Global Institute Of Health Science™

CERTIFICATE · PG DIPLOMA · FAST-TRACK PG DIPLOMA

CLINICAL TRIAL SUPPLY & INVESTIGATIONAL PRODUCT MANAGEMENT

A rare combination in one programme — and one fee.

NEW · AI MODULE **NOW WITH AI IN CLINICAL SUPPLY FORECASTING & IP MANAGEMENT**

What's inside this brochure

12 PAGES

01	**Programme Overview & Why It Stands Out** — What it covers, our teaching philosophy and the headline benefits	2
02	**Revised Syllabus — Module by Module** — All 9 modules, and what each one teaches	3–4
03	**NEW: Artificial Intelligence in Clinical Supply & Investigational Product Management** — 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

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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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★★★★★

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Recorded candidate satisfaction to date

PROGRAMME OVERVIEW

Build Future-Ready Skills in Clinical Supply & Investigational Product Management

Practical | Job-Oriented | ICH-GCP | GMP Interface | Forecasting | Cold Chain | IRT / IWRS | AI Literacy

The GIHS Clinical Trial Supply & Investigational Product Management programme covers the discipline that keeps a trial physically running: forecasting how much investigational product a study will need, getting it labelled, packaged, blinded, released, shipped and stored correctly, and accounting for every unit from manufacture to destruction.

Supply is where a well-designed trial meets logistics reality — customs, cold chain, expiry dates, randomisation systems and a site that has run out of kits on a Friday. The work is governed by GMP on one side and GCP on the other, and a single unexplained unit or temperature excursion can become an inspection finding.

> **"We Teach What the Job Requires."**
>
> GIHS teaches supply as the operational problem it is — demand planning under uncertainty, with regulatory constraints, real distribution networks and full accountability.

Why This Programme Stands Out

- Investigational product lifecycle from manufacture to destruction
- The GMP–GCP interface and QP release concepts
- Labelling, packaging, blinding and randomisation supply
- Demand forecasting, simulation and overage strategy
- Distribution, import–export, customs and cold-chain management
- IRT / IWRS systems and drug-supply triggers
- Accountability, reconciliation, returns and destruction
- Temperature excursions, deviations and quarantine decisions
- Practical AI use cases for forecasting and supply-risk detection
- NEW dedicated AI module for clinical supply and IP management
- Certificate, PG Diploma and Fast-Track learning pathways

REVISED SYLLABUS

Curriculum, Module by Module

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

MODULE 01 Foundations of Clinical Research

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

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

MODULE 02 Regulations & Key Stakeholders

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

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

MODULE 03 Investigational Product Fundamentals

What the product is, who releases it and under which rules it moves.

- Investigational medicinal product concepts and terminology
- The GMP and GCP interface in clinical supply
- Qualified Person release and certification concepts
- Comparators, placebo and auxiliary medicinal products
- Shelf life, expiry, retest dates and stability considerations

MODULE 04 Labelling, Packaging & Blinding

Getting the right kit, labelled correctly, into the right participant's hands.

- Clinical trial labelling requirements and local-language labels
- Packaging design, kit configuration and patient-ready supplies
- Blinding, unblinding and emergency code-break procedures
- Randomisation and kit-numbering concepts
- Booklet labels, expiry extension and re-labelling operations

MODULE 05 Forecasting, Planning & Supply Strategy

Deciding how much to make, for a study whose recruitment nobody can predict.

- Demand forecasting for clinical trials
- Overage, buffer stock and waste considerations
- Supply simulation and scenario planning
- Manufacturing campaign planning and lead times
- Budgeting and cost drivers in clinical supply

MODULE 06 Distribution, Import & Cold Chain

Moving a regulated product across borders without breaking the chain.

- Depot networks, direct-to-site and direct-to-patient models
- Import licences, export documentation and customs clearance
- Cold-chain packaging, qualification and temperature monitoring
- Shipment tracking, receipt and site storage requirements
- Temperature excursions, quarantine and disposition decisions

MODULE 07 IRT Systems, Accountability & Reconciliation

The systems and the paper trail that prove where every unit went.

