Global Institute Of Health Science™

CERTIFICATE · PG DIPLOMA · FAST-TRACK PG DIPLOMA

CLINICAL RESEARCH ASSOCIATE (CRA) & SITE MANAGEMENT

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 CLINICAL MONITORING & SITE 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 10 modules, and what each one teaches	3–4
03	**NEW: Artificial Intelligence in Clinical Monitoring & Site 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

Trusted by Healthcare Professionals Globally

Learn from global perspectives and prepare for impactful roles in hospitals, public health systems and healthcare organisations around the world.

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

15+ years of experience **2,000+** placed globally **Lifetime** free expert tutoring

ACCREDITED & CERTIFIED

- **ISO 21001:2018** Educational organisation management
- **UASL** Germany / UK — quality management
- **GSAAA** USA — global accreditation
- **IPR, Govt. of India** Copyright approved

ENROL ANY DAY · ADMISSIONS OPEN ALL YEAR

Scan to enrol — or apply online

www.gihsonline.com/enroll-now/

info@gihsonline.com · +91-9033768588 · www.gihsonline.com

★★★★★

300+ Google reviews

Recorded candidate satisfaction to date

PROGRAMME OVERVIEW

Build Future-Ready Skills in Clinical Monitoring & Site Management

Practical | Job-Oriented | ICH-GCP | Monitoring | Site Management | Source Data Verification | Risk-Based Monitoring | AI Literacy

The GIHS Clinical Research Associate (CRA) & Site Management programme prepares professionals for the role that holds a clinical trial together at the point where it actually happens — the investigator site. It covers the full monitoring cycle: selecting and qualifying a site, initiating it, monitoring it through recruitment and treatment, verifying that the data reflect the source, and closing it out to an inspection-ready standard.

A CRA is the sponsor's eyes at the site and the site's route back to the sponsor. The job is part verification, part relationship, part project management: confirming consent was taken properly, that the protocol is being followed, that the investigational product is accounted for, that deviations are recognised and reported — and doing it while keeping a busy clinical team willing to take your call.

> **"We Teach What the Job Requires."**
>
> GIHS builds the programme around what a monitor is actually asked to do in the working week — visit planning, source-data verification, query resolution, site training, escalation, follow-up letters and the reports that prove the visit happened.

Why This Programme Stands Out

- The complete monitoring cycle — selection, initiation, routine, close-out
- Source data verification and source data review, in practice
- Site relationship management, training and escalation handling
- Protocol deviations, non-compliance and corrective action at site level
- Informed consent verification and participant-protection checks
- Investigational product accountability and temperature-excursion handling
- Risk-based and remote monitoring concepts, and what they change
- Monitoring visit reports and follow-up letters that stand up to audit
- Practical AI use cases for visit preparation, data review and reporting
- NEW dedicated AI module for clinical monitoring and site management
- 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 Study Setup & Site Management

Getting a site open and recruiting — the stage where most trials lose time.

- Site selection and pre-study visits
- Site initiation
- Subject recruitment and retention planning
- Site contracts and budgeting

MODULE 05 Site Management & Investigator Relationships

The part of the job no checklist covers — keeping a site trained, motivated and responsive.

- Site qualification and feasibility assessment
- Site training, delegation log and role documentation
- Recruitment support, screening logs and retention discussions
- Communication, escalation and difficult-conversation handling
- Site payments, contracts and query follow-through

MODULE 06 Clinical Monitoring

The CRA's working week — verifying at the site that the data match the source.

- Roles of the lead and principal investigator
- Clinical monitoring activities
- CRF review and source data verification
- Drug safety reporting
- Drug accountability work
- Routine site monitoring and site close-out visit

MODULE 07 Source Data Verification & Data Quality

Proving that what the database says is what the patient's record says.

- Source data, source documents and the ALCOA+ principles
- Source data verification and source data review
- Query generation, resolution and data-clarification workflow
- Protocol deviations, violations and reporting obligations
- Informed consent verification at monitoring visits
- Investigational product accountability and reconciliation at site

MODULE 08 Monitoring Models, Reports & Close-Out

Writing the visit up, following it through, and closing the site properly.

- Risk-based monitoring and centralised monitoring concepts
- Remote and hybrid monitoring visits
- Monitoring visit report writing and follow-up letters
- Action-item tracking and issue escalation
- Site close-out visit, archiving and record retention

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 Clinical Monitoring & Site Management

Risk Signals · Visit Preparation · Source Review Support · Human Accountability

- AI across the monitoring lifecycle — planning, conduct, follow-up, close-out
- Risk-based quality management and AI-supported risk signal detection
- Centralised statistical monitoring and outlier identification
- … and 15 further topics — the full module is set out on page 5

MODULE 10 · NEW HIGHLIGHT MODULE

Artificial Intelligence in Clinical Monitoring & Site Management

Risk Signals · Visit Preparation · Source Review Support · 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 monitoring lifecycle — planning, conduct, follow-up, close-out
- Risk-based quality management and AI-supported risk signal detection
- Centralised statistical monitoring and outlier identification
- AI-assisted review of site performance, enrolment and query metrics
- AI support for monitoring visit preparation and checklist building
- AI-assisted drafting of visit reports and follow-up letters — with human review
- AI for protocol and essential-document familiarisation before a visit
- AI-assisted identification of documentation gaps and inconsistencies
- AI in site feasibility, selection and recruitment forecasting
- eSource, eConsent and decentralised-trial technology awareness
- What AI must not do — consent verification, source verification, judgement calls
- 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-drafted monitoring visit report for unsupported statements
- Build a risk-based monitoring plan for a mock study and test it against GCP
- Spot the documentation gaps in a set of mock site files
- Draft a follow-up letter and check every action item is traceable
- Compare manual and AI-assisted review of site performance metrics

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 on workplace competencies: qualifying a site, running an initiation visit, verifying consent and source data, resolving queries, handling deviations, managing investigational product accountability, writing reports that survive audit and closing a site out. Learners are trained to ask what a monitor asks — is the participant protected, does the data match the source, is the deviation reported, is the product accounted for, and is there evidence that I checked?

