Global Institute Of Health Science™

CERTIFICATE · PG DIPLOMA · FAST-TRACK PG DIPLOMA

CLINICAL RESEARCH & QUALITY ASSURANCE — GCP AUDITS & INSPECTIONS

A rare combination in one programme — and one fee.

NEW · AI MODULE **NOW WITH AI IN CLINICAL RESEARCH & QUALITY ASSURANCE**

What's inside this brochure

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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

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★★★★★

300+ Google reviews

Recorded candidate satisfaction to date

PROGRAMME OVERVIEW

Build Future-Ready Skills in Clinical Research, GCP Quality & AI

Practical | Job-Oriented | ICH-GCP | Quality Assurance | Audits | Inspections | 21 CFR Part 11 | AI Literacy

The GIHS Clinical Research & Quality Assurance / GCP Audits & Inspections programme is a practical, job-oriented professional programme for science, pharmacy, medical, life-science and healthcare professionals seeking specialised knowledge in clinical trials, Good Clinical Practice, quality assurance, monitoring, audits, inspections and regulatory compliance.

An audit is a systematic and independent examination of trial-related activities and documents, to determine whether those activities were conducted — and the data recorded, analysed and accurately reported — according to the protocol, the sponsor's SOPs, GCP and the applicable regulatory requirements. This programme teaches you to carry out and to withstand that examination.

> **"We Teach What the Job Requires."**
>
> GIHS connects core clinical-research knowledge with real workplace requirements — trial operations, quality systems, documentation, monitoring, audit readiness, regulatory expectations and emerging digital tools.

Why This Programme Stands Out

- Clinical Research + Quality Assurance + GCP Audits & Inspections in one programme
- ICH-GCP and regulatory-compliance orientation throughout
- Protocol, CRF, essential-document and informed-consent concepts
- Site selection, initiation, monitoring and close-out workflows
- Sponsor, investigator, vendor and ethics-committee responsibilities
- 21 CFR Part 11 and electronic-record compliance awareness
- Audit preparation, inspection readiness and CAPA-oriented thinking
- Practical AI use cases for document review, quality signals and operations
- Career-oriented for CRA, CTA, QA, audit and project-management pathways
- NEW dedicated AI module for clinical research & quality assurance
- 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 GCP Audits & Regulatory Inspections

The specialist end of the programme — planning an audit, gathering evidence and getting an organisation ready for a regulator at the door.

- Quality assurance — GCP audits and inspections
- Audit types: site, sponsor, vendor, process and system
- Audit evidence and findings
- Corrective and preventive action (CAPA) thinking
- Regulatory inspection readiness

MODULE 09 · NEW Artificial Intelligence in Clinical Research & Quality Assurance

AI-Assisted Document Review · Quality Signals · Audit Readiness · Human Accountability

- Introduction to AI, Machine Learning, Generative AI and LLMs in clinical research
- AI across the clinical-trial lifecycle — planning, conduct, monitoring, quality, reporting
- AI-assisted protocol and study-document review
- … and 16 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 Research & Quality Assurance

AI-Assisted Document Review · Quality Signals · Audit Readiness · Human Accountability

AI is increasingly relevant to modern clinical-development workflows. This module introduces practical AI concepts and responsible use cases across clinical research, QA, GCP audits and inspection readiness. The focus is AI as an assistive tool — never a substitute for qualified professional judgement, validated systems or regulatory accountability.

Key Topics Covered

- Introduction to AI, Machine Learning, Generative AI and LLMs in clinical research
- AI across the clinical-trial lifecycle — planning, conduct, monitoring, quality, reporting
- AI-assisted protocol and study-document review
- AI support for literature review, research summarisation and knowledge management
- AI-assisted identification of document gaps, inconsistencies and quality signals
- AI in Risk-Based Quality Management (RBQM) and risk-based monitoring
- AI for clinical data review, trend detection and anomaly identification
- AI-assisted audit planning, checklist development and inspection-readiness support
- AI support for SOP, CAPA and quality-system documentation — with human review
- AI in trial operations — site feasibility, recruitment, retention and workflow
- AI applications in safety surveillance and signal support
- Generative AI for audit questions, training content, summaries and structured reports
- Prompt engineering for clinical research and QA professionals
- AI hallucinations, bias, explainability and limitations
- Human-in-the-loop review and professional accountability
- Data privacy, confidentiality, cybersecurity and protection of trial information
- Data integrity, traceability, audit trails and validation considerations
- Responsible AI use under GCP and applicable regulatory requirements
- Practical AI case studies for CRA, QA auditor, CTA and project teams