- IRT / IWRS systems and drug-supply triggers
- Site-level accountability logs and dispensing records
- Drug reconciliation at monitoring visits and at close-out
- Returns, destruction and certificates of destruction
- Deviations, discrepancies and CAPA in supply operations

MODULE 08 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 09 · NEW Artificial Intelligence in Clinical Supply & Investigational Product Management

Demand Forecasting · Supply Risk · Cold-Chain Signals · Human Accountability

- AI across the supply lifecycle — forecasting, distribution, accountability
- AI-assisted demand forecasting and enrolment-driven supply modelling
- Simulation and scenario planning with AI support
- … and 15 further topics — the full module is set out on page 5

The Programme at a Glance

9
Modules, including the new AI module

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

100%
Online & distance, study at your own pace

Lifetime
Free personal expert assigned to you

MCQ + Project
Online exam plus a 30-page project

2
International certificates awarded and shipped

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

MODULE 09 · NEW HIGHLIGHT MODULE

Artificial Intelligence in Clinical Supply & Investigational Product Management

Demand Forecasting · Supply Risk · Cold-Chain Signals · 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 supply lifecycle — forecasting, distribution, accountability
- AI-assisted demand forecasting and enrolment-driven supply modelling
- Simulation and scenario planning with AI support
- AI for supply-risk detection — stock-outs, expiry exposure and waste
- AI-assisted analysis of temperature-excursion and shipment data
- AI support for label content review and translation checking
- AI-assisted reconciliation review and discrepancy identification
- AI in logistics routing, lead-time analysis and depot planning
- IRT data review and AI-supported anomaly detection
- What AI must not do — release decisions, unblinding and disposition calls
- Validated systems, GxP computerised-system expectations and change control
- 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:

- Review an AI demand forecast and challenge its recruitment assumptions
- Investigate a mock temperature excursion and decide the disposition question
- Reconcile a site accountability log against IRT records and explain the gap
- Assess an AI-assisted label review for regulatory completeness
- Build a supply risk register for a multi-country study

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 concentrates on the competencies a supply professional is judged on: forecasting demand, planning packaging and labelling campaigns, managing distribution and cold chain, configuring and reading IRT supply data, reconciling accountability records, and handling excursions, deviations and destruction correctly. Learners are trained to ask the supply questions — will the site run out, is the product still within expiry, was the chain maintained, can every unit be accounted for, and who is authorised to decide?

Career-Relevant Skills You Will Develop

- Clinical supply planning and forecasting
- Investigational product lifecycle management
- Labelling, packaging and blinding concepts
- Cold-chain and distribution management
- Import, export and customs documentation awareness
- IRT / IWRS supply configuration concepts
- Drug accountability and reconciliation
- Temperature excursion and deviation handling
- GMP–GCP interface awareness
- Supply risk assessment and contingency planning
- AI-enabled clinical supply literacy

Career & Professional Pathways

The current GIHS course page lists **Clinical Supply Associate, Investigational Product Coordinator, Clinical Logistics and trial-supply management pathways**. The programme can also support:

- Clinical Trial Supply Associate / Specialist
- Investigational Product Coordinator
- Clinical supply chain and logistics support
- Depot and distribution coordination
- IRT / IWRS supply configuration support
- Site pharmacy and dispensing support
- Drug accountability and reconciliation support
- Clinical operations and study management
- Cold-chain and quality-support roles

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. Product release, dispensing and regulated pharmacy activities remain restricted to appropriately qualified and authorised personnel.

PLACEMENT & CAREER ENHANCEMENT

What GIHS Does for Your Career After the Course

CAREER ENHANCEMENT INCLUDED FREE WITH EVERY ENROLMENT

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

FREE
worth ₹9,999

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

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

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

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

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

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

Where Our Graduates Are Making an Impact

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

Pharmaceutical companies
Clinical operations, drug safety and regulatory teams

Biotechnology & biosimilars
Development programmes and regulatory submissions

Contract Research Organisations
Monitoring, data management and safety services

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

Medical devices & diagnostics
Clinical evaluation, regulatory and quality functions

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

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

Academic & government research
Institutional trials, registries and regulatory bodies

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

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

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

Business & Entrepreneurship Value

Clinical supply sits between manufacturing, logistics and the trial, so independent expertise is valued by sponsors who have none of the three in-house.

- Provide clinical supply planning and forecasting support
- Offer depot, distribution and cold-chain coordination services
- Build an accountability and reconciliation review practice
- Consult on labelling, packaging and IRT supply configuration
- Deliver GMP-GCP interface and supply operations training

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 Trial Supply & Investigational Product Management** offered by Global Institute of Health Science, during **MONTH YEAR to MONTH YEAR**. Upon successful completion of the programme

POST GRADUATE DIPLOMA

IN

CLINICAL TRIAL SUPPLY & INVESTIGATIONAL PRODUCT MANAGEMENT

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

GLOBAL INSTITUTE OF HEALTH SCIENCE · GIHS · ACADEMIC OFFICE

Academic Head

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Global Institute Of Health Science

