Global Institute Of Health Science™

CERTIFICATE · PG DIPLOMA · FAST-TRACK PG DIPLOMA

PHARMACOVIGILANCE & MEDICAL / SCIENTIFIC WRITING

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 PHARMACOVIGILANCE, DRUG SAFETY & MEDICAL WRITING**

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

ENROL ANY DAY · ADMISSIONS OPEN ALL YEAR

Scan to enrol — or apply online

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★★★★★

300+ Google reviews

Recorded candidate satisfaction to date

PROGRAMME OVERVIEW

Build Future-Ready Skills in Drug Safety & Scientific Communication

Practical | Job-Oriented | Drug Safety | Case Processing | Signal Detection | MedDRA | Medical Writing | AI Literacy

Drug safety is only as good as the writing that carries it. A case narrative, an aggregate report or a safety summary has to be accurate, complete, consistent with its source data and readable by someone who was not there. The GIHS Pharmacovigilance & Medical/Scientific Writing programme deliberately teaches both halves of that job.

The syllabus builds practical knowledge in adverse-event collection and assessment, case processing, safety reporting, signal detection, risk management, MedDRA, pharmacovigilance regulations and audits — and pairs it with medical and scientific writing: research and referencing, copyediting, data consistency across text and graphics, and compliance with applicable guidelines and standards.

> **"We Teach What the Job Requires."**
>
> GIHS focuses on professional skill development that complements a learner's academic background with industry-oriented knowledge, practical workflows and career-relevant competencies.

Why This Programme Stands Out

- Pharmacovigilance + Medical / Scientific Writing in one programme
- Adverse event and ADR assessment and reporting
- Individual case processing and case narrative writing
- PV databases, signal detection and risk management
- PSUR and PBRER concepts
- MedDRA terminology and coding awareness
- PV regulations, SOPs, audits and inspections
- Scientific research, referencing, editing and data consistency
- NEW dedicated AI module for drug safety and medical / scientific writing
- Certificate, PG Diploma and Fast-Track learning pathways

REVISED SYLLABUS

Curriculum, Module by Module

The full GIHS syllabus, regrouped into 8 clearly sequenced modules so you can see exactly what each stage of the programme teaches and why it comes where it does. Module 8 is the new Artificial Intelligence module.

MODULE 01 Foundations of Pharmacovigilance

Drug safety as a discipline — what it monitors, why it exists and the language it uses.

- Introduction to pharmacovigilance
- Overview of pharmacovigilance
- Standard terms and terminology in pharmacovigilance
- Global perspective of pharmacovigilance

MODULE 02 Adverse Events & ADR Evaluation

Recognising a reaction, assessing it medically and knowing how fast it has to be reported.

- Medical evaluation of adverse events
- Adverse event reporting systems and forms
- Diagnosis and management of ADRs
- Expedited reporting criteria

MODULE 03 Case Processing & Narrative Writing

Turning a reported event into a complete, coded, chronological safety case.

- Case processing workflow
- Single case processing
- Case narrative writing
- Quality systems in pharmacovigilance

MODULE 04 Safety Databases, Signal Detection & Risk Management

Moving from individual cases to a population-level view of a medicine's safety profile.

- PV reporting databases
- Signal detection and signal management
- Risk assessment, evaluation and management
- PSUR & PBRER

MODULE 05 MedDRA, PV Laws, SOPs & Inspections

The terminology standard and the regulatory framework behind every safety submission.

- Medical Dictionary for Regulatory Activities (MedDRA)
- Regulatory guidelines and laws in PV
- SOPs in pharmacovigilance
- PV auditing and inspection
- Regulatory aspects in pharmacovigilance

MODULE 06

Medical & Scientific Writing — Purpose & Applications

Who the writing is for, and what each audience needs it to do.

- Disease-management related scientific content
- Continuing education and training content
- Medical reference and information gathering
- Practice management and monitoring

MODULE 07

Editing, Accuracy & Compliance

The craft end — where clarity, consistency and compliance are actually earned.

- Correct spelling, grammar and punctuation
- Copyediting for accuracy, clarity and flow
- Correct and consistent data across text and graphics
- Compliance with applicable guidelines and standards

MODULE 08 · NEW

Artificial Intelligence in Pharmacovigilance & Medical / Scientific Writing

Safety Intelligence · Writing Productivity · Responsible Human Oversight

- AI, Machine Learning, NLP and Generative AI fundamentals for PV professionals
- AI across the PV lifecycle — intake, triage, processing, review, reporting, surveillance
- AI-assisted extraction of adverse-event information from structured and unstructured text
- … and 19 further topics — the full module is set out on page 5

The Programme at a Glance

8
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 08 · NEW HIGHLIGHT MODULE

Artificial Intelligence in Pharmacovigilance & Medical / Scientific Writing

Safety Intelligence · Writing Productivity · Responsible Human Oversight

This future-ready module introduces responsible applications of Artificial Intelligence and Generative AI in pharmacovigilance and medical / scientific writing. AI is positioned as an assistive technology for efficiency, review, research and structured drafting — not as a replacement for qualified safety assessment, medical judgement, regulatory accountability or professional authorship.