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 technology. Qualified professionals remain responsible for judgement, verification, participant protection, compliance and every decision within their role — and only approved, validated tools may be used where procedures or regulations require it.

What This Module Adds to Your Career

The module is designed to show where AI improves efficiency and where strict professional oversight remains essential. You learn to evaluate AI output critically, protect confidential trial information, recognise hallucinations and bias, document human review, and use only approved tools. The aim is AI literacy for clinical research — not training on one software product — so you can adapt as technology and regulatory expectations evolve.

AI-Enhanced Practice Exercises

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

- Review an AI-generated audit checklist for gaps and unsupported items
- Identify quality signals in a set of mock trial documents
- Draft a risk-based monitoring concept and test it against GCP
- Produce a structured inspection-readiness summary
- Compare human and AI-assisted essential-document review

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 focuses on workplace-relevant competencies: protocol and CRF understanding, essential-document review, site processes, monitoring, source-data verification, quality systems, audit planning, inspection readiness, SOPs and compliance. Learners are trained to ask the questions a quality professional asks — what is the requirement, what evidence demonstrates compliance, what is the risk, is the data reliable, is the process traceable, and what corrective action is appropriate?

Career-Relevant Skills You Will Develop

- Clinical trial operations across the lifecycle
- ICH-GCP and regulatory compliance
- Clinical quality assurance systems
- GCP audit concepts and audit evidence
- Regulatory inspection readiness
- Clinical monitoring and source-data verification
- Essential documents and trial documentation
- SOP and quality-system development
- 21 CFR Part 11, electronic records and data integrity
- AI-enabled clinical research literacy
- Risk-based quality thinking

Career & Professional Pathways

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

- Clinical Research Associate (CRA)
- Clinical Trial Assistant (CTA)
- Clinical Research Coordinator
- QA Auditor / Clinical QA Professional
- GCP quality & compliance support
- Clinical trial / project management pathways
- Site, sponsor and vendor quality coordination
- Audit & inspection readiness support
- Clinical operations & quality systems

Job title and seniority depend on your original qualification, relevant experience, demonstrated competencies and employer requirements. The programme supports professional upskilling and does not itself guarantee appointment to a particular designation. GCP and clinical-quality principles are internationally relevant, but candidates seeking work abroad must also meet destination-country employer, immigration and regulatory 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

Clinical quality work is one of the most outsourced functions in the industry — sponsors and sites both buy audit, SOP and inspection-readiness support from small independent specialists rather than carrying that capacity in-house.

- Offer independent GCP audit and inspection-readiness services to sites and sponsors
- Build an SOP, quality-manual and CAPA documentation practice for CROs and research sites
- Provide site quality-system setup and staff GCP training programmes
- Consult on vendor qualification and process-compliance reviews
- Develop quality-training content for clinical research teams

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

OF POST GRADUATE DIPLOMA

This is to certify that

CANDIDATE NAME

has successfully completed the one year programme of **Post Graduate Diploma in Clinical Research & Quality Assurance — GCP Audits & Inspections** offered by Global Institute of Health Science, during **MONTH YEAR to MONTH YEAR**. Upon successful completion of the programme

POST GRADUATE DIPLOMA

IN

CLINICAL RESEARCH & QUALITY ASSURANCE — GCP AUDITS & INSPECTIONS

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

GLOBAL INSTITUTE OF HEALTH SCIENCE · GIHS · ACADEMIC OFFICE

Academic Head

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Global Institute Of Health Science