Career-Relevant Skills You Will Develop

- Clinical monitoring across the visit cycle
- Site selection, initiation and close-out
- Source data verification and source data review
- ICH-GCP and regulatory compliance
- Informed consent verification
- Protocol deviation identification and reporting
- Investigational product accountability
- Monitoring visit report writing
- Risk-based and remote monitoring concepts
- Site relationship and escalation management
- AI-enabled clinical monitoring literacy

Career & Professional Pathways

The current GIHS course page lists **Clinical Research Associate, Site Management Associate, Clinical Trial Assistant and clinical operations pathways**. The programme can also support:

- Clinical Research Associate (CRA)
- Site Management Associate (SMA)
- Clinical Trial Assistant (CTA)
- Clinical Research Coordinator (CRC)
- In-house / centralised monitoring support
- Clinical operations and study management
- Site quality and compliance support
- Trial master file and documentation support
- Investigator-site study coordination

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. ICH-GCP principles are internationally relevant, but candidates seeking work abroad must also meet destination-country employer, immigration and regulatory 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

Sponsors and CROs buy monitoring capacity by the visit, which makes site-facing work one of the easiest clinical-research skills to offer independently.

- Provide contract CRA and site-monitoring support to CROs
- Offer site start-up, training and coordination services to investigator sites
- Build a site quality and inspection-readiness review practice
- Consult on protocol deviation handling and CAPA at site level
- Deliver ICH-GCP and monitoring training for site and sponsor teams

Any regulated clinical service must only be provided within your lawful professional scope and applicable registration requirements. Career titles, eligibility and the right to practise depend on your prior qualification, experience, employer criteria and local regulation.

Global Institute Of Health Science

Certificate

OF POST GRADUATE DIPLOMA

This is to certify that

CANDIDATE NAME

has successfully completed the one year programme of **Post Graduate Diploma in Clinical Research Associate (CRA) & Site 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 RESEARCH ASSOCIATE (CRA) & SITE MANAGEMENT

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

SCAN TO VERIFY

Verify Certification : **https://www.gihsonline.com/verify-certification/** | info@gihsonline.com | +91-9033768588

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

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 Research Associate (CRA) & Site 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	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	**Study Setup & Site Management** Site selection and pre-study visits; Site initiation; Subject recruitment and retention planning; Site contracts and budgeting	45	50	47	94%
05	**Site Management & Investigator Relationships** Site qualification and feasibility assessment; Site training, delegation log and role documentation; Recruitment support, screening logs and retention discussions; Communication, escalation and difficult-conversation handling; Site payments, contracts and query follow-through	45	50	47	94%
06	**Clinical Monitoring** Roles of the lead and principal investigator; Clinical monitoring activities; CRF review and source data verification; Drug safety reporting; Drug accountability work; Routine site monitoring and site close-out visit	45	50	48	96%
07	**Source Data Verification & Data Quality** Source data, source documents and the ALCOA+ principles; Source data verification and source data review; Query generation, resolution and data-clarification workflow; Protocol deviations, violations and reporting obligations; Informed consent verification at monitoring visits; Investigational product accountability and reconciliation at site	45	50	48	96%
08	**Monitoring Models, Reports & Close-Out** Risk-based monitoring and centralised monitoring concepts; Remote and hybrid monitoring visits; Monitoring visit report writing and follow-up letters; Action-item tracking and issue escalation; Site close-out visit, archiving and record retention	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 Clinical Monitoring & Site Management** NEW Risk Signals · Visit Preparation · Source Review Support · Human Accountability; AI across the monitoring lifecycle — planning, conduct, follow-up, close-out; Risk-based quality management and AI-supported risk signal detection; Centralised statistical monitoring and outlier identification	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

SCAN TO VERIFY

Verify Certification : **https://www.gihsonline.com/verify-certification/** | info@gihsonline.com | +91-9033768588

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

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 a Clinical Research Associate actually do?

A CRA monitors investigator sites on behalf of a sponsor or CRO: qualifying and initiating sites, verifying consent and source data, resolving queries, checking product accountability, reporting deviations and closing sites out.

Is this different from a Clinical Research Coordinator?

Yes. A coordinator works at the site and runs the study day to day; a CRA visits the site on the sponsor's behalf and verifies that it is being run correctly. The programme covers both sides of that relationship.

Does the programme cover source data verification in detail?

Yes — source documents, ALCOA+ principles, source data verification and review, query workflow, deviation handling and consent verification are a full module.

What is risk-based monitoring, and is it covered?

It is an approach that concentrates monitoring effort where the risk to participants and data is highest, supported by centralised data review. It is covered along with remote and hybrid visit models.

Why is AI included in a monitoring programme?

Monitoring is increasingly supported by centralised data review, risk signals and AI-assisted document work. AI literacy helps you use these tools and recognise where they are unreliable.

Does AI replace a CRA?

No. Consent verification, source verification and judgement calls at a site remain human responsibilities; AI assists with preparation, review support and drafting.

Will this help me move into a CRA role?

It strengthens the knowledge relevant to CRA, SMA, CTA and coordination pathways. Actual appointment depends on your qualifications, experience and employer requirements.

Can working clinical-research professionals enrol?

Yes. The online and distance-learning format suits working professionals moving from coordination, data or safety roles into monitoring.

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