Bealink

Callimedia
by Bealink

Présentation de notre partenaire

GXP training

2025

NOUVEAU PARTENARIAT

GXP training x Callimedia by Bealink

Des solutions de formation certifiées et performantes pour les professionnels des secteurs des sciences de la vie.

GxP Training en quelques mots

GxP Training est un catalogue de cours en ligne **accrédités**, conçu pour aider **les professionnels de la qualité** à répondre à leurs exigences réglementaires.

20
ans d'activité

+2000
clients

1.5M
apprenants formés

98%
de satisfaction

Les secteurs professionnels concernés :

- Pharmacie
- Agroalimentaire
- Excipients
- Recherche clinique
- Dispositifs médicaux

Ils leur font déjà confiance :

AstraZeneca · Audika · IQVIA · MERCK · NHS · NOVARTIS

Une expertise reconnue

La conception des modules suit une méthodologie éprouvée afin d'assurer **une conformité avec les normes internationales.**

CONCEPTION PAR DES EXPERTS

- Nos cours sont écrits par **des professionnels expérimentés** en affaires réglementaires, médecine ou pharmacologie.
- Le contenu est validé par **un conseil scientifique** pour garantir la qualité.

CONTRÔLE QUALITÉ SYSTÉMATIQUE

- Intégration des **contraintes locales** pour assurer la conformité.
- Process de validation **rigoureux** à 3 étapes.
- Mises à jour du contenu des modules **2 fois par an.**

CERTIFICATIONS ET ACCRÉDITATIONS

Accredited cpd

IACET The Standard for Lifelong Learning ACCREDITED

FDA 21 CFR Part 11

Information Security Management ISO 27001 Certified

MISES À JOUR

Les cours sont mis à jour régulièrement en fonction des **nouvelles normes de conformité** émises par les organismes de santé publique.
Fréquence : 1 fois par an ou par trimestre

Évaluations

Utilisez les évaluations intelligentes pour **tester réellement** les connaissances acquises, tout au long de la formation :

AVANT → **PENDANT** → **APRÈS**

Au choix parmi 7 types d'évaluations :

- Vrai / Faux
- QCM
- QRM
- Classez les réponses
- Clics sur images
- Associations
- Textes à trous

Certifications

Grâce à l'acccréditation du Groupe CPD, obtenez **la certification CPD/CEU** grâce aux modules GxP.

- Tous les certificats sont générés depuis la plateforme LMS GxP
- L'apprenant obtient le certificat automatiquement à la fin d'un module s'il valide l'évaluation.
- Ils sont uniques, authentiques, traçables et conformes à la norme **FDA 21 CFR PART 11.**

Catalogue de formations

Catalogue de formations
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Un catalogue riche, qui évolue constamment

+85
modules

+130
heures de formation

24/7
accessibilité du catalogue

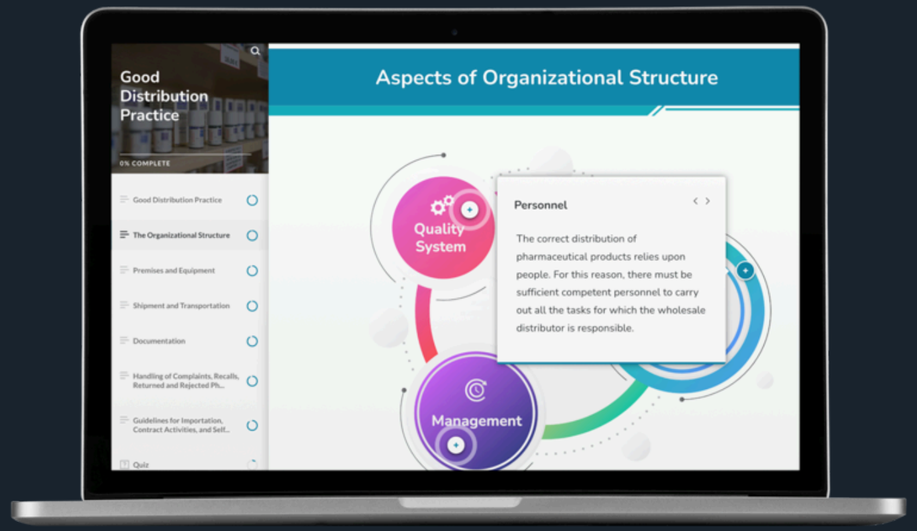
+12
mises à jour par mois

Choisissez le mode de déploiement adapté à vos besoins

Intégration des modules SCORM

Possibilité d'intégrer des cours via SCORM dans votre propre LMS, SIRH ou logiciel de formation dédié.

