Global Institute Of Health Science™

CERTIFICATE · PG DIPLOMA · FAST-TRACK PG DIPLOMA

CLINICAL STUDY START-UP & REGULATORY SITE ACTIVATION

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 STUDY START-UP & REGULATORY SUBMISSIONS**

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 10 modules, and what each one teaches	3–4
03	**NEW: Artificial Intelligence in Study Start-Up & Regulatory Submissions** — 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.

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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 Study Start-Up & Site Activation

Practical | Job-Oriented | ICH-GCP | Start-Up | Ethics Submissions | Site Contracts | Trial Master File | AI Literacy

The GIHS Clinical Study Start-Up & Regulatory Site Activation programme covers the phase where most trials lose time: everything between a protocol being finalised and the first participant being consented. It teaches feasibility and site selection, regulatory and ethics submissions, contracts and budgets, essential-document collection, trial master file setup and the green-light process that activates a site.

Start-up is a documentation and coordination discipline. A site cannot screen a single participant until the approvals are in place, the agreement is signed, the product is released, the staff are trained and the essential documents are filed — and every one of those steps has an owner, a dependency and a date. This programme is built around managing exactly that.

> **"We Teach What the Job Requires."**
>
> GIHS teaches start-up the way it is run in a real study team — as a tracked list of dependencies with regulatory deadlines attached, not as an abstract process diagram.

Why This Programme Stands Out

- The full start-up path from protocol final to site green light
- Feasibility assessment and evidence-based site selection
- Regulatory submissions and ethics committee / IRB requirements
- Informed consent document development and local adaptation
- Site contracts, budgets, negotiation and payment schedules
- Essential documents, trial master file and investigator site file setup
- Site activation criteria, green light and readiness verification
- Start-up metrics, cycle times and timeline management
- Practical AI use cases for document preparation and tracking
- NEW dedicated AI module for start-up and regulatory submissions
- Certificate, PG Diploma and Fast-Track learning pathways

The Programme at a Glance

10	3 / 4 / 12	100%	Lifetime	MCQ + Project	2
Modules, including the new AI module	Months — Certificate, Fast Track or PG Diploma	Online & distance, study at your own pace	Free personal expert assigned to you	Online exam plus a 30-page project	International certificates awarded and shipped

REVISED SYLLABUS

Curriculum, Module by Module

The full GIHS syllabus, regrouped into 10 clearly sequenced modules so you can see exactly what each stage of the programme teaches and why it comes where it does. Module 10 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 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 04 Feasibility & Site Selection

Choosing sites that can actually recruit — and evidencing the choice.

- Country and site feasibility assessment
- Investigator and site qualification criteria
- Patient-population and competing-trial assessment
- Pre-study visits and site selection documentation
- Site start-up risk assessment

MODULE 05 Regulatory & Ethics Submissions

The approvals without which nothing else matters.

- Regulatory authority submissions and approval pathways
- Ethics Committee / IRB submission requirements
- Import licences, test licences and applicable permissions
- Informed consent document development and translation
- Protocol amendments and re-submission workflow
- Clinical trial registration requirements

MODULE 06 Contracts, Budgets & Site Agreements

Turning a willing site into a contracted one.

- Clinical trial agreements and site contracts
- Budget development, fair market value and negotiation
- Payment schedules, milestones and invoicing
- Confidentiality agreements and indemnity concepts
- Vendor and central-lab contracting interfaces

MODULE 07 Essential Documents, TMF & Site Files

The evidence file that the whole trial is later judged on.

- Essential documents before, during and after a trial
- Trial Master File structure and the DIA TMF Reference Model
- Investigator Site File setup and maintenance
- Document collection, quality review and filing standards
- eTMF systems, completeness and inspection readiness

MODULE 08 Site Activation & Start-Up Management

The green light, and the tracking that gets you there on time.

