Global Institute Of Health Science™

CERTIFICATE · PG DIPLOMA · FAST-TRACK PG DIPLOMA

REGULATORY AFFAIRS, IPR & PATENTS

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 REGULATORY AFFAIRS, INTELLECTUAL PROPERTY & PATENT INTELLIGENCE**

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

www.gihsonline.com/enroll-now/

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★★★★★

300+ Google reviews

Recorded candidate satisfaction to date

PROGRAMME OVERVIEW

Build Future-Ready Skills in Regulatory Affairs, IPR & Patents

Practical | Job-Oriented | USFDA | ICH | EMA / UK MHRA | CTD & DMF | IPR & Patents | AI Literacy

A medicine or device reaches a patient only after it has satisfied a regulator, and the organisation that owns it stays viable only if its intellectual property is protected. Both jobs are done by the same kind of professional — someone who can read a requirement precisely, assemble the evidence for it and defend that evidence under scrutiny.

The GIHS Regulatory Affairs, IPR & Patents programme covers Indian pharmaceutical regulation together with international regulatory concepts involving USFDA, ICH, EMA and European authorities, UK MHRA and WHO, and connects product development, pre-clinical and clinical requirements, submissions, quality operations and legislation with intellectual property, patents, trademarks and copyrights.

> **"We Teach What the Job Requires."**
>
> The programme complements academic qualifications with specialised, industry-oriented knowledge relevant to regulatory documentation, compliance, product registration, quality systems, IPR and global market requirements.

Why This Programme Stands Out

- Regulatory Affairs + IPR + Patents in one professional programme
- Indian and international pharmaceutical regulatory frameworks
- Pre-clinical and clinical regulatory considerations
- Drug, medical-device and cosmetics registration concepts
- Generic, biosimilar, herbal and orphan-product regulation
- Drug Master File and Common Technical Document concepts
- USFDA, ICH, EMA / European authorities, UK MHRA and WHO coverage
- Manufacturing-facility auditing and quality-assurance orientation
- NEW dedicated AI module for regulatory intelligence and patent research
- 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 Regulatory Foundations & the Indian Framework

Who regulates what in India, and the statutes every pharmaceutical professional works under.

- Pre-clinical trial and clinical trial requirements
- Regulatory bodies in India
- Central Drug Regulatory System
- Drug & Cosmetics Act and Schedule Y
- Medical device registration in India
- Product development protocol
- Environmental Protection Act, 1986

MODULE 02 Pharmaceutical Legislation in India

The wider body of law that shapes how a medicine may be made, priced, named and sold.

- Narcotic Drugs & Psychotropic Substances Act
- Medicinal & Toilet Preparations
- Drug Price Control Order
- Laws on trademarks and copyrights
- Prevention of Cruelty to Animals Act
- Consumer Protection Act, 1986

MODULE 03 The Pharmaceutical Regulation Process in India

How a product actually moves through registration — and what each market expects.

- Regulatory considerations for pre-clinical and clinical testing
- Regulation and registration of medical devices
- Regulation and registration of cosmetics
- General drug regulatory approach
- New-drug development procedures in India and other markets
- WHO certification concepts, import and export policy

MODULE 04 Generic, Biosimilar & Specialised Products

The product categories with their own rulebooks — and the dossiers that carry them.

- Regulation of generic pharmaceutical products
- Biosimilars in India and Europe
- Worldwide regulation of herbal products
- Introduction and regulation of orphan drugs
- Submission of the Drug Master File to USFDA
- Common Technical Document (CTD)
- Legal environment of business

MODULE 05 Global Regulatory Authorities & Submissions

The agencies whose expectations determine whether a product crosses a border.

- Introduction to USFDA
- ICH guidelines
- European Union authorities — EMA and UK MHRA
- WHO guidelines
- Guidelines for Rest of World markets
- Development of orphan drugs

MODULE 06 Quality, Auditing & Regulatory Projects

Proving in practice what the dossier claims on paper.

- Auditing of manufacturing facilities
- Quality assurance in regulatory operations
- Regulatory documentation and project work
- Compliance and inspection readiness

MODULE 07 IPR, Patents, Trademarks & Copyrights

Protecting the asset — what can be owned, for how long, and on what evidence.

