GLOBAL GxP TRAINING

MASTER TOPICS CATALOGUE

Pharmaceuticals • Biotechnology • Medical Devices • Cosmetics • Nutraceuticals

Regulatory compliance | Quality systems | Laboratories | Engineering | Manufacturing | Clinical research

294	18	2026
TRAINING TOPICS	SPECIALIST CATEGORIES	CATALOGUE EDITION

PURPOSE

A structured list of available training subject areas for client needs assessment, annual training plans, professional development programmes, and customised onsite/online workshops. Select topics according to the client's product type, jurisdiction, job roles and actual regulatory obligations. This catalogue is a proposed training menu, not a claim that every standard applies to every facility.

TRAINING DELIVERY OPTIONS

Awareness session (2–4 hours) • One-day interactive course • Two-day detailed workshop • Multi-day practical masterclass • Role-specific onsite coaching • Mock audit and case-study exercises.

For in-house teams, regulated laboratories, technical leadership and project delivery teams.

ENSURING GLOBAL GMP COMPLIANCE

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CATALOGUE INDEX

18 focused programmes • 294 modules • Sector-specific regulatory references

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01	Global GxP & Regulatory Fundamentals	14
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KEY TERMINOLOGY

GMP = Good Manufacturing Practice; cGMP = current GMP; GLP = Good Laboratory Practice (OECD nonclinical safety studies when specified); GCP = Good Clinical Practice; GEP = Good Engineering Practice; GDocP = Good Documentation Practice; GDP = Good Distribution Practice; GSP = Good Storage Practice; GCLP = Good Clinical Laboratory Practice; GVP = Good Pharmacovigilance Practice; QMS/PQS = (Pharmaceutical) Quality Management System; QRM = Quality Risk Management.

Choose current, jurisdiction-specific rules for each client. US FDA device QMSR took effect on 2 Feb 2026; ICH E6(R3) Annex 2 will become effective in the EU on 15 Jan 2027.

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01 | Global GxP & Regulatory Fundamentals

APPLICATION: All sectors | Core regulatory foundation

KEY REFERENCES: WHO GMP; EU GMP (EudraLex Vol. 4); US FDA 21 CFR; PIC/S; ICH; Indian Schedule M; applicable national legislation

Sr. No.	TRAINING TOPIC
001	Introduction to global GxP: GMP, cGMP, GLP, GCP, GEP, GDP, GSP, GDocP and GVP
002	WHO GMP fundamentals and inspection expectations
003	India revised Schedule M: GMP requirements and implementation
004	EU GMP Part I, Part II and relevant annexes
005	US FDA 21 CFR Parts 210 and 211: pharmaceutical cGMP
006	ICH Q7 GMP for active pharmaceutical ingredients
007	ICH Q8, Q9(R1), Q10, Q11 and Q12: lifecycle quality framework
008	PIC/S GMP and international inspection harmonisation
009	MHRA, EMA, US FDA, TGA and CDSCO regulatory expectations
010	GMP organisational roles, key personnel and quality culture
011	Product classification and choice of applicable regulatory framework
012	GxP compliance gap-assessment techniques
013	Global regulatory intelligence and change monitoring
014	Ethics, professional conduct and patient/consumer safety

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02 | Pharmaceutical Quality Management Systems (PQS/QMS)

APPLICATION: All regulated manufacturing; emphasis pharmaceuticals/biotech

KEY REFERENCES: ICH Q10; EU GMP Chapter 1; WHO GMP; ISO 9001; ISO 13485 where applicable

