Global Institute Of Health Science™

CERTIFICATE · PG DIPLOMA · FAST-TRACK PG DIPLOMA

BIOSTATISTICS & CLINICAL STATISTICAL PROGRAMMING

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 BIOSTATISTICS & CLINICAL STATISTICAL PROGRAMMING**

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 Biostatistics & Clinical Statistical Programming** — 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 Biostatistics & Clinical Statistical Programming

Practical | Job-Oriented | Biostatistics | SAS | R | CDISC | SDTM & ADaM | AI Literacy

The GIHS Biostatistics & Clinical Statistical Programming programme covers how clinical trial data becomes evidence: the statistical thinking behind trial design, the analysis plan that commits to it in advance, the CDISC standards that carry the data, and the programming that produces the tables, listings and figures a regulator actually reads.

Statistics is what stops a trial from telling us what we hoped rather than what happened. Randomisation, sample size, pre-specified endpoints, analysis populations and a locked statistical analysis plan exist to protect the result from the enthusiasm of the people producing it — and every programmer working on a submission is part of that protection.

> **"We Teach What the Job Requires."**
>
> GIHS teaches the statistics a trial team actually uses and the programming that delivers it — specification, standard-compliant datasets, reproducible output and documented validation.

Why This Programme Stands Out

- Statistical foundations built for clinical trials, not general theory
- Randomisation, blinding, sample size and power in practice
- Endpoints, estimands, analysis populations and missing data
- Statistical analysis plan structure and pre-specification discipline
- CDISC standards — SDTM, ADaM, define.xml and submission datasets
- Statistical programming in SAS, with R awareness
- Tables, listings and figures for clinical study reports
- Validation, double programming and reproducibility
- Practical AI use cases for code drafting, review and documentation
- NEW dedicated AI module for biostatistics and statistical programming
- 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 Statistical Foundations for Clinical Trials

The statistical vocabulary every clinical team is expected to share.

- Types of data, distributions and descriptive statistics
- Probability, confidence intervals and p-values in context
- Hypothesis testing, type I and type II error
- Common tests — t-test, chi-square, ANOVA and non-parametric alternatives
- Correlation, regression and survival-analysis concepts

MODULE 03 Trial Design, Randomisation & Sample Size

Designing a study that can answer its question, before a single participant is enrolled.

- Parallel, crossover, factorial and adaptive designs
- Superiority, non-inferiority and equivalence framing
- Randomisation methods, stratification and blinding
- Sample size, power and assumptions
- Bias, confounding and multiplicity

MODULE 04 Endpoints, Estimands & Analysis Populations

Deciding exactly what will be measured, in whom, and how deviations are handled.

- Primary, secondary and exploratory endpoints
- The estimand framework and intercurrent events
- ITT, per-protocol and safety analysis populations
- Missing data, imputation strategies and sensitivity analyses
- Interim analysis and data monitoring committee concepts

MODULE 05 Statistical Analysis Plan & Reporting

Committing to the analysis in writing, then producing the output that is actually read.

- Statistical analysis plan structure and content
- Pre-specification, amendments and documented rationale
- Table, listing and figure (TLF) shells and specifications
- Clinical study report statistical sections
- Regulatory expectations for analysis and presentation

MODULE 06 CDISC Standards — SDTM & ADaM

The data standards every submission is expected to speak.

- CDISC overview and the submission data journey
- SDTM domains, structure and mapping concepts
- ADaM datasets, traceability and analysis-ready structure
- define.xml, annotated CRF and submission documentation
- Controlled terminology and standards compliance checks

MODULE 07 Clinical Statistical Programming

Writing the code that produces the numbers — and proving the numbers are right.

- SAS foundations for clinical programming — DATA step and procedures
- Producing analysis datasets from raw clinical data
- Generating tables, listings and figures to specification
- Programming in R for clinical analysis — an introduction
- Validation, double programming, QC and reproducibility
- Version control, documentation and handover standards

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 Biostatistics & Clinical Statistical Programming

Code Drafting · Output Review · Documentation · Human Accountability

- AI across the analysis lifecycle — specification, programming, validation, reporting
- AI-assisted drafting of SAS and R code, and why it must be reviewed line by line
- AI support for interpreting specifications and TLF shells
- … 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 Biostatistics & Clinical Statistical Programming

Code Drafting · Output Review · Documentation · 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 analysis lifecycle — specification, programming, validation, reporting
- AI-assisted drafting of SAS and R code, and why it must be reviewed line by line
- AI support for interpreting specifications and TLF shells
- AI-assisted code review, debugging and efficiency improvement
- AI for documentation, program headers and handover notes
- AI-assisted SDTM and ADaM mapping support, with traceability preserved
- Machine learning in clinical research — prediction versus inference
- AI-supported data review, outlier detection and quality checks
- Why statistical validity is not a coding question — pre-specification and bias
- Reproducibility, environment control and validated systems
- What AI must not do — choose the analysis, interpret the result, sign the output
- 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 AI-generated SAS code against a TLF specification and find the errors
- Explain why a plausible AI-suggested analysis would break pre-specification
- Map a small raw dataset to SDTM and check traceability to ADaM
- Validate an AI-assisted table against an independently programmed version
- Write program documentation that a reviewer could follow without you

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 is built around the deliverables a statistical team is measured on: a clear statistical analysis plan, standard-compliant datasets, tables, listings and figures that match their specifications, and validation evidence that the output is correct. Learners are trained to ask the statistical questions — what was pre-specified, which population is this, how is missing data handled, is the result traceable to the source data, and has anyone independently reproduced it?

