



2026 EDITION

Life Sciences & Healthcare *Training* Programs

Equipping life sciences and healthcare teams to develop, deliver and stay compliant

INSTRUCTOR-LED

ENTERPRISE READY

CUSTOMIZABLE

GLOBAL DELIVERY

45

COURSES

13

CATEGORIES

5,000+

EXPERT TRAINERS

Building *compliant* life sciences teams

Edstellar is an enterprise learning partner that helps organizations build future-ready workforces. Our 2,000+ instructor-led programs are delivered onsite or live online by 5,000+ practitioner trainers, and each one is tailored to your goals. This catalog brings together 45 life sciences & healthcare programs that span pharma and medtech regulatory compliance, digital health and health informatics, clinical development and data, hospital operations and revenue cycle, and medical affairs and market access.

Global leaders including Visa, ABB and Aditya Birla Group trust Edstellar to build capability at enterprise scale. Before any program starts, we look at the real capability gap and the business goal behind it, so every module ties back to outcomes your leaders can measure, not just attendance. The result is training your clinical, quality and commercial teams put to work straight away.

Learning paths for *every* role



Regulatory & Quality Affairs

COMPLIANCE & QUALITY

Keep products compliant through GxP, submissions and quality systems across pharma and medical devices.

START WITH

- Good Manufacturing Practice for Pharma & Biotech
- Regulatory Affairs: FDA, EMA, MHRA & CDSCO Submissions
- ISO 13485 Medical Device Quality Management System



Digital Health & Informatics Teams

DATA & INTEROPERABILITY

Modernize clinical systems and move health data safely between platforms and care settings.

START WITH

- Clinical Informatics & EHR Optimization
- HL7 FHIR for Health Technology Teams
- Healthcare Data Governance & Interoperability



Clinical Research & Development

TRIALS & EVIDENCE

Run trials to good practice, manage clinical data and generate the evidence that supports approvals.

START WITH

- Good Clinical Practice (GCP) ICH E6(R3)
- Clinical Trial Operations & Decentralised Trials
- Biostatistics for Clinical Research



Hospital & Commercial Leaders

OPERATIONS & ACCESS

Improve patient flow, revenue cycle and market access so services run efficiently and sustainably.

START WITH

- Hospital Operations & Patient Flow Management
- Market Access, Pricing & Reimbursement (HTA)
- Medical Coding, Billing & Revenue Cycle Management

Why *choose* us

01

Practitioner trainers

5,000+ verified clinicians, regulatory specialists and quality leaders who work in real life sciences and healthcare settings.

02

Customizable programs

Curriculum built around your therapeutic areas, regulatory landscape, systems and organizational maturity.

03

Global delivery

Onsite or live virtual in any time zone, with in-person delivery across 1,000+ locations.

04

ISO 9001 & ISO 27001 certified

ISO 9001:2015 for quality and ISO 27001:2022 for information security.

How we *work*

01

Submit your brief

Share your training needs, team size and skill gaps.

02

Custom plan

Receive a tailored curriculum and trainer recommendations.

03

Approve & schedule

Finalize content, dates and delivery mode for your team.

04

Train & measure

Expert-led sessions with progress tracking and ROI reporting.