Plateforme LMS GxP



- 100% en ligne
- Accès unique par apprenant conforme RGPD
- Reportings détaillés
- Accès 24/7, appli mobile

Les catégories de modules en ligne

- ⚙️ GMP : Good Manufacturing Practices
- 🔬 GCP : Clinical Trials
- 🚚 GDP : Good Distribution Practices
- 🧪 GLP : Good Laboratory Practices
- 💊 Pharmacovigilance
- 🔎 Quality Assurance
- 🖋️ Regulatory Affairs

Les modules disponibles

Name	Categories
Selection of Trial Sites	GCP : Clinical Trials
Registration of Trials on clinicaltrials.gov	GCP : Clinical Trials
Management of Clinical Trial Protocols	GCP : Clinical Trials
Management of Externally Sponsored Studies	GCP : Clinical Trials
Investigational Medicinal Product Order for Clinical Studies	GCP : Clinical Trials
Management of The TMF and ISF	GCP : Clinical Trials
First in Human Safety Assessment (FIHSA) Dossier	GCP : Clinical Trials
Management of Clinical Study Reports	GCP : Clinical Trials
Clinical Trials : Preparing for an audit or an inspection	GCP : Clinical Trials
Good Clinical Practice Training Refresher 2025: ICH-GCP E6(R3) Revision	GCP : Clinical Trials
Human Biological Samples Handling	GCP : Clinical Trials, GDP : Good Distribution Practices
Anti-Bribery and Anti-Corruption Certified Training Course : ISO37001	GCP : Clinical Trials, GDP : Good Distribution Practices, GLP : Good Laboratory Practices, GMP : Good Manufacturing Practices, Pharmacovigilance, Quality Assurance, Regulatory Affairs
Good Clinical Laboratory Practice (GCLP)	GCP : Clinical Trials, GLP : Good Laboratory Practices
21 CFR Part 58: GLP for Nonclinical Laboratories	GCP : Clinical Trials, GLP : Good Laboratory Practices
Laboratory Management Systems – ISO/IEC 17025:2017	GCP : Clinical Trials, GLP : Good Laboratory Practices
General Data Protection Regulation (GDPR) for Pharmaceuticals and Clinical Trials	GCP : Clinical Trials, GLP : Good Laboratory Practices
Regulatory Compliance Inspections and External Audits	GCP : Clinical Trials, GMP : Good Manufacturing Practices, Medical Devices, Regulatory Affairs
GMP for clinical trials manufacture and supply	GCP : Clinical Trials, GMP : Good Manufacturing Practices, Regulatory Affairs

Les modules disponibles

Name	Categories
Pharmacovigilance Case Handling	GCP : Clinical Trials, Pharmacovigilance
Introduction to Non-Interventional Studies (NIS)	GCP : Clinical Trials, Pharmacovigilance
Clinical Management of Non-Interventional Studies	GCP : Clinical Trials, Pharmacovigilance, Quality Assurance
Quality Management System for Pharmaceuticals Course	GCP : Clinical Trials, Pharmacovigilance, Quality Assurance
Clinical Data Management	GCP : Clinical Trials, Quality Assurance, Regulatory Affairs
ICH-GCP: Introduction to Good Clinical Practice	GCP : Clinical Trials, Regulatory Affairs
21 CFR PART 50 Informed Consent of Human Subjects	GCP : Clinical Trials, Regulatory Affairs
Randomization & Blinding in Clinical Research Trials	GCP : Clinical Trials, Regulatory Affairs
Introduction to Current Good Practice (CGXP)	GDP : Good Distribution Practices, GLP : Good Laboratory Practices, Quality Assurance, Regulatory Affairs
Pharmaceutical Warehouses / Storage Practices	GDP : Good Distribution Practices, GMP : Good Manufacturing Practices
Management of Service Providers	GDP : Good Distribution Practices, GMP : Good Manufacturing Practices, Quality Assurance
Introduction to FDA 21 CFR PART 11 Online Course	GDP : Good Distribution Practices, GMP : Good Manufacturing Practices, Quality Assurance, Regulatory Affairs
Good Distribution Practice	GDP : Good Distribution Practices, Medical Devices
Introduction to Good Laboratory Practice (GLP)	GLP : Good Laboratory Practices
OSHA Lab Safety Certified Training	GLP : Good Laboratory Practices
Cleaning Validation Course	GLP : Good Laboratory Practices, GMP : Good Manufacturing Practices