Career-Relevant Skills You Will Develop

- Biostatistics for clinical trials
- Trial design, randomisation and sample size
- Endpoints, estimands and analysis populations
- Statistical analysis plan development
- CDISC SDTM and ADaM concepts
- SAS programming for clinical data
- R programming awareness for clinical analysis
- Tables, listings and figures production
- Validation, double programming and QC
- Data integrity and reproducibility
- AI-enabled statistical programming literacy

Career & Professional Pathways

The current GIHS course page lists **Clinical SAS Programmer, Statistical Programmer, Biostatistics Associate and clinical data-science pathways**. The programme can also support:

- Clinical SAS Programmer
- Statistical Programmer / Analyst
- Biostatistics Associate
- Clinical data science and analytics support
- CDISC / standards support roles
- Clinical data management interface roles
- Statistical QC and validation support
- Medical writing statistical support
- Real-world evidence and observational analysis 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. Responsibility for the statistical design and interpretation of a trial rests with the qualified statistician accountable for the study.

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

Statistical programming is contracted per study across the industry, and a validated, reproducible deliverable is worth more than hours on a timesheet.

- Provide contract clinical SAS and statistical programming
- Offer CDISC SDTM and ADaM mapping and conversion services
- Build an independent validation and double-programming practice
- Consult on statistical analysis plans and TLF specifications
- Deliver clinical programming and CDISC standards 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 Biostatistics & Clinical Statistical Programming** offered by Global Institute of Health Science, during **MONTH YEAR to MONTH YEAR**. Upon successful completion of the programme

POST GRADUATE DIPLOMA

IN

BIOSTATISTICS & CLINICAL STATISTICAL PROGRAMMING

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

GLOBAL INSTITUTE OF HEALTH SCIENCE · GIHS · ACADEMIC OFFICE

Academic Head

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

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Global Institute Of Health Science

STATEMENT OF RESULT & STUDY

Grade Sheet & Transcript

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

PROGRAMME

Post Graduate Diploma in Biostatistics & Clinical Statistical Programming

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	**Statistical Foundations for Clinical Trials** Types of data, distributions and descriptive statistics; Probability, confidence intervals and p-values in context; Hypothesis testing, type I and type II error; Common tests — t-test, chi-square, ANOVA and non-parametric alternatives; Correlation, regression and survival-analysis concepts	54	50	46.5	93%
03	**Trial Design, Randomisation & Sample Size** Parallel, crossover, factorial and adaptive designs; Superiority, non-inferiority and equivalence framing; Randomisation methods, stratification and blinding; Sample size, power and assumptions; Bias, confounding and multiplicity	54	50	48	96%
04	**Endpoints, Estimands & Analysis Populations** Primary, secondary and exploratory endpoints; The estimand framework and intercurrent events; ITT, per-protocol and safety analysis populations; Missing data, imputation strategies and sensitivity analyses; Interim analysis and data monitoring committee concepts	54	50	47	94%
05	**Statistical Analysis Plan & Reporting** Statistical analysis plan structure and content; Pre-specification, amendments and documented rationale; Table, listing and figure (TLF) shells and specifications; Clinical study report statistical sections; Regulatory expectations for analysis and presentation	54	50	47	94%
06	**CDISC Standards — SDTM & ADaM** CDISC overview and the submission data journey; SDTM domains, structure and mapping concepts; ADaM datasets, traceability and analysis-ready structure; define.xml, annotated CRF and submission documentation; Controlled terminology and standards compliance checks	45	50	48	96%
07	**Clinical Statistical Programming** SAS foundations for clinical programming — DATA step and procedures; Producing analysis datasets from raw clinical data; Generating tables, listings and figures to specification; Programming in R for clinical analysis — an introduction; Validation, double programming, QC and reproducibility; Version control, documentation and handover standards	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 Biostatistics & Clinical Statistical Programming** NEW Code Drafting · Output Review · Documentation · Human Accountability; AI across the analysis lifecycle — specification, programming, validation, reporting; AI-assisted drafting of SAS and R code, and why it must be reviewed line by line; AI support for interpreting specifications and TLF shells	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

Do I need a statistics degree to take this programme?

No. The statistical foundations are taught from the beginning, in clinical-trial context. A comfort with numbers and logical structure helps more than prior statistical training.

Which software does the programme cover?

SAS is the primary focus because it remains the standard for regulatory submissions, with an introduction to R for clinical analysis.

What are SDTM and ADaM?

CDISC data standards: SDTM organises the collected trial data, ADaM organises the analysis-ready datasets derived from it. Both are covered along with define.xml and controlled terminology.

What is an estimand?

A precise statement of what the trial is estimating — the population, the endpoint, the treatment condition and how intercurrent events are handled. It is covered in full.

Is the statistical analysis plan covered?

Yes — its structure, the discipline of pre-specification, amendments with documented rationale, and the TLF specifications that follow from it.

Why is AI included in a statistics programme?

AI now drafts code quickly and confidently, including code that is subtly wrong. The module teaches how to use it for speed while validating every output that carries a result.

Does AI replace a statistical programmer?

No. Choosing the analysis, interpreting the result and signing off output remain human responsibilities, and validated, reproducible programming is a regulatory expectation.

Can working data-management or pharma professionals enrol?

Yes. The online and distance-learning format suits professionals moving into programming and analysis from data management, clinical operations or life-science backgrounds.

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

INDIA | UK | AUSTRALIA | RUSSIA | CANADA | USA | NEW ZEALAND | UAE | AFRICA

CHAT ON WHATSAPP