Sr. No.	TRAINING TOPIC
015	Pharmaceutical Quality System establishment and governance
016	Quality policy, objectives, management review and KPIs
017	Quality assurance versus quality control responsibilities
018	Quality Risk Management (QRM) under ICH Q9(R1)
019	Risk assessment tools: FMEA, FTA, HACCP and risk ranking
020	Deviation and nonconformance management
021	Root cause analysis: 5 Whys, Ishikawa and fault-tree methods
022	Corrective and Preventive Action (CAPA) lifecycle
023	Change control: planned, emergency and temporary changes
024	Out-of-Specification (OOS) investigation and laboratory failures
025	Out-of-Trend (OOT) and atypical trend investigations
026	Complaints, product defects and quality investigations
027	Product recall, mock recall and traceability exercises
028	Annual Product Quality Review / Product Quality Review (APQR/PQR)
029	Supplier qualification, vendor audits and quality agreements
030	Outsourced activities, contract manufacturing and testing controls
031	Continual improvement, knowledge management and management review
032	Quality metrics, inspection signals and escalation management

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03 | GDocP, Records, Data Integrity & Digital GxP

APPLICATION: All sectors and laboratories

KEY REFERENCES: EU GMP Chapter 4 and Annex 11; FDA 21 CFR Part 11; WHO data integrity guidance; ALCOA+

Sr. No.	TRAINING TOPIC
033	Good Documentation Practice (GDocP): principles and common errors
034	ALCOA+ data integrity principles and practical examples
035	Controlled document hierarchy: policies, SOPs, forms and records
036	SOP writing, approval, effective date and periodic review
037	Good recording practice: corrections, dates, time stamps and signatures
038	Batch Manufacturing and Batch Packaging Records (BMR/BPR)
039	Electronic records and electronic signatures: 21 CFR Part 11
040	GxP computerised systems under EU GMP Annex 11
041	Audit trails: review strategy, exception management and retention
042	Data governance, metadata, access management and backup
043	Laboratory raw-data control and chromatographic integration
044	eQMS, LIMS, ERP, MES and electronic batch records
045	AI-enabled tools: risk assessment, human oversight and validation
046	Document lifecycle, retention, archiving and secure disposal
047	Data integrity inspection case studies and remediation

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04 | GLP, Quality Control & Analytical Laboratories

APPLICATION: Pharma, biotech, medical device, cosmetics, nutraceutical testing

KEY REFERENCES: OECD GLP for nonclinical studies; GMP QC; ISO/IEC 17025; pharmacopoeias

Sr. No.	TRAINING TOPIC
048	OECD Good Laboratory Practice (GLP) principles for nonclinical safety studies
049	Distinguishing OECD GLP, GMP quality-control laboratories and ISO/IEC 17025
050	Laboratory organisation, independent QA unit and study director roles
051	Study plans, amendments, raw data, final reports and archiving
052	Analytical method development and analytical control strategy
053	Analytical method validation (ICH Q2(R2)) and verification
054	Analytical procedure development (ICH Q14)
055	HPLC, UPLC, GC, UV-Vis, FTIR and dissolution system practices
056	Calibration, measurement uncertainty and metrological traceability
057	Specifications, reference standards and working standards
058	Pharmacopoeial compliance: USP, Ph. Eur., BP and IP
059	OOS/OOT investigation and atypical analytical result handling
060	Good Microbiology Laboratory Practices (GMLP) and aseptic techniques
061	Microbial limits, bioburden, sterility and bacterial endotoxin testing
062	Environmental and personnel monitoring for laboratory operations
063	Laboratory reagent, culture media and reference culture management
064	Stability studies, sample management and retention samples
065	NABL / ISO/IEC 17025 laboratory accreditation readiness

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05 | Good Engineering Practice (GEP), Facilities & Utilities

APPLICATION: Manufacturing sites, projects, cleanrooms and labs

KEY REFERENCES: GEP industry practice; EU GMP Annex 1; ISO 14644; WHO HVAC/water GMP guidance

