacovigilance & Drug Safety Exposure

The safety side of a trial — enough drug-safety grounding to understand what the trial owes its participants and the regulator.

- Introduction and overview of pharmacovigilance
- Standard terms and terminology in PV
- Medical evaluation of adverse events; ADR diagnosis and management
- Case processing and case narrative writing
- PV database, signal detection and risk management
- MedDRA; PV laws, SOPs, auditing and inspection

MODULE 09 · NEW Artificial Intelligence in Clinical Trial Management & Operations

AI-Assisted Planning · Risk & Forecasting · Monitoring Support · GCP Accountability

- AI, Machine Learning, Generative AI and LLMs in clinical development
- AI across the trial lifecycle — planning, startup, conduct, monitoring, close-out
- AI-assisted protocol and study-document review
- … and 19 further topics — the full module is set out on page 5

The Programme at a Glance

9
Modules, including the new AI module

3 / 4 / 12
Months — Certificate, Fast Track or PG Diploma

100%
Online & distance, study at your own pace

Lifetime
Free personal expert assigned to you

MCQ + Project
Online exam plus a 30-page project

2
International certificates awarded and shipped

Every module is backed by a personal subject expert assigned to you for life at no cost — email or call whenever a topic needs explaining, and a session can be arranged around your queries.

MODULE 09 · NEW HIGHLIGHT MODULE

Artificial Intelligence in Clinical Trial Management & Operations

AI-Assisted Planning · Risk & Forecasting · Monitoring Support · GCP Accountability

This module introduces AI, Machine Learning and Generative AI across clinical-development operations — planning, feasibility, recruitment, forecasting, monitoring, documentation and quality. AI is taught as an assistive capability within a GCP environment, never as a replacement for accountable professional judgement.

Key Topics Covered

- AI, Machine Learning, Generative AI and LLMs in clinical development
- AI across the trial lifecycle — planning, startup, conduct, monitoring, close-out
- AI-assisted protocol and study-document review
- AI support for literature review and clinical-research knowledge management
- AI concepts for study feasibility and country/site selection
- AI-assisted patient recruitment and retention strategy
- AI for project planning, timeline forecasting and milestone-risk identification
- AI-assisted vendor-performance and operational KPI analysis
- AI concepts for trial budgeting, resource and workload forecasting
- AI support for Risk-Based Quality Management and risk-based monitoring
- AI-assisted review of data trends, anomalies and quality signals
- AI support for monitoring preparation, visit reports and follow-up documentation
- AI-assisted essential-document and Trial Master File review
- AI for audit and inspection-readiness checklists
- AI-assisted identification of missing, inconsistent or overdue documentation
- AI support for adverse-event extraction and safety-workflow prioritisation
- Generative AI for SOPs, meeting summaries, training content and checklists
- Prompt engineering for trial managers, CRAs and clinical operations
- AI hallucinations, fabricated citations and inaccurate regulatory information
- Participant privacy, confidential trial information and secure AI use
- Data integrity, audit trails, validation and 21 CFR Part 11 considerations
- Human-in-the-loop review and responsible AI governance in GCP environments

THE RESPONSIBLE BOUNDARY

Anyone can generate an answer with AI. The professional skill — and the one employers are now paying for — is knowing how to find what the AI got wrong.

AI is an assistive capability. Trial managers, investigators, monitors and sponsors remain accountable for GCP compliance, participant safety, source verification and every regulatory decision.

What This Module Adds to Your Career

Clinical trial management runs on volume — documents, timelines, risks, site data, communications and operational metrics. AI literacy helps professionals understand modern approaches to information review, forecasting, risk identification, monitoring support and workflow productivity, while keeping accountability exactly where GCP places it.

AI-Enhanced Practice Exercises

Applied work that builds the verification habit, not just the prompting habit:

- Build a feasibility and site-selection rationale from mock data
- Draft a timeline-risk register and test the assumptions behind it
- Review an AI-generated monitoring visit report against source
- Identify missing or overdue documents in a mock Trial Master File
- Prepare an inspection-readiness checklist for a study team

HOW YOU STUDY & WHERE IT TAKES YOU

Study Method, Certification, Skills & Career Pathways

Study Methodology

- Freelance, self-paced method of study around your work schedule.
- Study the books and course material provided with your enrollment.
- Note down your queries as you progress through each module.
- Contact your assigned expert by email or phone at any time.
- Experts respond by mail, by call, or arrange a session based on your queries.
- Lifetime expert support to keep raising your knowledge and expertise.
- All content is based on live cases and industry best practice.

