Training Calendar

Courses

20 and 24 February 2025 *Online*	**Reporting Requirements in Veterinary Pharmacovigilance** Animal Health / Pharmacovigilance
20 and 21 February 2025 *online*	**Labelling Requirements for Medical Devices** *Understanding the regulatory labelling requirements for medical devices in the context of the MDR 2017/745* *Medical Device*
25 and 26 March 2025 *Online*	**How to Implement the EU CTR in Sponsor Organisations to Ensure Compliance: A Practical Approach** Clinical Research
11, 18 e 20 marzo 2025 *Online*	**La Qualifica dei Fornitori GMP** *Strategie efficaci per il controllo e la gestione ottimali dei propri fornitori* *GMP (Good Manufacturing Practices)*
12 and 13 March 2025 *Online*	**Mastering MDCG 2024-3: Comprehensive Training on Clinical Investigation Plans** *Ensuring Compliance and Excellence in Clinical Investigations of Medical Devices* *Medical Device / Medical Writing*
12 e 14 marzo 2025 *Online*	**La Ricerca della Letteratura Scientifica: dal Quesito ai Risultati** *Principali funzionalità delle banche dati biomediche e degli strumenti di gestione delle ricerche bibliografiche* *Medical Device / Pharmacovigilance / Medical Affairs / Medical Writing / Clinical Research*
19 marzo 2025 *Online*	**Il Consenso quale Base Giuridica per il Trattamento Dati nella Ricerca Clinico-Scientifica** *Eccezioni e regole; legittimo interesse e riutilizzo dei dati* *Medical Affairs / Clinical Research*

20 e 21 marzo 2025 *Milano*	**Dalla Comunicazione Professionale alla Negoziazione** *Tecniche di base per migliorare le interazioni* *Medical Affairs / Clinical Research / Soft Skill*
25 e 26 marzo 2025 *Online*	**Annex 1 e Contamination Control Strategy** *Approcci e applicazioni tecnico-operative per la conformità allo standard* *GMP (Good Manufacturing Practices)*
26 March 2025 *Online*	**Attributability and Causality in Pharmacovigilance: Still a Hot Topic** *How to navigate the different algorithms and approaches used to evaluate the causality of an adverse event/reaction* *Pharmacovigilance*
27 and 28 March 2025 *Online*	**Medical Writing Course: Improve your Writing & Reviewing Skills** *Writing, editing & proofreading tips for medical writers: a standardised process to make your message effective; review & ensure document quality* *Medical Affairs / Medical Writing / Clinical Research*
31 March and 03 April 2025 *Online*	**Searching the Medical Literature** *Best resources and tips for finding medical information* *Medical Device / Pharmacovigilance / Market Access / Medical Affairs / Medical Writing / Clinical Research*
01 and 02 April 2025 *Online*	**Medical Devices Periodic Safety Update Report (PSUR)** *What's behind the MDCG guidelines* *Medical Device*
08 e 10 aprile 2025 *Online*	**Il Responsabile del Servizio Scientifico** *Consapevolezza di un ruolo articolato* *Medical Affairs*
08 April 2025 *Online*	**Select the Ideal PMCF Strategy for a Medical Device** *An advanced training on post-market requirements under the MDR 2017/745* *Medical Device*
08 e 10 aprile 2025 *Online*	**Normativa della Ricerca Clinica tra Presente e Futuro** *Pillole di regolatorio* *Regulatory / Clinical Research*