Sr. No.	TRAINING TOPIC
066	Good Engineering Practice (GEP) and engineering quality management
067	User Requirement Specifications (URS) and Design Basis Reports (DBR)
068	Site master planning and GMP-compliant facility layouts
069	Personnel, material, waste and equipment movement flows
070	Cleanroom classification, ISO 14644 and zoning concepts
071	HVAC fundamentals: temperature, humidity, pressure and airflow
072	Air handling units, HEPA filters, airflow patterns and recovery
073	Cleanroom pressure cascades, airlocks and interlocks
074	Cleanroom qualification and requalification strategy
075	Purified Water (PW), Water for Injection (WFI) and water-loop basics
076	Clean steam, compressed air, nitrogen and process gases
077	Utilities commissioning, operation, maintenance and monitoring
078	Building/Environmental Monitoring Systems (BMS/EMS)
079	Preventive maintenance, breakdown management and spare parts
080	Facility contamination control, pest prevention and cleaning
081	Energy-efficient HVAC, utility optimisation and GMP sustainability

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06 | Qualification, Validation & Quality by Design

APPLICATION: Pharma, biotech, devices and regulated systems

KEY REFERENCES: EU GMP Annex 15; FDA process validation; ICH Q8/Q9/Q10; ASTM E2500 concepts

Sr. No.	TRAINING TOPIC
082	Validation Master Plan (VMP) preparation and implementation
083	Validation lifecycle and risk-based validation planning
084	Design Qualification (DQ), Installation Qualification (IQ)
085	Operational Qualification (OQ) and Performance Qualification (PQ)
086	Factory Acceptance Testing (FAT) and Site Acceptance Testing (SAT)
087	Equipment qualification and commissioning-to-qualification transfer
088	Process validation: lifecycle stages and continued process verification
089	Cleaning validation and cleaning verification
090	Health-Based Exposure Limits (HBEL), PDE and MACO calculations
091	HVAC qualification and cleanroom performance testing
092	Water, clean steam and compressed gas system qualification
093	Autoclave, dry-heat, depyrogenation and sterilisation validation
094	Computerised System Validation (CSV) and CSA approaches
095	Spreadsheet validation and laboratory software qualification
096	Analytical method transfer and method validation
097	Packaging process and container-closure integrity validation
098	Hold-time studies and transport qualification
099	Continued verification, revalidation and change-impact assessment

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07 | Pharmaceutical Manufacturing: Non-Sterile Dosage Forms

APPLICATION: Tablets, capsules, liquids, ointments, creams, powders, inhalations

KEY REFERENCES: EU GMP Parts I/Annex 9; WHO GMP; US FDA 21 CFR 210/211; revised Schedule M

Sr. No.	TRAINING TOPIC
100	Solid oral dosage manufacturing: tablets and hard capsules
101	Wet granulation, dry granulation and direct compression
102	Blending, segregation control and blend uniformity
103	Tablet compression, coating, friability and dissolution
104	Capsule filling: powders, pellets, granules and liquids
105	Oral liquid manufacturing: syrups, suspensions and emulsions
106	Topical semisolids: creams, ointments, gels and lotions
107	Powders, sachets, effervescent tablets and oral granules
108	In-process controls and yield/reconciliation management
109	Cross-contamination prevention and campaign production
110	Line clearance, status labelling and mix-up prevention
111	Scale-up, technology transfer and process troubleshooting
112	Packaging: blisters, strips, bottles, tubes and sachets
113	Extractables/leachables and packaging compatibility basics
114	Product complaints related to dosage-form failures
115	Dose uniformity, dissolution and stability failure case studies

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08 | Sterile Products, Aseptic Processing & Contamination Control

APPLICATION: IV fluids, ampoules, vials, lyophilised products and sterile medical products

KEY REFERENCES: EU GMP Annex 1; WHO TRS 1044 Annex 2; ISO 14644; applicable FDA aseptic guidance