Mode of Examination

- Login details for the online examination are issued on completion of the programme duration.
- A 7-day window is given to appear for the online examination.
- The examination pattern is Multiple Choice Questions (MCQ).
- A PowerPoint project of minimum 30 pages on a topic of your own choice.
- Successful candidates are awarded two certifications (Degree and Transcriptional Degree).

Learning Mode, Assessment & Certification

Mode	Online and distance learning, designed for working professionals.
Assessment	Online and distance examinations (MCQ / theoretical) and/or projects.
Certification	GIHS certificate awarded to successful candidates on completion of applicable requirements.
Career support	Placement support and CV-forwarding assistance for successful candidates.
Verification	Employers can confirm certificate authenticity through the official GIHS verification process.

Verified by GIHS itself — not by a third party. Every certificate carries its own certificate ID. An employer, university or registration body can confirm it directly at www.gihsonline.com/verify-certification — no third-party verification agency and no fee.

Practical & Job-Oriented Learning

The programme connects theory with the operational work of running a study: feasibility, startup, site contracts and budgets, vendor coordination, recruitment planning, monitoring visits, source-data verification, drug accountability, documentation and close-out.

Career-Relevant Skills You Will Develop

- Clinical trial lifecycle management
- ICH-GCP and regulatory compliance
- Trial design and protocol understanding
- Project planning and timeline management
- Vendor selection, contracts and budgeting
- Site management and recruitment planning
- Clinical monitoring and source-data verification
- Essential documents and TMF awareness
- QC, compliance and audit readiness
- Drug-safety and pharmacovigilance awareness
- AI literacy for clinical operations

Career & Professional Pathways

The current GIHS course page lists **Clinical Research Associate, Clinical Trial Assistant, Clinical Research Coordinator and clinical project-management pathways**. The programme can also support:

- Clinical Research Associate (CRA)
- Clinical Trial Assistant (CTA)
- Clinical Research Coordinator (CRC)
- Clinical operations support
- Clinical trial / study management support
- Assistant Clinical Project Manager
- Clinical Project Manager pathway
- Site management and monitoring roles
- Clinical trial quality & compliance support

Management and senior clinical-research positions generally require relevant practical experience in addition to education. GIHS professional education supports skill development and career progression rather than guaranteeing a specific job title, and eligibility depends on prior qualification, competency and employer requirements.

PLACEMENT & CAREER ENHANCEMENT

What GIHS Does for Your Career After the Course

CAREER ENHANCEMENT INCLUDED FREE WITH EVERY ENROLMENT

Exclusive FREE CV & Interview Coaching — worth ₹9,999

FREE
worth ₹9,999

Join a GIHS programme now and unlock complimentary access to TopCV Coach, giving you the professional resume and interview skills you need to launch your career with confidence. Your profile is also posted on the TopCV Coach website, where a large recruiter and employer network is active.

Professional resume building
Build a CV that reads the way recruiters in healthcare actually screen.

Interview skills coaching
Practise and sharpen how you present your qualification and your new AI literacy.

Profile posted on TopCV
Your profile goes live on topcvcoach.com, in front of a large employer network.

No extra cost
Worth ₹9,999, included free with your GIHS enrolment.

TopCV Interview — Delivered with **www.topcvcoach.com** · included free with every GIHS programme enrolment

Where Our Graduates Are Making an Impact

Clinical research is a global industry, and GIHS learners work right across it — in pharmaceutical and biotech companies, CROs, trial sites, regulatory bodies, medical devices, healthcare IT and academic research.

Pharmaceutical companies
Clinical operations, drug safety and regulatory teams

Biotechnology & biosimilars
Development programmes and regulatory submissions

Contract Research Organisations
Monitoring, data management and safety services

Hospitals & trial sites
Coordination, documentation and ethics-committee support

Medical devices & diagnostics
Clinical evaluation, regulatory and quality functions

Healthcare IT & analytics
Clinical data platforms, SAS programming and reporting

Regulatory & consulting firms
Submissions, audits, IPR and compliance services

Academic & government research
Institutional trials, registries and regulatory bodies

Independent practice & ventures
Writing, data, PV and regulatory support services

WORKING GLOBALLY — India · UK · USA · Canada · Australia · New Zealand · UAE & Gulf · Saudi Arabia · Qatar · Oman · Russia · Germany · Ireland · Singapore · Malaysia · Africa · Europe

Placement support. More than 2,000 GIHS professionals are now working in well-known organisations worldwide, many of them after building CV and interview skills through the free TopCV Coach programme. GIHS shares your profile globally for career growth. Successful candidates receive placement support and their CVs may be forwarded to various organisations. Final employment remains subject to employer selection and role-specific eligibility.