STATEMENT OF RESULT & STUDY

Grade Sheet & Transcript

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

PROGRAMME

Post Graduate Diploma in Clinical Trial Supply & Investigational Product Management

NO.	MODULE & TOPICS COVERED	HOURS	MAX	OBTAINED	%
01	**Foundations of Clinical Research** Introduction to clinical research; Clinical trial phases; Pharmacological principles of clinical research; Drug development and launch	54	50	48.5	97%
02	**Regulations & Key Stakeholders** Indian regulation (domestic regulation); ICH GCP / Schedule Y / ICMR / Indian GCP; Key stakeholders in clinical research; Ethics Committees and Institutional Review Board; Responsibilities of the sponsor and the investigator; Sponsor–investigator and sponsor–vendor responsibilities	54	50	46.5	93%
03	**Investigational Product Fundamentals** Investigational medicinal product concepts and terminology; The GMP and GCP interface in clinical supply; Qualified Person release and certification concepts; Comparators, placebo and auxiliary medicinal products; Shelf life, expiry, retest dates and stability considerations	54	50	48	96%
04	**Labelling, Packaging & Blinding** Clinical trial labelling requirements and local-language labels; Packaging design, kit configuration and patient-ready supplies; Blinding, unblinding and emergency code-break procedures; Randomisation and kit-numbering concepts; Booklet labels, expiry extension and re-labelling operations	54	50	47	94%
05	**Forecasting, Planning & Supply Strategy** Demand forecasting for clinical trials; Overage, buffer stock and waste considerations; Supply simulation and scenario planning; Manufacturing campaign planning and lead times; Budgeting and cost drivers in clinical supply	54	50	47	94%
06	**Distribution, Import & Cold Chain** Depot networks, direct-to-site and direct-to-patient models; Import licences, export documentation and customs clearance; Cold-chain packaging, qualification and temperature monitoring; Shipment tracking, receipt and site storage requirements; Temperature excursions, quarantine and disposition decisions	45	50	48	96%
07	**IRT Systems, Accountability & Reconciliation** IRT / IWRS systems and drug-supply triggers; Site-level accountability logs and dispensing records; Drug reconciliation at monitoring visits and at close-out; Returns, destruction and certificates of destruction; Deviations, discrepancies and CAPA in supply operations	45	50	48	96%
08	**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	49	98%
09	**Artificial Intelligence in Clinical Supply & Investigational Product Management** NEW Demand Forecasting · Supply Risk · Cold-Chain Signals · Human Accountability; AI across the supply lifecycle — forecasting, distribution, accountability; AI-assisted demand forecasting and enrolment-driven supply modelling; Simulation and scenario planning with AI support	45	50	46.5	93%
	Module Examinations — Sub-total	**450**	**450**	428.5	**95.2%**
10	**Project Work** 30-page applied project submitted at the end of the programme	150	200	185	92.5%
	GRAND TOTAL	**600**	**650**	**613.5**	**94.4%**

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

Academic Head

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

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ENROLMENT

Three Flexible Learning Options

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

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

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

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

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

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

What Our Candidates Say

★★★★★ 300+ Google reviews

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

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

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

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

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

Admission Procedure START ANY DAY

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

Who Can Enrol

LISTED ON THE GIHS COURSE PAGE

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

ALSO OPEN TO

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

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

QUESTIONS

Frequently Asked Questions

What does clinical trial supply management involve?

Forecasting how much investigational product a study needs, then labelling, packaging, releasing, shipping, storing, dispensing, reconciling and destroying it under GMP and GCP requirements.

Is this a pharmacy qualification?

No. It is professional education in clinical supply operations. Dispensing and regulated pharmacy activities remain restricted to appropriately qualified and authorised personnel.

Does it cover cold chain and temperature excursions?

Yes — cold-chain packaging and qualification, temperature monitoring, excursion handling, quarantine and disposition decisions are covered in a dedicated module.

What is an IRT or IWRS system?

An interactive response system that randomises participants and triggers drug supply to sites. The programme covers how supply settings, resupply triggers and its data are used.

Are import and export requirements included?

Yes, at an operational level — import licences, export documentation, customs clearance and depot-network models, with the reminder that requirements differ by country.

Why is AI included in a supply programme?

Forecasting, risk detection and excursion analysis are exactly the problems AI is now applied to. The module teaches how to use it and how to challenge its assumptions.

Does AI replace a supply professional?

No. Release, unblinding and disposition decisions stay with authorised people; AI assists with forecasting, analysis and review support.

Can working pharma or CRO professionals enrol?

Yes. The online and distance-learning format suits professionals moving into clinical supply from pharmacy, logistics, manufacturing or clinical operations.

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