Sr. No.	TRAINING TOPIC
116	Sterile manufacturing fundamentals and microbiological contamination risks
117	Contamination Control Strategy (CCS) design and governance
118	EU GMP Annex 1: technical expectations and implementation
119	Cleanroom Grades A/B/C/D and ISO cleanroom relationships
120	Aseptic technique, gowning qualification and behaviour
121	Aseptic process simulation / media fill design and review
122	Sterilising filtration and filter integrity testing
123	Autoclave moist-heat sterilisation and biological indicators
124	Dry-heat sterilisation and depyrogenation tunnel validation
125	Vial, ampoule and prefilled syringe washing and preparation
126	Vial filling, stoppering, crimping and container closure integrity
127	Lyophilisation process, equipment and cycle development
128	Terminal sterilisation versus aseptic processing decisions
129	Environmental monitoring: viable/non-viable trends and investigations
130	Personnel monitoring, intervention management and aseptic observations
131	Barrier technology: RABS, isolators and glove integrity
132	Water-for-injection, clean steam and clean utility controls
133	Sterility assurance investigations and regulatory observations

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09 | API, Chemical Synthesis & Highly Potent Products

APPLICATION: APIs, KSMs, intermediates, beta-lactams, oncology and hormones

KEY REFERENCES: ICH Q7/Q11; EU GMP Part II; WHO API GMP; occupational safety frameworks

Sr. No.	TRAINING TOPIC
134	API GMP under ICH Q7 and EU GMP Part II
135	Starting-material selection, KSM definition and supplier controls
136	Chemical synthesis and reaction process controls
137	Crystallisation, filtration, centrifugation and drying
138	Solvent recovery, solvent residues and ICH Q3C
139	Elemental impurities, genotoxic impurities and nitrosamine control
140	Process impurities, degradation products and impurity profiling
141	API process validation, critical process parameters and control strategy
142	High Potency API (HPAPI) containment and occupational exposure
143	Beta-lactam, cephalosporin, hormonal and cytotoxic product segregation
144	Cleaning verification and cross-contamination risk in multipurpose sites
145	API packaging, storage, stability and retest periods
146	DMF/ASMF/CEP concepts and API regulatory documentation
147	API technology transfer, lifecycle change control and audits

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10 | Biotechnology & Biopharmaceutical GMP

APPLICATION: Vaccines, mAbs, recombinant proteins, biosimilars, cell/gene therapy

KEY REFERENCES: EU GMP Annex 2; ICH Q5 series/Q6B; WHO biological product GMP; applicable ATMP guidance

Sr. No.	TRAINING TOPIC
148	Biotechnology GMP principles and biologics regulatory framework
149	Cell line development, characterisation and cell bank management
150	Upstream processing: bioreactors, fermentation and perfusion
151	Downstream processing: chromatography and ultrafiltration
152	Bioprocess aseptic operations and contamination prevention
153	Bioburden control and closed single-use processing
154	Single-use systems: extractables, leachables and integrity
155	Viral safety: viral clearance, adventitious agents and biosafety
156	Process analytical technology (PAT) in bioprocessing
157	Biologics process validation and comparability assessments
158	Biosimilars: quality, analytical similarity and comparability
159	Vaccine manufacture, aseptic filling and cold-chain integration
160	Monoclonal antibody manufacturing and critical quality attributes
161	Cell therapy and gene therapy GMP / ATMP concepts
162	Bioburden, endotoxin and sterility assurance in biologics
163	Biological potency testing, bioassays and assay validation
164	Biological product stability and freeze-thaw studies
165	Biotechnology batch records, deviations and investigation scenarios

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11 | Research, GCP & Clinical Laboratory Practice

APPLICATION: Clinical trials, CROs, R&D and nonclinical study teams

KEY REFERENCES: ICH E6(R3); ICH E8(R1); OECD GLP; GCLP best-practice frameworks; local clinical-trial rules

Sr. No.	TRAINING TOPIC
166	Good Clinical Practice (GCP) principles under ICH E6(R3)
167	ICH E8(R1): quality by design in clinical studies
168	Sponsor, investigator, CRO and ethics committee responsibilities
169	Clinical trial protocol design, amendments and deviations
170	Informed consent, participant rights and research ethics
171	Clinical trial monitoring, risk-based quality management and oversight
172	Clinical study data integrity, eSource, eCRF and eTMF
173	Safety reporting, SAEs, SUSARs and protocol safety oversight
174	Clinical trial investigational product GMP and accountability
175	Decentralised and pragmatic clinical trial considerations
176	Clinical study audits, inspections and CAPA
177	Good Clinical Laboratory Practice (GCLP): sample handling and testing
178	OECD GLP: nonclinical toxicology study conduct
179	Bioanalytical method validation and regulated bioanalysis
180	CTRI registration and India clinical-trial regulatory orientation