08, 10, 15 e 17 aprile 2025 *Online*	**Il Sistema di Qualità Applicato alla Farmacovigilanza** *Approfondire il ruolo centrale svolto dall'Assicurazione della Qualità nelle attività di Farmacovigilanza* *Pharmacovigilance*
10 April 2025 *Online*	**Advertisement and Promotion Claims for Medical Devices** *How to ensure regulatory compliance of advertisement and promotion claims and activities in the MedTech industry* *Medical Device / IVDs (In-Vitro Diagnostics) / Regulatory*
11 April 2025 *Online*	**Master the New Swiss Clinical Trial Framework and its Interplay with the EU CTR - Similarities and Differences** Clinical Research
11 aprile 2025 *Online*	**ICH GCP (R3)** *Innovazione, Flessibilità e Qualità nella Ricerca Clinica* *Clinical Research*
14 and 15 April 2025 *Online*	**Mastering the Essentials of Marketing Authorisation Application (MAA): A Path to Regulatory Success in the EU, UK and US** Regulatory
15 aprile 2025 *Online*	**Progettare un Patient Support Program (PSP): Configurazione del Processo e Strumenti** Medical Affairs
15 e 16 aprile 2025 *Online*	**Protezione dei dati personali (GDPR) e Ricerca Scientifica** *L'impatto del GDPR e delle norme locali sulla ricerca clinica; evoluzioni e orizzonti normativi* *Clinical Research*
16 April 2025 *Online*	**From Good to Excellent: The Summary of Safety and Clinical Performance (SSCP)** *An advanced training to improve your SSCP and write it in the most efficient way* *Medical Device*
17 April 2025 *Online*	**Introduction to Aseptic Process Simulation (APS)** *Fundamentals to Designing Compliant Media Fills for Sterility Assurance* *GMP (Good Manufacturing Practices)*

06 maggio 2025 *Online*	**ICH Q9 (R1) Quality Risk Management** *Strumenti per affrontare la gestione del rischio nell'industria farmaceutica* *GMP (Good Manufacturing Practices)*
07 e 08 maggio 2025 *Online*	**Clinical Support Programs: come Sperimentare la Partnership Pubblico Privato** *Strategie e progetti per favorire le sinergie tra soggetti privati e pubblici a supporto degli HCP e del percorso di cura* *Market Access*
07 and 09 May 2025 *Online*	**Pharmacovigilance Agreements** *What pharmaceutical companies should consider when contracting with partners who may receive safety information relevant to their active substances* *Pharmacovigilance*
07 May 2025 *Online*	**Effective Management of SAE/SUSAR Reporting and CIOMS Forms** *Essential Skills for Accurate and Compliant Adverse Reaction Reporting in Medicinal Products* *Pharmacovigilance*
07 e 09 maggio 2025 *Online*	**La Sorveglianza Post-Market dei Dispositivi Diagnostici in Vitro** *Come creare e gestire un sistema efficace ed utilizzare al meglio i dati ottenuti* *IVDs (In-Vitro Diagnostics)*
08 maggio 2025 *Milano*	**Generare Real World Evidence (RWE) con Metodi Osservazionali** *Metodologia e Normativa* *Evidence Generation / Medical Affairs / Clinical Research*
27 e 28 maggio 2025 *Online*	**I Risvolti Pratici del Regolamento (UE) n. 536/2014 sulle Sperimentazioni Cliniche (CTR 536/2014)** *Cosa è cambiato nello sviluppo e nella produzione GMP dei medicinali sperimentali (IMP)* *GMP (Good Manufacturing Practices)*
13 e 15 maggio 2025 *Online*	**Patient Support Program (PSP) e Compliance: quali Normative Considerare?** *Il codice deontologico Farmindustria e i requisiti di privacy e farmacovigilanza per un PSP* *Pharmacovigilance / Market Access / Medical Affairs / Regulatory*