- Site activation criteria and green-light process
- Site initiation visit planning and delivery
- Investigational product release and first shipment
- Start-up cycle times, metrics and timeline tracking
- Start-up risk, escalation and contingency planning

MODULE 09 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 10 · NEW Artificial Intelligence in Study Start-Up & Regulatory Submissions

Document Preparation · Submission Tracking · Feasibility Analysis · Human Accountability

- AI across the start-up lifecycle — feasibility, submission, contracting, activation
- AI-assisted feasibility analysis and site-selection support
- AI support for regulatory and ethics submission document preparation
- … and 15 further topics — the full module is set out on page 5

MODULE 10 · NEW HIGHLIGHT MODULE

Artificial Intelligence in Study Start-Up & Regulatory Submissions

Document Preparation · Submission Tracking · Feasibility Analysis · Human Accountability

AI is now part of everyday clinical-development work. This module introduces practical AI concepts and responsible use cases for this role, with one consistent principle: AI is an assistive tool, never a substitute for qualified professional judgement, validated systems or regulatory accountability.

Key Topics Covered

- AI across the start-up lifecycle — feasibility, submission, contracting, activation
- AI-assisted feasibility analysis and site-selection support
- AI support for regulatory and ethics submission document preparation
- AI-assisted review of informed consent documents for readability and completeness
- AI in contract and budget review — clause comparison and gap spotting
- AI-assisted essential-document and TMF completeness checks
- AI for translation support and local-language document review
- AI-supported start-up timeline forecasting and bottleneck identification
- eTMF, eISF and start-up technology awareness
- Version control, document provenance and audit-trail discipline with AI drafts
- What AI must not do — approvals, submissions and regulatory commitments
- Introduction to AI, Machine Learning, Generative AI and LLMs in clinical research
- Prompt engineering for clinical research 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 ICH-GCP and applicable regulatory requirements

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. The skill that matters most is finding what the AI got wrong.

What This Module Adds to Your Career

The module shows where AI saves time and where professional oversight stays essential. You learn to evaluate AI output critically, protect confidential trial and participant 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 your skills survive the next change of technology and regulatory expectation.

AI-Enhanced Practice Exercises

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

- Check an AI-drafted consent document against protocol and local requirements
- Run a completeness check on a mock essential-document list
- Build a start-up timeline and identify the critical path
- Review an AI-summarised site feasibility pack for unsupported claims
- Draft a submission checklist and verify it against the applicable guidance

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 the competencies a start-up specialist is measured on: assessing feasibility, assembling submission packages, adapting consent documents, negotiating and tracking contracts, collecting and quality-reviewing essential documents, building and maintaining the TMF, and driving a site to green light against a timeline. Learners are trained to ask the start-up questions — what is outstanding, who owns it, what is it waiting on, and what does the regulation require before this site can screen a participant?

Career-Relevant Skills You Will Develop

- Study start-up planning and tracking
- Feasibility and site selection assessment
- Regulatory and ethics submission preparation
- Informed consent document development
- Clinical trial agreements and budget concepts
- Essential documents and TMF management
- Investigator site file setup and quality review
- Site activation and green-light process
- Start-up metrics and cycle-time management
- ICH-GCP and regulatory compliance
- AI-enabled start-up and submission literacy

Career & Professional Pathways

The current GIHS course page lists **Study Start-Up Specialist, Regulatory Start-Up Associate, Site Activation Coordinator and clinical operations pathways**. The programme can also support:

- Study Start-Up Specialist / Associate
- Site Activation Coordinator
- Regulatory Start-Up Associate
- Clinical Trial Assistant (CTA)
- Trial Master File / eTMF specialist
- Contracts and budgets support
- Ethics and regulatory submission support
- Clinical operations and study management
- Site contracts and feasibility support

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. Submission requirements differ by country, so candidates working across regions must also meet local regulatory 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

Every study needs start-up capacity in bursts, and few organisations keep enough of it on staff — which is exactly what makes it contractable work.