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12 | Good Distribution, Storage, Pharmacovigilance & Supply Chain

APPLICATION: Pharma, biotech, devices and temperature-sensitive products

KEY REFERENCES: EU GDP; WHO storage/distribution guidance; GVP where applicable

Sr. No.	TRAINING TOPIC
181	Good Distribution Practice (GDP) and Good Storage Practice (GSP)
182	Warehouse qualification, mapping and zoning
183	Temperature mapping, monitoring and excursion investigation
184	Cold chain logistics for vaccines, biologics and diagnostics
185	Good transportation practice and route risk assessment
186	Transport packaging validation and data logger management
187	Inventory management: FEFO/FIFO, segregation and status control
188	Counterfeit prevention, serialisation and traceability
189	Distributor qualification, third-party logistics and quality agreements
190	Import/export quality management and border shipment controls
191	Good Pharmacovigilance Practice (GVP) system fundamentals
192	Adverse event case processing, signal detection and reporting
193	Post-marketing surveillance, recalls and field safety notices
194	Product complaint and returns management in distribution

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13 | Medical Devices & IVD Quality and Regulatory Training

APPLICATION: Medical devices, implants, IVDs, sterile/single-use products

KEY REFERENCES: ISO 13485; ISO 14971; US FDA QMSR; EU MDR/IVDR; India MDR 2017

Sr. No.	TRAINING TOPIC
195	ISO 13485:2016 medical device QMS implementation
196	US FDA Quality Management System Regulation (QMSR): 2026 requirements
197	India Medical Devices Rules, 2017: licensing and compliance
198	EU MDR 2017/745 regulatory and conformity assessment overview
199	EU IVDR 2017/746 regulatory and conformity assessment overview
200	Medical device and IVD classification and intended purpose
201	ISO 14971:2019 medical device risk management lifecycle
202	ISO/TR 24971: risk management application guidance
203	Design and development controls, design reviews and design transfer
204	Design verification, design validation and usability engineering
205	ISO 62366-1 usability engineering and human factors
206	IEC 62304 software lifecycle and software validation basics
207	Medical device cybersecurity and software as a medical device
208	ISO 10993 biological evaluation and biocompatibility planning
209	Sterile device packaging and ISO 11607 validation
210	Sterilisation validation: ISO 11135, 11137 and 17665 overview
211	Cleanroom controls and particulate contamination for devices
212	UDI, labelling, IFU and traceability requirements
213	Clinical evaluation, PMCF, PMS and vigilance
214	Device technical documentation, CER and performance evaluation
215	Device CAPA, complaints and field safety corrective actions

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Sr. No.	TRAINING TOPIC
216	Supplier controls, production process validation and audit readiness

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14 | Cosmetics & Personal Care GMP / Compliance

APPLICATION: Cosmetics, skin care, hair care and personal-care plants

KEY REFERENCES: ISO 22716; EU Regulation 1223/2009; India Cosmetics Rules; applicable US requirements

Sr. No.	TRAINING TOPIC
217	ISO 22716 cosmetics Good Manufacturing Practices
218	India Cosmetics Rules and manufacturing/import licensing overview
219	EU Cosmetics Regulation (EC) No 1223/2009 requirements
220	US cosmetics regulation and MoCRA compliance awareness
221	Cosmetic product classification, intended use and claims
222	Cosmetic ingredient selection, restrictions and safety review
223	Cosmetic Product Safety Report (CPSR) and safety assessment
224	Product Information File (PIF) and CPNP notification
225	Cosmetic formulation development, scale-up and process controls
226	Preservative efficacy / challenge testing and microbiological quality
227	Stability, packaging compatibility and shelf-life studies
228	Manufacturing controls for creams, lotions, shampoos and gels
229	Cosmetic labelling, INCI names and artwork approval
230	Claims substantiation, marketing claims and advertising compliance
231	Organic/natural cosmetics: COSMOS / ECOCERT concepts
232	Cosmetovigilance, complaints, recalls and adverse event handling