Business & Entrepreneurship Value

Trial operations run on outsourced capacity — site management organisations, feasibility consultants, vendor coordinators and project-management specialists — and almost all of them started as small teams with operational expertise.

- Set up or manage a site management / clinical trial support organisation
- Offer feasibility, site-selection and recruitment-planning consulting
- Provide trial project management, timeline and vendor coordination services
- Build trial budgeting, contracting and investigator-payment support
- Deliver clinical operations training for new research staff

Any regulated clinical service must only be provided within your lawful professional scope and applicable registration requirements. Career titles, eligibility and the right to practise depend on your prior qualification, experience, employer criteria and local regulation.

Global Institute Of Health Science

Certificate

This is to certify that

CANDIDATE NAME

completed the one year programme of

Post Graduate Diploma in

Clinical Trials Management

offered by

Global Institute of Health Science.

during MONTH YEAR to MONTH YEAR

Upon successful completion of the programme

POST GRADUATE DIPLOMA

IN

CLINICAL TRIALS MANAGEMENT

is awarded to him/her this day the XX MONTH YEAR.

Copyright Office
Government of India

MHRD (Higher Education)
Reg No: SW - 12032/2018
CR Act 1957

UASL

ISO 21001:2018

Cert ID :gihs-XXXX-XXXXX

Academic Head

Verify Certification: https://www.gihsonline.com/verify-certification/ | info@gihsonline.com |+91- 9033768588
INDIA | UK | AUSTRALIA | RUSSIA | CANADA | USA | NEW-ZEALAND | UAE | AFRICA

Global Institute Of Health Science

POST GRADUATE DIPLOMA
IN
Clinical Trials Management

MONTH YEAR to MONTH YEAR

CANDIDATE NAME

GRADE SHEET & TRANSCRIPT

Introduction | Clinical Trial Phases | Principal | Drug Development And Launch | ICH GCP | Schedule Y
Indian Regulation (domestic regulation) | ICMR | Indian GCP | Key Stakeholders
Ethics Committees And Institutional Review Board | Vendor Selection And Managements
Responsibilities Of Sponsor, Investigator, Sponsor – Investigator & Sponsor – Vendor
Clinical Trial Design And Project Managements & Planning | Principles of good clinical practices
Protocol Design | CRF Design | Essential documents | Clinical Study Report | NDA Application
Informed Consent process And Documentation | Study setup process | Site Selection And Pre- Study VIsits
Subject Recruitment And Retention Planning | Site Contract And Budgeting | 21 CRF Part 11 | Site Auditing
QC, Compliance & Auditing in Clinical Research | Sponsor Compliance And Auditing | Clinical Monitoring
CRF Review & Source Data Verification | Drug Safety Reporting | Drug Accountability Work
Routine Site Monitoring | Site Close Out Visit
Overview & Standard Terms And Terminology | Medical Evaluation Of Adverse Events
Adverse Event Reporting System And Form | Diagnosis And Managements Of ADRs
Medical Evaluation Of AE | Case Processing |Global Perspective | Single Case Processing
Case Narrative Writing | Reporting Database, Signal Detection | Managements And Risk Assessments & Evaluation
Quality System In PV | Expedited Reporting Criteria | PSUR & PBRER | Database And Signal Detection
Risk Assessments & Management | Medical Dictionary For Regulatory Activities medDRA
PV laws And Guideline | Regulatory Guideline @ Laws | SOPS | PV Auditing & Inspection | Regulatory Aspects

Examination: 41/50 **Project Work: 43/50**

Total: 84/100

GRADE: A+

UASL QUALITY MANAGEMENT SYSTEM

Quality Control Certification

ISO 21001:2018

Copyright Office Government of India

MHRD (Higher Education)
Reg No: SW - 12032/2018
CR Act 1957

Academic Head

XX/XX/XXXX

Cert. ID :gihs-XXXX-XXXXX

Verify Certification: https://www.gihsonline.com/verify-certification/ | info@gihsonline.com |+91- 9033768588
INDIA | UK | AUSTRALIA | RUSSIA | CANADA | USA | NEW-ZEALAND | UAE | AFRICA