Key Topics Covered

- AI, Machine Learning, NLP and Generative AI fundamentals for PV professionals
- AI across the PV lifecycle — intake, triage, processing, review, reporting, surveillance
- AI-assisted extraction of adverse-event information from structured and unstructured text
- AI concepts for case intake, duplicate detection and case prioritisation
- AI-assisted case narrative drafting and quality review — with human verification
- AI support for MedDRA coding suggestions and terminology mapping
- AI-assisted literature screening and safety-information surveillance
- AI concepts for signal detection, trend identification and safety-pattern recognition
- AI support for risk-management documentation and aggregate-report workflows
- AI-assisted PSUR / PBRER organisation, summarisation and consistency review
- AI support for PV SOP, checklist, audit and inspection-readiness documentation
- Generative AI for outlines, first drafts, summaries and plain-language adaptation
- AI-assisted literature search strategy, evidence organisation and reference management
- AI for grammar, readability, consistency, terminology and data cross-checks
- AI-assisted comparison of text, tables, figures and source data
- Prompt engineering for pharmacovigilance and medical-writing professionals
- AI hallucinations, fabricated references and inaccurate medical claims
- Bias, explainability and the limits of AI-generated safety conclusions
- Patient privacy, confidential safety data, cybersecurity and secure AI use
- Data integrity, audit trails, traceability, validation and quality-system considerations
- Human-in-the-loop review and accountable medical / regulatory sign-off
- Responsible authorship, transparency and ethical use of AI in scientific communication

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 can draft; it cannot sign off. Safety assessment, medical judgement, coding decisions and regulatory submissions remain the responsibility of qualified professionals, and every claim and citation produced with AI must be verified against reliable source material.

What This Module Adds to Your Career

Pharmacovigilance and medical writing are information-intensive professions, so AI literacy translates directly into productivity in text processing, literature surveillance, case-review support, data consistency and structured drafting. The module also confronts the single biggest risk in AI-assisted medical writing — a fabricated citation or a plausible but scientifically wrong statement — and trains the verification habit that prevents it reaching a regulator.

AI-Enhanced Practice Exercises

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

- Evaluate an AI-generated case narrative against the source data
- Identify missing safety information in an AI-assisted summary
- Review an AI literature summary against the original source material
- Detect fabricated references in AI-generated scientific text
- Prepare a structured PV checklist with human verification points

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 workplace activities: adverse-event assessment, case-processing logic, narrative writing, MedDRA awareness, safety-reporting concepts, signal and risk-management thinking, SOP and audit readiness, literature research and medical-writing quality review. Learners are trained to write to a source, not from memory — and to make the trail from statement back to evidence visible to any reviewer.

Career-Relevant Skills You Will Develop

- Adverse-event and ADR terminology
- ICSR / single-case processing concepts
- Case narrative writing
- Safety reporting and expedited-reporting awareness
- PSUR / PBRER concepts
- PV database and signal-detection awareness
- Risk-management concepts
- MedDRA terminology
- PV quality systems, SOPs, audits and inspections
- Medical and scientific research and writing
- Editing, clarity and data consistency
- AI literacy for PV and scientific-writing workflows

Career & Professional Pathways

The current GIHS course page lists **Drug Safety Associate, Pharmacovigilance Associate and Medical / Scientific Writer pathways**. The programme can also support:

- Drug Safety Specialist
- Pharmacovigilance Associate
- Drug Safety Associate
- PV case-processing roles
- Safety operations / PV support
- Medical / Scientific Writer
- Drug safety and medical writing roles
- PV quality / compliance support
- Literature surveillance support
- Signal management support

Medical Monitor and other medically accountable roles normally require appropriate medical qualifications and employer-specific experience. GIHS professional education supports upskilling and does not replace statutory or employer qualification requirements; actual appointment depends on prior qualification, competencies, experience and employer criteria.

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

Medical and scientific writing is one of the clearest routes from employment to independent practice in the life sciences: the work is project-based, remote and priced on expertise rather than headcount.