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15 | Nutraceuticals, Dietary Supplements & Food Safety

APPLICATION: Nutraceuticals, supplements, botanical products, food and herbal processing

KEY REFERENCES: FSSAI requirements; US dietary supplement 21 CFR Part 111; ISO 22000; FSSC 22000; BRCGS

Sr. No.	TRAINING TOPIC
233	FSSAI nutraceuticals and health supplements regulatory framework
234	Food safety fundamentals and Good Hygiene Practices (GHP)
235	Hazard Analysis and Critical Control Points (HACCP)
236	ISO 22000:2018 Food Safety Management Systems
237	FSSC 22000 certification scheme fundamentals
238	BRCGS Global Standard for Food Safety awareness
239	US dietary supplement CGMP: 21 CFR Part 111
240	Ingredient qualification: botanicals, vitamins and minerals
241	Allergen management and cross-contact prevention
242	Mycotoxins, heavy metals, pesticides and contaminant control
243	Food microbiology, sanitation, CIP and hygiene monitoring
244	Pre-requisite programmes (PRPs) and operational PRPs
245	Nutraceutical manufacturing: capsules, tablets, powders and liquids
246	Herbal extraction, solvent handling and botanical traceability
247	Stability, shelf-life testing and packaging protection
248	Labelling, nutrition/supplement facts and claims compliance
249	Food recall, food defence and food fraud vulnerability assessment
250	HALAL, KOSHER and organic certification readiness

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16 | Regulatory Affairs, Approvals & Certification Readiness

APPLICATION: All sectors; India, EU, UK, US and export markets

KEY REFERENCES: CDSCO; MHRA; EMA; US FDA; TGA; FSSAI; EU MDR/IVDR and related regulatory authorities

Sr. No.	TRAINING TOPIC
251	GMP inspection readiness for US FDA, WHO, EMA, MHRA and TGA
252	CDSCO / State FDA pharmaceutical manufacturing and testing licences
253	WHO GMP and export-market inspection preparedness
254	EU GMP certification, Qualified Person and batch release concepts
255	Common Technical Document (CTD), eCTD and ACTD fundamentals
256	Regulatory dossier modules and CMC documentation
257	EU/US pharmacopoeial and product registration strategy
258	Medical device regulatory submissions and technical files
259	Cosmetics registration, notification, responsible person and PIF
260	FSSAI licensing, product standards and compliance documentation
261	NABL / ISO/IEC 17025 accreditation preparation
262	DSIR in-house R&D recognition and R&D documentation readiness
263	DSIR-SIRO recognition and scientific research organisation governance
264	BIS standards, WPC/ETA and relevant product approvals overview
265	MSME / Udyam, IEC, DPIIT and ZED awareness for regulated firms
266	Environmental approvals, factory licence, Fire NOC and pollution control
267	Regulatory correspondence, deficiency responses and commitments
268	Master audit roadmap, inspection war room and remediation governance

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17 | EHS, Biosafety, Environmental & Occupational Compliance

APPLICATION: All manufacturing, engineering and laboratory sites

KEY REFERENCES: Applicable local EHS law; ISO 14001/45001; biosafety and pollution control guidance

Sr. No.	TRAINING TOPIC
269	Occupational health, safety and environmental compliance fundamentals
270	ISO 14001 Environmental Management System awareness
271	ISO 45001 Occupational Health & Safety Management System
272	Chemical safety: SDS, GHS labelling and safe storage
273	Laboratory biosafety, biosecurity and biological waste
274	Fire safety, emergency planning and evacuation drills
275	Hazardous waste, effluent treatment and emissions control
276	Containment and exposure controls for potent/sensitising compounds
277	Confined space, permit-to-work, lockout/tagout and contractor safety
278	Ergonomics, behaviour-based safety and incident investigation
279	Environmental monitoring, water conservation and energy management
280	Risk assessments for change projects and construction in live plants