ENROLMENT

Three Flexible Learning Options

Programme	Duration	Indian Fee	International	Positioning
Certificate Program	3 Months	**₹22,999**	**450 USD**	Basic foundation programme — core knowledge first
Post Graduate Diploma Program	1 Year	**₹32,999**	**699 USD**	Advanced and career-oriented — full depth at a steady pace
Fast Track PG Diploma Program MOST CHOSEN	4 Months	**₹36,999**	**750 USD**	Same advanced syllabus — flexible, finish at your own pace

Both advanced options carry the same career-oriented syllabus and the same two international certificates. The Fast Track is simply flexible — finish as quickly as your own pace allows and step into a fast-growing job market sooner. The Certificate Program is the basic, foundation-level option.

Printed books are optional
Online study material is included with every enrolment. Add the printed books at enrolment if you would like them shipped.

₹2,999 additional (India) | **99 USD** additional (International)

EMI available on credit cards Convert your fee into easy monthly instalments — ask admissions when you enrol.

All modes of payment are available. Fees may be subject to applicable taxes, exchange rates or future revision.

What Our Candidates Say

★★★★★ 300+ Google reviews

"I recently completed my Post Graduation in Pregnancy and Child Care from GIHS and it was a wonderful experience."
Dr Fatema Ebrahim

"One of the best institutions... Course is carefully structured and detailed. Management is very well organised and prompt in assistance."
A V

"This course has been a game-changer for me, providing the knowledge and skills necessary to succeed on a global platform."
Akram Sabiq

"They have a very good independent study methodology and the subject matter is really good... the Institute of Health was a wise choice."
Andrea Del Pilar

"From the day 1 to till date Chethan assisted very well... the studies are very detailed and made interesting."
Sarah Lilly, MBA

Admission Procedure START ANY DAY

1. **Apply online — any day**
   Enrol at www.gihsonline.com / Apply Online. Admissions stay open all year — no batch to wait for.
2. **Confirm your fee**
   No hidden charges. Payment within 7 days of enrollment.
3. **Receive your study material**
   Online access within 48 hours; printed books shipped if you add them.
4. **Get your personal expert**
   A professional subject expert is assigned to you for lifetime support at no cost.
5. **Complete exam and project**
   Online examination plus a project submission.
6. **Receive your certificates**
   Two international certificates are awarded and shipped.

Who Can Enrol

LISTED ON THE GIHS COURSE PAGE

- Science Graduates
- MBBS / BDS / BHMS / BAMS / BPT / BUMS
- B.Pharm / M.Pharm / Pharmacy Professionals
- Occupational Therapists
- B.Sc. / M.Sc. & Life-Science Graduates
- Medical IT Professionals
- Medical Laboratory Professionals
- Nursing Professionals
- Microbiology Professionals
- Biomedical / Biotechnology Professionals
- Clinical Research / CRO / Pharma Professionals
- Professionals working with hospitals & healthcare organisations

ALSO OPEN TO

- Any graduate from any discipline
- Working professionals with relevant work experience
- Entrepreneurs planning to start a healthcare or allied venture
- Managers and administrators running a healthcare organisation

Any graduate, and any professional with relevant work experience, may enrol — including those who intend to start, run or manage a healthcare or allied organisation.

QUESTIONS

Frequently Asked Questions

How is clinical trial management different from clinical practice?

In practice you use established treatments. In a trial you collect the evidence that establishes one — which brings protocol, GCP, regulatory and documentation obligations that ordinary practice does not carry.

Does the programme cover project management as well as GCP?

Yes. Trial design, project planning, project management, vendor selection, contracts and budgeting are all part of the curriculum.

Does it include pharmacovigilance?

Yes. The syllabus gives drug-safety exposure — AE evaluation, case processing, signal detection, MedDRA and PV laws — for a fuller view of participant protection.

How can AI help a clinical trial manager?

With feasibility, recruitment planning, timeline forecasting, vendor KPIs, risk identification, document review and monitoring support. Accountability for GCP compliance stays with qualified professionals.

Does it cover the Trial Master File?

Yes — essential documents and TMF review concepts, including identifying missing, inconsistent or overdue documentation.

Does the course cover monitoring and source-data verification?

Yes. CRF review, source-data verification, drug accountability, routine monitoring and site close-out are covered in a dedicated module.

Can this programme lead to a Clinical Project Manager role?

It supports the pathway. Project-management roles normally also require demonstrated operational experience, and appointment remains an employer decision.

Can working professionals enrol?