STATEMENT OF RESULT & STUDY

Grade Sheet & Transcript

CANDIDATE NAME	PROGRAMME PERIOD	TOTAL LEARNING HOURS
CANDIDATE NAME	MONTH YEAR – MONTH YEAR	600 Hours

PROGRAMME

Post Graduate Diploma in Clinical Research & Quality Assurance — GCP Audits & Inspections

NO.	MODULE & TOPICS COVERED	HOURS	MAX	OBTAINED	%
01	**Foundations of Clinical Research** Introduction to clinical research; Clinical trial phases; Pharmacological principles of clinical research; Drug development and launch	54	50	48.5	97%
02	**Regulations & Key Stakeholders** 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	54	50	46.5	93%
03	**Trial Design & Project Management** Clinical trial design; Vendor selection and management; Project planning; Project management	54	50	48	96%
04	**GCP, Protocol & Essential Documents** 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	54	50	47	94%
05	**Study Setup & Site Management** Site selection and pre-study visits; Site initiation; Subject recruitment and retention planning; Site contracts and budgeting	54	50	47	94%
06	**Quality Control, Compliance & Auditing** 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	45	50	48	96%
07	**Clinical Monitoring** 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	45	50	48	96%
08	**GCP Audits & Regulatory Inspections** Quality assurance — GCP audits and inspections; Audit types: site, sponsor, vendor, process and system; Audit evidence and findings; Corrective and preventive action (CAPA) thinking; Regulatory inspection readiness	45	50	49	98%
09	**Artificial Intelligence in Clinical Research & Quality Assurance** NEW AI-Assisted Document Review · Quality Signals · Audit Readiness · Human Accountability; Introduction to AI, Machine Learning, Generative AI and LLMs in clinical research; AI across the clinical-trial lifecycle — planning, conduct, monitoring, quality, reporting; AI-assisted protocol and study-document review	45	50	46.5	93%
	Module Examinations — Sub-total	**450**	**450**	428.5	**95.2%**
10	**Project Work** 30-page applied project submitted at the end of the programme	150	200	185	92.5%
	GRAND TOTAL	**600**	**650**	**613.5**	**94.4%**

TOTAL MARKS	PERCENTAGE	GRADE	RESULT	DATE OF AWARD	CERT. ID
613.5 / 650	94.4%	A+	PASSED	XX-XX-XXXX	gihs-XXXX-XXXXX

Academic Head

MHRD (Higher Education) Reg No: SW - 12032/2018 · CR Act 1957 | ISO 21001:2018 · GSAAA & UASL

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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

What is clinical research quality assurance?

QA uses planned and systematic quality activities to support participant protection, GCP compliance, reliable data and adherence to protocols, SOPs and applicable regulations.

Why is AI included in a GCP and quality assurance course?

Clinical research increasingly uses digital and data-driven workflows. AI literacy helps professionals understand efficiency gains while recognising risks around privacy, bias, hallucinations, validation, traceability and regulatory compliance.

Does AI replace a QA auditor or CRA?

No. AI is assistive. Qualified professionals remain responsible for judgement, verification, participant protection, compliance and decisions within their roles.

Does the course cover 21 CFR Part 11?

Yes, as part of electronic-record, compliance and quality-system awareness, alongside data integrity and audit-trail concepts.

Does it cover clinical monitoring?

Yes — CRF review, source-data verification, drug-safety reporting, drug accountability, routine monitoring and site close-out.

Does the programme cover GCP audits and regulatory inspections?

Yes. GCP audits, inspections, site auditing, sponsor compliance, QA/QC and inspection readiness are major areas of the curriculum.

Can this programme help me become a CRA or QA auditor?

It can strengthen the knowledge relevant to CRA, CTA, QA, compliance and audit pathways. Actual appointment depends on your qualifications, experience and employer requirements.

Can working pharma, CRO or clinical-research professionals enrol?

Yes. The online and distance-learning format suits working professionals strengthening GCP, QA, audit and AI competencies.

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

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CHAT ON WHATSAPP