13, 15 e 20 maggio 2025 *Online*	**Patient Advocacy ed Engagement in Ambito Farmaceutico ed Healthcare** *Ruolo e mission in un approccio cross-funzionale* *Medical Affairs*
15, 20 and 22 May 2025 *Online*	**Pharmacovigilance and Safety in Clinical Trials under the Clinical Trial Regulation (EU) No 536/2014** *First experience with CTR/CTIS, regulatory expectations for safety documents, CTR assessment procedures and CTR Q&A document, typical issues and how to avoid it.* *Pharmacovigilance / Regulatory*
15 and 16 May 2025 *Online*	**Labelling Requirements for Medical Devices** *Understanding the regulatory labelling requirements for medical devices in the context of the MDR 2017/745* *Medical Device*
19 and 21 May 2025 *Online*	**Veterinary Marketing Authorisation (MAA) and Variations (MAV) in the EU: Regulatory Requirements and Best Practices** Animal Health / Regulatory
21 maggio 2025 *Online*	**ICH Q14: Lo Sviluppo di Metodi Analitici** *Concetti e approcci per ottenere metodi fit for purpose* *GMP (Good Manufacturing Practices)*
22 maggio 2025 *Online*	**Etichettatura degli Integratori Alimentari - Applichiamo la Normativa ad Esempi Reali** Regulatory
22 and 23 May 2025 *Online*	**Person Responsible for Regulatory Compliance (PRRC) – An MDR/IVDR Requirement** *Responsibilities and challenges for medical device companies within MDR & IVDR* *Medical Device / IVDs (In-Vitro Diagnostics) / Regulatory*
27 e 29 maggio 2025 *Online*	**La Valutazione di Impatto della Protezione dei Dati (DPIA) nella Ricerca Clinica** *Suggerimenti pratici per la redazione della DPIA nello studio clinico* *Clinical Research*
03, 05 e 10 giugno 2025 *Online*	**La Statistica Medica per Non Statistici** *I principi statistici dalla pianificazione della ricerca alla pubblicazione sulle riviste scientifiche* *Medical Device / Medical Affairs / Medical Writing / Clinical Research*

03, 04 e 10 giugno 2025 *Online e in presenza, Milano*	**ICH Q2 (R2): Le Novità nella Convalida dei Metodi Analitici** *Integrare Standard Avanzati e Approcci Statistici per ottenere il metodo fit for purpose* *GMP (Good Manufacturing Practices)*
03 and 05 June 2025 *Online*	**Cosmetics in the EU, US and Canada** *What you need to know to reach the market in compliance* *Regulatory*
04 e 06 giugno 2025 *Online*	**La Gestione del Case Processing in Farmacovigilanza** *Gestione delle segnalazioni di sospette reazioni avverse e aspetti di Qualità nel case processing* *Pharmacovigilance*
05 and 06 June 2025 *Online*	**Clinical Evaluation for Medical Devices** *Understanding the clinical evaluation requirements for the MedTech industry in the context of MDR 2017/745* *Medical Device / Regulatory*
05 June 2025 *online*	**Electronic Submissions and Data Management in Regulatory Affairs** *From eCTD to xEVMPD and IDMP, and how to manage your regulatory information fit for purpose* *Regulatory*
09, 13 e 16 giugno 2025 *Online*	**Computer System Validation (CSV) - GxP Process Owner and Quality Assurance: In or Out?** *Il ruolo del QA e del Process Owner nella convalida dei sistemi computerizzati GxP* *Regulatory / Clinical Research*
11 e 12 giugno 2025 *Online*	**Digital Support Services (D-SS): l'Evoluzione Digitale dei Servizi di Supporto al Paziente per una Migliore Assistenza Terapeutica** Medical Affairs
17 and 19 June 2025 *Online*	**Pharmacovigilance System: Audit & Inspection Readiness** Pharmacovigilance
17 giugno 2025 *Online*	**ICH GCP (R3)** *Innovazione, Flessibilità e Qualità nella Ricerca Clinica* *Clinical Research*