- Provide contract study start-up and submission support
- Offer ethics and regulatory documentation preparation services
- Build a trial master file quality-review and remediation practice
- Consult on site contracts, budgets and feasibility assessment
- Deliver start-up process and TMF training for sponsors and sites

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 Study Start-Up & Regulatory Site Activation** offered by Global Institute of Health Science, during **MONTH YEAR to MONTH YEAR**.
Upon successful completion of the programme

POST GRADUATE DIPLOMA

IN

CLINICAL STUDY START-UP & REGULATORY SITE ACTIVATION

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

Academic Head

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

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

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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 Study Start-Up & Regulatory Site Activation

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	45	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	45	50	46.5	93%
03	**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	45	50	47.5	95%
04	**Feasibility & Site Selection** Country and site feasibility assessment; Investigator and site qualification criteria; Patient-population and competing-trial assessment; Pre-study visits and site selection documentation; Site start-up risk assessment	45	50	47	94%
05	**Regulatory & Ethics Submissions** Regulatory authority submissions and approval pathways; Ethics Committee / IRB submission requirements; Import licences, test licences and applicable permissions; Informed consent document development and translation; Protocol amendments and re-submission workflow; Clinical trial registration requirements	45	50	47	94%
06	**Contracts, Budgets & Site Agreements** Clinical trial agreements and site contracts; Budget development, fair market value and negotiation; Payment schedules, milestones and invoicing; Confidentiality agreements and indemnity concepts; Vendor and central-lab contracting interfaces	45	50	48	96%
07	**Essential Documents, TMF & Site Files** Essential documents before, during and after a trial; Trial Master File structure and the DIA TMF Reference Model; Investigator Site File setup and maintenance; Document collection, quality review and filing standards; eTMF systems, completeness and inspection readiness	45	50	48	96%
08	**Site Activation & Start-Up Management** Site activation criteria and green-light process; Site initiation visit planning and delivery; Investigational product release and first shipment; Start-up cycle times, metrics and timeline tracking; Start-up risk, escalation and contingency planning	45	50	49	98%
09	**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	46	92%
10	**Artificial Intelligence in Study Start-Up & Regulatory Submissions** NEW Document Preparation · Submission Tracking · Feasibility Analysis · Human Accountability; AI across the start-up lifecycle — feasibility, submission, contracting, activation; AI-assisted feasibility analysis and site-selection support; AI support for regulatory and ethics submission document preparation	45	50	48.5	97%
	Module Examinations — Sub-total	**450**	**500**	476	**95.2%**
11	**Project Work** 30-page applied project submitted at the end of the programme	150	200	185	92.5%
	GRAND TOTAL	**600**	**700**	**661**	**94.4%**

TOTAL MARKS	PERCENTAGE	GRADE	RESULT	DATE OF AWARD	CERT. ID
661 / 700	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 study start-up?

Everything between protocol finalisation and a site being able to screen its first participant: feasibility, submissions and approvals, contracts and budgets, essential documents, training, product release and the activation green light.

Why does start-up get its own programme?

Because it is where trials most often lose time and money, and because it is a distinct skill set — regulatory documentation and coordination rather than monitoring.

Does it cover ethics committee and regulatory submissions?

Yes — submission requirements, approval pathways, consent document development and translation, amendments and trial registration are a dedicated module.

Is the trial master file covered?

Yes. Essential documents, TMF structure and the reference model, investigator site files, quality review and eTMF inspection readiness are covered in detail.

Are contracts and budgets included?

Yes — clinical trial agreements, budget development and negotiation concepts, payment schedules and confidentiality and indemnity awareness.

Why is AI included in a start-up programme?

Start-up is document-heavy, and AI is increasingly used for drafting, comparison and completeness checks. The module teaches where that helps and where it must not be trusted.

Does AI replace a start-up specialist?

No. Submissions, approvals and regulatory commitments remain human responsibilities; AI assists with preparation, comparison and tracking.

Can working clinical-research professionals enrol?

Yes. The online and distance-learning format suits professionals moving into start-up from coordination, regulatory or administrative roles.

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