Les modules disponibles

Name	Categories
Corrective Action and Preventive Action (CAPA) Training	GMP : Good Manufacturing Practices
Introduction To Good Manufacturing Practice (GMP)	GMP : Good Manufacturing Practices
GMP for Food Safety and ISO 22000	GMP : Good Manufacturing Practices
Computer System Validation (CSV)	GMP : Good Manufacturing Practices
GMP for Cosmetics and ISO 22716	GMP : Good Manufacturing Practices
CFRs Part 210 and 211 : cGMPs for Finished Pharmaceuticals	GMP : Good Manufacturing Practices
GMP for Medical Devices and FDA 21 CFR PART 820	GMP : Good Manufacturing Practices, Medical Devices
Quality Management for Medical Devices - ISO 13485:2016	GMP : Good Manufacturing Practices, Medical Devices, Quality Assurance
IVDR Training : In Vitro Diagnostic Medical Devices Regulation 2017/746	GMP : Good Manufacturing Practices, Medical Devices, Regulatory Affairs
Signal Detection & Management in Pharmacovigilance	GMP : Good Manufacturing Practices, Pharmacovigilance
Overview of Qualification and Validation	GMP : Good Manufacturing Practices, Quality Assurance
Introduction to Pharmaceutical Quality	GMP : Good Manufacturing Practices, Quality Assurance
GMP Documentation and Record Retention	GMP : Good Manufacturing Practices, Quality Assurance
Quality Risk Management	GMP : Good Manufacturing Practices, Quality Assurance
Good Documentation Practice (GDocP)	GMP : Good Manufacturing Practices, Quality Assurance
HACCP : Hazard Analysis and Critical Control Point System	GMP : Good Manufacturing Practices, Quality Assurance
GMP for Pharmaceutical Packaging and ISO 15378	GMP : Good Manufacturing Practices, Quality Assurance
CMC: Chemistry, Manufacturing, and Controls	GMP : Good Manufacturing Practices, Quality Assurance
An Introduction to Good Manufacturing Practices in the EU	GMP : Good Manufacturing Practices, Quality Assurance, Regulatory Affairs
Internal GMP Audit & Role of GMP Auditor	GMP : Good Manufacturing Practices, Regulatory Affairs
GMP Refresher 2025	GMP : Good Manufacturing Practices, Regulatory Affairs

Les modules disponibles

Name	Categories
Manufacturing (GMP) and Quality	Learning Paths
Medical Devices	Learning Paths
Distribution (GDP)	Learning Paths
Pharmacovigilance Bundle	Learning Paths
Clinical Trials (GCP)	Learning Paths
GxP Bundle	Learning Paths
Clinical Research Associate (CRA)	Learning Paths
Regulatory Affairs Specialist	Learning Paths
Validation Engineer	Learning Paths
Medical Devices : ISO 13485 Internal Auditor	Medical Devices, Quality Assurance
Medical Device Regulation - MDR 2017/745	Medical Devices, Quality Assurance, Regulatory Affairs
ISO 14971: Risk Management For Medical Devices	Medical Devices, Quality Assurance, Regulatory Affairs
GMP for Medical Devices: EU versus FDA	Medical Devices, Regulatory Affairs