Course *categories*

01 Pharma & MedTech Regulatory Compliance

11

03 Clinical Development & Data

4

05 Medical Affairs, Market Access & Commercial

4

07 Patient Safety, Infection Control & Public Health

3

09 Medical Device Quality & Risk

2

11 Biotech & Bio-Manufacturing

1

13 Computer System Validation

1

02 Digital Health & Health Informatics

7

04 Hospital Operations, Quality & Revenue Cycle

4

06 Biomanufacturing & Process Development

3

08 Clinical Research

2

10 Pharma Quality & Drug Safety

2

12 Clinical Leadership

1

45 courses in total · full list on the pages that follow

All 45 courses

01 Pharma & MedTech Regulatory Compliance

11

EU MDR & IVDR Medical Device Compliance	→
FDA 510(k), De Novo & PMA Device Pathways	→
Good Distribution Practice (GDP) for Pharmaceuticals	→
Good Laboratory Practice (GLP) & Data Integrity (ALCOA+)	→
Good Manufacturing Practice for Pharma & Biotech	→
GxP Data Integrity & ALCOA+	→
Pharmaceutical Sales Compliance & Promotional Practices	→
Pharmaceutical Serialisation & Track-&-Trace Compliance	→
Regulatory Affairs: FDA, EMA, MHRA & CDSCO Submissions	→
Software as a Medical Device (SaMD) & AI/ML Device Regulation	→
Software as a Medical Device (SaMD) & IEC 62304	→

02 Digital Health & Health Informatics

7

Clinical Informatics & EHR Optimization	→
Digital Health Product Management & Clinical Safety (DCBO129)	→
Digital Health Strategy & Transformation	→
Genomics & Precision Medicine for Health Systems	→
Healthcare Data Governance & Interoperability	→
HL7 FHIR for Health Technology Teams	→
Telehealth & Remote Patient Monitoring Operations	→

03 Clinical Development & Data

4

Bioinformatics & Genomic Data Analysis	→
Biostatistics for Clinical Research	→
Clinical Data Management (CDM) & EDC Systems	→
Clinical Trial Operations & Decentralised Trials	→

04 Hospital Operations, Quality & Revenue Cycle

4

Healthcare Accreditation: JCI, NABH & CQC Readiness	→
Hospital Operations & Patient Flow Management	→
Lean Healthcare & Clinical Process Improvement	→
Medical Coding, Billing & Revenue Cycle Management	→

05 Medical Affairs, Market Access & Commercial

4

Market Access, Pricing & Reimbursement (HTA)	→
Medical Affairs & Scientific Communication	→
Real-World Evidence & Health Economics (HEOR)	→
Veeva CRM for Life Sciences	→

06 Biomanufacturing & Process Development

3

Biologics & Biosimilars Manufacturing	→
Biopharmaceutical Process Development & Scale-Up	→

06 Biomanufacturing & Process Development (cont.)

3

Cell & Gene Therapy Manufacturing & Supply	→
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07 Patient Safety, Infection Control & Public Health

3

Infection Prevention & Control	→
Patient Safety & Clinical Quality Improvement	→
Public Health Emergency Preparedness	→

08 Clinical Research

2

Good Clinical Practice (GCP) ICH E6(R3)	→
Medical Writing & Clinical Documentation	→

09 Medical Device Quality & Risk

2

ISO 13485 Medical Device Quality Management System	→
Medical Device Risk Management with ISO 14971	→

10 Pharma Quality & Drug Safety

2

Pharmaceutical Quality Systems & CAPA	→
Pharmacovigilance & Drug Safety (GVP)	→

11 Biotech & Bio-Manufacturing

1

Synthetic Biology & Bio-Manufacturing for Industry	→
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12 Clinical Leadership

1

Clinical Leadership & Physician Management Development	→
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13 Computer System Validation

1

Computer System Validation (CSV) & GAMP 5	→
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Trusted by *enterprises*

“*Edstellar understood that pharma regulatory training cannot be generic. The GxP and MedTech compliance modules mapped directly to our submissions, and the clinical data sessions gave our teams a shared standard before an audit, not after.*”

VP OF REGULATORY AFFAIRS · GLOBAL PHARMACEUTICAL COMPANY

“*The digital health and health informatics program moved our hospital teams from paper habits to confident use of our clinical systems. Patient safety outcomes followed.*”

CHIEF MEDICAL INFORMATION OFFICER ·
MULTINATIONAL HEALTHCARE PROVIDER

“*We needed quality and drug safety capability at scale across sites. The training was rigorous, practical and built around our own SOPs.*”

DIRECTOR OF QUALITY & COMPLIANCE ·
INTERNATIONAL BIOTECHNOLOGY GROUP

TRUSTED BY **Visa** **ABB** **Aditya Birla Group** & OTHER FORTUNE 500 ORGANIZATIONS

Let's *connect*

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[Schedule a consultation →](#)



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