- Introduction to Intellectual Property Rights
- Patents and patentability concepts
- Prior art and patent searching
- Trademarks and copyrights
- IPR in the pharmaceutical and biotechnology industry

MODULE 08 · NEW Artificial Intelligence in Regulatory Affairs, IPR & Patents

Regulatory Intelligence · Dossier Support · Patent Intelligence · Human Oversight

- AI, Machine Learning, Generative AI and LLMs for regulatory professionals
- AI-assisted global regulatory intelligence and monitoring of regulatory change
- AI support for searching, organising and comparing regulatory guidance
- … 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 Regulatory Affairs, IPR & Patents

Regulatory Intelligence · Dossier Support · Patent Intelligence · Human Oversight

This future-ready module introduces responsible applications of Artificial Intelligence and Generative AI across regulatory and intellectual-property workflows. AI is presented as an assistive capability for research, document review, intelligence and productivity — not as a replacement for regulatory professionals, patent specialists, legal counsel, validated systems or accountable human decision-making.

Key Topics Covered

- AI, Machine Learning, Generative AI and LLMs for regulatory professionals
- AI-assisted global regulatory intelligence and monitoring of regulatory change
- AI support for searching, organising and comparing regulatory guidance
- AI-assisted gap analysis between internal documents and regulatory requirements
- Generative AI for first-draft regulatory summaries, checklists and structured content
- AI support for CTD / eCTD content review and dossier-readiness concepts
- AI-assisted consistency checks across submission documents and labels
- AI for regulatory document classification, metadata and knowledge management
- AI-assisted inspection-readiness and compliance-checklist development
- AI support for regulatory risk identification and issue prioritisation
- AI in medical-device and pharmaceutical regulatory intelligence
- AI-assisted patent landscape and prior-art research concepts
- AI for patent classification, technology landscaping and competitor intelligence
- AI-assisted summarisation of patent claims and technical disclosures
- AI support for trademark, copyright and broader IPR research workflows
- Generative AI for invention-disclosure structuring and patent-drafting support
- Prompt engineering for regulatory affairs and IPR professionals
- AI hallucinations, inaccurate citations and fabricated regulatory or legal information
- Confidentiality, trade secrets, unpublished inventions and secure AI use
- Data integrity, traceability, version control, validation and auditability
- Human-in-the-loop review and accountable regulatory / legal decision-making
- Ethical and responsible AI governance in regulated industries

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.

Patentability, freedom-to-operate and regulatory conclusions require qualified professional review and authoritative searches. AI must never be treated as an autonomous regulatory decision-maker, and submission content must remain accurate, traceable, reviewed and compliant with applicable requirements.

What This Module Adds to Your Career

Regulatory and patent work is research- and document-intensive, so properly governed AI genuinely helps you search, compare, summarise and structure information faster. The module also teaches the failure modes that matter most here — hallucinated regulations, invented references, inaccurate patent interpretations, outdated guidance and confidentiality leakage of unpublished inventions — because in this field an unverified output can cost a filing date or an approval.

AI-Enhanced Practice Exercises

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

- Compare regulatory requirements across two markets and check the result against guidance
- Generate a first-pass regulatory checklist and audit it for unsupported items
- Review an AI-produced summary against the authoritative source
- Build a patent-landscape search framework
- Identify hallucinations in AI-generated regulatory content

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 regulatory theory with workplace activities: regulatory pathway comparison, guidance review, CTD and DMF awareness, compliance checklists, product-registration concepts, manufacturing-audit thinking, IPR research and structured regulatory documentation. The discipline being trained is precision — reading a requirement as written, and being able to show exactly which document satisfies it.

Career-Relevant Skills You Will Develop

- Regulatory intelligence and compliance awareness
- Indian and international regulatory frameworks
- Product registration and submission-document awareness
- CTD and DMF concepts
- Medical-device and cosmetics regulatory awareness
- Generic, biosimilar and orphan-drug regulatory concepts
- IPR, patent, trademark and copyright awareness
- Manufacturing-audit and quality-assurance concepts
- AI-assisted regulatory research and document-review literacy
- AI-assisted patent and IP intelligence with human verification

Career & Professional Pathways

The current GIHS course page lists **Regulatory Affairs Executive / Specialist, Regulatory Operations and IPR research pathways**. The programme can also support:

- Regulatory Affairs Specialist
- Regulatory Affairs Executive
- Regulatory Affairs Manager pathway
- Regulatory operations / submission support
- Regulatory compliance specialist
- Compliance officer
- Regulatory intelligence support
- IPR / patent research support
- Pharma, biotech and medical-device regulatory roles
- Quality and regulatory coordination

Senior, legal, patent-agent, attorney and formally licensed roles require specific education, professional registration, examinations or jurisdiction-specific authorisation. This programme develops professional regulatory and IPR knowledge; it does not confer patent-agent or attorney status, and appointment depends on prior qualification, experience 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

Regulatory and IP work is bought project by project — a registration, a dossier, a landscape search — which is precisely how a small consultancy can compete with a large department.

- Offer regulatory documentation, dossier and product-registration support
- Provide regulatory intelligence and market-entry pathway consulting
- Build a patent-landscape and prior-art research service for R&D teams
- Support medical-device and cosmetics registration for manufacturers and importers
- Deliver regulatory and IPR awareness training for pharma and biotech 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

This is to certify that

CANDIDATE NAME

completed the four months programme of

Post Graduate Diploma in

Regulatory Affairs, IPR & Patent

offered by

Global Institute of Health Science.

during MONTH YEAR to MONTH YEAR

Upon successful completion of the programme

POST GRADUATE DIPLOMA

IN

REGULATORY AFFAIRS IPR & PATENT

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:ghs-XXXX-XXXXX

Academic Head

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Global Institute Of Health Science

POST GRADUATE DIPLOMA
IN
Regulatory Affairs, IPR & Patent

MONTH YEAR to MONTH YEAR

CANDIDATE NAME

GRADE SHEET & TRANSCRIPT

Regulatory Affairs | Pre-Clinical Trial | Clinical Trial | Regulatory Bodies in India
Central Drug Regulatory System | Drug & Cosmetics Act | Schedule – Y
Medical Device Registration in India | Product Development Protocol
Environmental Protection Act –1986 | Pharmaceutical Legislation in India
The Narcotic drug & psychotropic substance act | Medicinal & Toilet preparation
Drug price control order in force | Laws on trademarks and copy rights
Prevention of cruelty to animal act | Consumer protection Act 1986
Pharmaceutical Regulation Process in India | General drug approach
Regulatory consideration for pre-clinical and clinical testing
Regulation and registration of medical devices & Cosmeticcs
New drug development procedure in different market
WHO Guidelines & certification in import export policy
Regulation of Generic Pharmaceutical and Bio Similar Product
Introduction and regulation of biosimilar in Indian & Europe
Reg.of herbal product & Orphan drug | Submission of Drug Master File to USFDA
Legal environment of business | Common Technical Documents
USFDA | ICH Guidelines | DRA in European union with special reference to EMA & UKMHRA
Auditing of Manufacturing Facility | IPR, Patents, Quality Assurance, Projects

Examination: 45/50 **Project Work: 35/50**

Total: 80/100

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

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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 regulatory affairs?

It is the profession that helps organisations understand and meet the regulatory requirements that apply to products and activities in regulated industries such as pharmaceuticals, biotechnology and medical devices.

Does the programme include IPR and patents?

Yes — IPR and patents, together with trademarks, copyrights and related regulatory and legal concepts.

Does it cover USFDA, ICH, EMA and UK MHRA?

Yes. USFDA, ICH guidelines, European Union authorities with reference to EMA and UK MHRA, WHO guidelines and Rest of World markets are all in the syllabus.

Does it cover CTD and Drug Master Files?

Yes. The Common Technical Document and submission of the Drug Master File to USFDA are both included.

Does it cover medical devices and biosimilars?

Yes — medical device registration in India, regulation of devices and cosmetics, generic products, and biosimilars in India and Europe.

Does this make me a registered patent agent or attorney?

No. Patent-agent and attorney status is governed by jurisdiction-specific legal and professional requirements. The programme builds professional IPR and patent knowledge but does not replace statutory registration.

Can AI prepare a regulatory submission automatically?

No. AI can support drafting and review, but submission content must be accurate, traceable, reviewed and compliant — and the accountable decision stays with a qualified professional.

Can working pharma professionals enrol?

Yes. The online and distance-learning format suits eligible working professionals seeking regulatory, quality, compliance or IPR 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