Les modules disponibles

Name	Categories
Introduction to Good Pharmacovigilance Practice (GVP)	Pharmacovigilance
Management of Drug Safety Concern Through a Pharmacovigilance Steering Committee	Pharmacovigilance, Quality Assurance
Management of Periodic Safety Update Reports (PSURs)	Pharmacovigilance, Quality Assurance
Global Pharmacovigilance Awareness	Pharmacovigilance, Quality Assurance
Medical Writing of Periodic Benefit-Risk Evaluation Report (PBRER)	Pharmacovigilance, Quality Assurance
Screening and Management of Literature for Pharmacovigilance Activities	Pharmacovigilance, Quality Assurance, Regulatory Affairs
Introduction to GxP	Pharmacovigilance, Quality Assurance, Regulatory Affairs
Risk Analysis Model for Management of Quality Risk	Quality Assurance
Records, Archiving, and Retention	Quality Assurance
Pharmaceutical Data Integrity : ALCOA and ALCOA+	Quality Assurance
Introduction to Software Validation Assurance	Quality Assurance
Statistical Process Control	Quality Assurance, Regulatory Affairs
How to write effective Standard Operating Procedures (SOPs)	Quality Assurance, Regulatory Affairs
Cybersecurity in Life Sciences	Quality Assurance, Regulatory Affairs
Procedure for Handling Cases of Suspected Serious Breach, Misconduct & Fraud	Regulatory Affairs
Pharmaceutical Regulatory Affairs	Regulatory Affairs
Submitting a New Drug Application in the USA	Regulatory Affairs

Les parcours de formation

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Des packs formation par thématiques

Nos experts ont préparé **des parcours d'apprentissage** regroupant plusieurs modules, sélectionnés pour les professionnels souhaitant **valider leur expertise** dans un domaine.

- 🔬 Clinical Trials
- 🔎 Manufacturing and Quality
- 🚚 Distribution
- 💊 Pharmacovigilance

Clinical Trials

14 h Training Path Director : Dr. Patricia Kay

1. ICH-GCP : Introduction to Good Clinical Practice
2. ICH-GCP R3 Training Refresher 2025
3. Introduction to clinical trial protocols
4. Selection of trial sites
5. Management of the TMF and ISF
6. Management of Clinical Study reports

Recommended additional courses

- Registration of trials on clinicaltrials.gov
- Clinical Data Management
- Clinical trials : Preparing for an audit or an inspection
- Management of externally sponsored studies
- Human Biological Samples Handling
- Clinical Management of NIS
- GMP for Clinical Trials manufacture and supply
- FIHSA Dossier
- Management of Periodic Safety Update Reports

🔎Manufacturing and Quality

12 h Training Path Director : Dr. Ellena J. Jefferson

1. Introduction to Pharmaceutical Quality
2. Introduction to GMP
3. Corrective and Preventive Action (CAPA)
4. Quality Risk Management
5. Overview of qualification and validation
6. Good Documentation Practice

Recommended additional courses

- Records, achiving and retention
- Pharmaceutical Data Integrity
- Regulatory Compliance Inspections and External Audits
- Management of Service Providers
- GMP Documentation and record retention
- GMP for clinical trials manufacture and supply
- Statiscal Process Control
- Introduction to GMP in the EU

🚚 Distribution

10 h Training Path Director : Khurram Rehman

1. GDP : Introduction to Good Distribution Practice
2. Human Biological Samples Handling
3. Management of Service Providers
4. Introduction to Current Good Practices (cGxP)
5. Pharmaceutical Warehouses Practicesew of qualification and validation

Recommended additional courses

- Regulatory compliance inspections and external audits
- Good Documentation Practice
- Pharmaceutical Data Integrity : ALCOA and ALCOA+
- Records, Archiving and Retention

🧪 Pharmacovigilance

14 h Training Path Director : Marc Estrow

1. Global Pharmacovigilance Awareness
2. Introduction to Good Pharmacovigilance Practice (GVP)
3. Pharmacovigilance Case Handling
4. FIHSA Dossier
5. Management of Clinical Study Reports
6. How to setup a Pharmacovgilance Steering Committee
7. Management of Literature for Pharmacovigilance Activities

Recommended additional courses

- Introduction to cGxP
- Clinical Data Management
- Clinical trials : Preparing for an audit or an inspection
- Management of externally sponsored studies
- Human Biological Samples Handling
- Clinical Management of NIS
- GMP for Clinical Trials manufacture and supply
- FIHSA Dossier
- Management of Periodic Safety Update Reports

Merci.

Je reste à votre disposition.

Christophe FABRE

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