Yes. The programme is delivered online and by distance learning, with a Fast-Track option.

What is included in the programme fee?

Course material with the latest updates, online access within 48 hours, printed books shipped with your enrolment, a personal subject expert for life at no cost, the online examination and project assessment, and two international certificates on successful completion. There are no hidden charges.

Is the TopCV Coach access really free?

Yes. Every GIHS enrolment now includes complimentary TopCV Coach access worth ₹9,999 for professional resume building and interview skills, and your profile is posted on the TopCV website where a large recruiter network is active.

Can I use this programme to start my own practice or venture?

Yes, within your lawful professional scope. The Business & Entrepreneurship section of this brochure sets out the non-clinical services, education and consulting work the knowledge supports. Regulated clinical services require the appropriate qualification and authorisation.

I am not from a clinical background. Can I still enrol?

Yes. Any graduate from any discipline may enrol, as can working professionals with relevant experience, entrepreneurs and managers of healthcare or allied organisations.

No batch to wait for — start the day you enrol Admissions stay open every day of the year — enrol today and begin at once.

ABOUT THE INSTITUTE

Global Institute of Health Science

Global Institute of Health Science is a pioneer in health science education, especially in distance and online learning, with more than 15 years of experience and international exposure. GIHS is an ISO 21001:2018 certified health science institute, globally acclaimed, and certified with copyright approval from IPR, Government of India. GIHS also holds international certification from UASL UK for its Quality Control Management System.

- **Excellence for health professionals** — Programmes built around what healthcare employers actually ask for.
- **15+ years with international exposure** — A student and professional base across India, UK, USA, Canada, UAE, Russia and Australia.
- **ISO certified and government approved** — ISO 21001:2018, IPR Government of India copyright approval and UASL UK quality certification.
- **Lifetime free personalised tutoring** — A dedicated subject expert assigned to you at no cost, for life.

More Ways to Build Your Career

GIHS programmes help professionals get a strong start in the healthcare field. These courses are ad-hoc to a professional career and provide the additional benefit of being part of a global learning organisation.

Health Management Programs

- Health Informatics
- Hospital Administration
- Pharma Management
- Hospital Planning & Management
- Medical Transcription
- Nursing Management
- Public Health Administration
- Health Insurance
- Blood Banking Management
- Medico-Legal Systems
- Medical Laboratory Technology
- Medical Radiology Technology
- Quality Management for Hospitals
- Disaster Management for Hospitals
- Hospital Infection Control & Patient Safety
- Healthcare & Hospital Administration
- Medical Record Management
- Medical Scribe

Clinical Research Programs

- Clinical Research & Quality Assurance
- Pharmacovigilance & Medical Writing
- Clinical Research & Pharmacovigilance
- Clinical Data Management & SAS
- Regulatory Affairs, IPR & Patent
- Clinical Trials Management
- CRA & Site Management
- Study Start-Up & Site Activation
- Trial Supply & IP Management
- Biostatistics & Statistical Programming

Allied Medical Programs

- Mother & Child Health
- Mental Health & Addictions
- Nutrition & Dietetics
- Pregnancy Education
- Gynaecology & Obstetrics
- Electroencephalogram (EEG)
- Nuclear Medicine Technology
- Anaesthesia Management
- First Aid (CPR / Emergency Care)
- Electrocardiogram (ECG)

Environment, Health & Safety Programs

- Industrial Health & Hygiene
- Environment Health Safety & RM
- Occupational Health & Safety
- Industrial Safety
- Food Safety & Quality Management
- Oil & Gas Safety Engineering
- Process Safety & Engineering
- Fire & Industrial Safety Management
- Environment Social Governance (ESG)

Ayurveda & Homeopathic Programs

- Panchkarma: Ayurveda
- Garbhsanskar: Ayurveda
- Naturopathy & Yoga
- Siddha: Medicine
- Ksharsutra: Ayurveda
- Classical Homeopathy
- Materia Medica: Homeopathy
- Repertory: Homeopathy
- Organon: Homeopathy
- Ayurvedic Nutrition & Dietetics

Ready to enrol?

Apply online at https://www.gihsonline.com/enroll-now/ · Email info@gihsonline.com · Call +91-9033768588

Watch our programmes: youtube.com/@globalinstituteofhealthscience · Verify a certificate: www.gihsonline.com/verify-certification

INDIA | UK | AUSTRALIA | RUSSIA | CANADA | USA | NEW ZEALAND | UAE | AFRICA

CHAT ON WHATSAPP