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18 | Auditing, Training Effectiveness & Leadership

APPLICATION: QA/QC, production, engineering, regulatory and management teams

KEY REFERENCES: ISO 19011; applicable sector QMS/GMP inspection expectations

Sr. No.	TRAINING TOPIC
281	Internal quality auditing based on ISO 19011 principles
282	GMP self-inspections and mock regulatory audits
283	Supplier audits and contract manufacturer audits
284	Regulatory inspection hosting, interviews and document retrieval
285	Regulator observation response writing and evidence packages
286	CAPA effectiveness checks, verification and closure
287	GMP training needs analysis and role-based training matrices
288	Competency assessment, train-the-trainer and requalification
289	Human error prevention, human factors and error-proofing
290	Quality culture, leadership accountability and escalation
291	Practical deviation/OOS/CAPA workshop with case studies
292	GDocP and data integrity hands-on error-identification workshop
293	GMP risk management exercises using FMEA and fishbone
294	KPI dashboards, visual management and operational excellence

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REGULATORY REFERENCE SOURCES

Reference websites retained from the source catalogue. Check the current regulatory editions and applicability before each delivery.

International / GMP sources	Sector and national sources
WHO GMP | who.int/publications/i/item/9789240086081	FDA Device QMSR | fda.gov/medical-devices/quality-and-compliance-medical-devices
EU GMP | health.ec.europa.eu/medicinal-products/eudralex/eudralex-volume-4_en	EU MDR/IVDR | health.ec.europa.eu/medical-devices-new-regulations
FDA Drug GMP | fda.gov/drugs/pharmaceutical-quality-resources	EU Cosmetics | eur-lex.europa.eu (Regulation 1223/2009)
ICH Guidelines | admin.ich.org/page/quality-guidelines	FDA Dietary CGMP | fda.gov (21 CFR Part 111)
OECD GLP | oecd.org/en/topics/sub-issues/testing-of-chemicals	FSSAI Supplements | fssai.gov.in/standards/health-supplements
EMA GCP | ema.europa.eu/en/scientific-guidelines/ich-e6-good-clinical-practice	India Schedule M | pib.gov.in (Press release PRID 2244523)

IMPORTANT NOTE

Training does not confer accreditation or approval. OECD GLP covers regulated nonclinical studies; GMP laboratory controls, GCLP and GEP have different scopes. Confirm latest rules and standards for each market.

CONTACT

ISHA'S GMP Consultancy Pvt. Ltd. (India) | ISHA'S GMP Turnkey Consultancy Europe Ltd. (Ireland)

www.ishasgmp.com | info@ishasgmp.com | India & Ireland

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THANK YOU | GET IN TOUCH

India and Ireland - Global GMP and turnkey project partners

INDIA

ISHA'S GMP CONSULTANCY PRIVATE LIMITED

Chennai Head Office
Regency Flat, Sri Hi-Tech Construction
55/66, 1st Cross Street, CRR Puram
Manapakkam, Chennai – 600125, India

PHONE +91 86675 89984 | +91 99764 91578

EMAIL info@ishasgmp.com

WEB www.ishasgmp.com

Regional office: 338, 3rd Floor, Shoppers Gate,
Vapi–Chala Road, Gujarat – 396191

IRELAND / EUROPE

ISHA'S GMP TURNKEY CONSULTANCY EUROPE LTD

Ireland Office
3 Graydon Green
Newcastle, Dublin, D22 Y9V9
Ireland

PHONE +353 89 989 1842

EMAIL info@ishasgmpeu.ie

WEB www.ishasgmpeu.ie

European GMP, regulatory compliance
and turnkey project consultancy

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