- Build a freelance medical and scientific writing practice
- Offer case-narrative, aggregate-report and safety-document writing support
- Provide literature-surveillance and evidence-summary services
- Develop continuing-education, patient-information and training content
- Consult on editorial quality, data consistency and document compliance

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 Pharmacovigilance & Medical / Scientific Writing** offered by Global Institute of Health Science, during **MONTH YEAR to MONTH YEAR**. Upon successful completion of the programme

POST GRADUATE DIPLOMA

IN

PHARMACOVIGILANCE & MEDICAL / SCIENTIFIC WRITING

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

Academic Head

CERT. ID : **gihs-XXXX-XXXXX**

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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 Pharmacovigilance & Medical / Scientific Writing

NO.	MODULE & TOPICS COVERED	HOURS	MAX	OBTAINED	%
01	**Foundations of Pharmacovigilance** Introduction to pharmacovigilance; Overview of pharmacovigilance; Standard terms and terminology in pharmacovigilance; Global perspective of pharmacovigilance	63	50	48.5	97%
02	**Adverse Events & ADR Evaluation** Medical evaluation of adverse events; Adverse event reporting systems and forms; Diagnosis and management of ADRs; Expedited reporting criteria	63	50	46.5	93%
03	**Case Processing & Narrative Writing** Case processing workflow; Single case processing; Case narrative writing; Quality systems in pharmacovigilance	54	50	47.5	95%
04	**Safety Databases, Signal Detection & Risk Management** PV reporting databases; Signal detection and signal management; Risk assessment, evaluation and management; PSUR & PBRER	54	50	47	94%
05	**MedDRA, PV Laws, SOPs & Inspections** Medical Dictionary for Regulatory Activities (MedDRA); Regulatory guidelines and laws in PV; SOPs in pharmacovigilance; PV auditing and inspection; Regulatory aspects in pharmacovigilance	54	50	47	94%
06	**Medical & Scientific Writing — Purpose & Applications** Disease-management related scientific content; Continuing education and training content; Medical reference and information gathering; Practice management and monitoring	54	50	48	96%
07	**Editing, Accuracy & Compliance** Correct spelling, grammar and punctuation; Copyediting for accuracy, clarity and flow; Correct and consistent data across text and graphics; Compliance with applicable guidelines and standards	54	50	48	96%
08	**Artificial Intelligence in Pharmacovigilance & Medical / Scientific Writing** NEW Safety Intelligence · Writing Productivity · Responsible Human Oversight; AI, Machine Learning, NLP and Generative AI fundamentals for PV professionals; AI across the PV lifecycle — intake, triage, processing, review, reporting, surveillance; AI-assisted extraction of adverse-event information from structured and unstructured text	54	50	49	98%
	Module Examinations — Sub-total	**450**	**400**	381.5	**95.4%**
09	**Project Work** 30-page applied project submitted at the end of the programme	150	200	185	92.5%
	GRAND TOTAL	**600**	**600**	**566.5**	**94.4%**

TOTAL MARKS	PERCENTAGE	GRADE	RESULT	DATE OF AWARD	CERT. ID
566.5 / 600	94.4%	A+	PASSED	XX-XX-XXXX	gihs-XXXX-XXXXX

Academic Head

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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 pharmacovigilance?

Commonly called drug safety, it covers the collection, detection, assessment, understanding, monitoring and prevention of adverse effects and other medicine-related safety problems.

Why is medical writing taught alongside it?

Safety work is communicated in writing — narratives, aggregate reports, SOPs and summaries. Accuracy in the assessment is wasted if the document that carries it is unclear, inconsistent or unsupported.

Can AI write a case narrative?

It can produce a first draft or organise case information, but the narrative must be checked against source data for medical accuracy, completeness, chronology and regulatory compliance by a competent professional.

Can AI help with MedDRA coding?

It can suggest terminology or support the workflow, but final coding must follow current MedDRA conventions, organisational procedures and qualified human review.

What is the biggest risk of AI in medical writing?

Plausible but incorrect scientific statements and fabricated references. Every AI-assisted claim and citation should be verified against reliable source material.

Does it cover signal detection, PSUR/PBRER and MedDRA?

Yes. PV databases, signal detection, risk management, PSUR, PBRER and MedDRA are all specific parts of the syllabus.

Does it cover PV audits and inspections?

Yes — PV auditing and inspection, SOPs, quality systems and regulatory aspects.

Can working pharma or CRO professionals enrol?

Yes. The online and distance-learning format suits eligible working professionals seeking PV, drug-safety or medical-writing upskilling, with a Fast-Track option available.

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