19, 23 and 25 June 2025 *Online*	**A Practical Guide to Innovative Trial Design** Clinical Research / Statistics and Data Management
19 e 20 giugno 2025 *Online*	**I Combination Products ai Sensi del Regolamento UE sui Dispositivi Medici (MDR)** *Una formazione mirata per comprendere il quadro normativo per i prodotti combinati alla luce dell'Art.117 dell'MDR 2017/745 e delle linee guida EMA* *Medical Device / Regulatory*
19 June 2025 *Online*	**Advertisement and Promotion Claims for Medical Devices** *How to ensure regulatory compliance of advertisement and promotion claims and activities in the MedTech industry* *Medical Device / IVDs (In-Vitro Diagnostics) / Regulatory*
23 June 2025 *Online*	**Food Supplement Regulations in the EU** *An introduction to the regulatory landscape* *Regulatory*
24, 25 and 26 June 2025 *Online*	**Tips and Tricks to Improve your Technical/Scientific Writing** *Learn the basic techniques to effectively write technical/scientific documents* *Medical Writing*
27 giugno 2025 *Online*	**Qualifica dei Vendor nella Ricerca Clinica - Approfondimenti** *Esempi e aspetti pratici* *Clinical Research*
02 July 2025 *Online*	**Human Error: the True Root Cause of a Deviation?** *Tips and strategies to understand and manage why human errors occur and to minimize them* *GMP (Good Manufacturing Practices)*
22 and 24 July 2025 *Online*	**Good Distribution Practices (GDP) of Medicinal Products and Medical Devices** *Ensuring compliance from the warehouse to the end user* *Medical Device / GMP (Good Manufacturing Practices) / Regulatory / Clinical Research*
17, 18 and 22 September 2025 *Online*	**How to Write a Clinical Evaluation Plan and Report** *Find your way around the clinical evaluation of a medical device from concept to market approval and beyond* *Medical Device*

17 September 2025 *Online*	**Pharmacovigilance Documents - Basic Concepts and Definitions for Pharmacovigilance Writing** Pharmacovigilance / Medical Writing
18 e 19 settembre 2025 *online*	**La Pubblicità del Farmaco** *Dall'informazione scientifica alla pubblicità al pubblico* *Regulatory*
18, 22 and 23 September 2025 *Online*	**Pharmacovigilance and Safety in Clinical Trials under the Clinical Trial Regulation (EU) No 536/2014** *First experience with CTR/CTIS, regulatory expectations for safety documents, CTR assessment procedures and CTR Q&A document, typical issues and how to avoid it.* *Pharmacovigilance / Regulatory*
22 e 23 settembre 2025 *Online*	**EU GMP Annex 15 - La Qualifica e la Convalida nell'Industria Farmaceutica** *Panorama Normativo, linee guida applicabili e approcci per la conformità di impianti, attrezzature e sistemi* *GMP (Good Manufacturing Practices)*
22 September 2025 *Online*	**The "Global" Qualified Person Responsible for Pharmacovigilance (QPPV) Workshop** *Evolution of the role, challenges and opportunities in a global environment* *Pharmacovigilance*
22 e 23 settembre 2025 *Online*	**La Biocompatibilità e la Caratterizzazione Chimica: Dispositivi Medici Sicuri** *Requisiti normativi, gestione del rischio e valutazione dei dati* *Medical Device / Regulatory*
23, 25 and 30 September 2025 *Online*	**Signal Detection and Signal Management** *Research, identification, interpretation, and management of safety signals* *Pharmacovigilance*
24 September 2025 *Online*	**Pharmacovigilance Documents - Focus on Signal Management and Development Safety Update Reports (DSUR)** Pharmacovigilance / Medical Writing

24 e 25 settembre 2025 *Online*	**ISO 14155/2020 - Come Svolgere uno Studio Clinico con Dispositivi Medici** *Implementazione, Ottemperanza e Chiavi per il Successo* *Medical Device*
24 e 26 settembre 2025 *Online*	**Buone Pratiche per la Gestione dei Documenti Cartacei e dei Dati Elettronici in Ambito GxP** *Strumenti per la conformità alle Good Documentation Practice (GDocP) e ai requisiti di Data Integrity* *Pharmacovigilance / GMP (Good Manufacturing Practices) / Clinical Research*
25 and 29 September 2025 *Online*	**Writing Clinical Study Reports** *Planning and Authoring CSRs in Accordance with Public Disclosure Requirements* *Medical Writing / Clinical Research*
29 September 2025 *Online*	**Veterinary QPPV Oversight** *Best Practices to ensure Compliance & Safety* *Animal Health / Pharmacovigilance*
30 settembre e 01 ottobre 2025 *Milano*	**Comunicare con l'Intelligenza Relazionale** *La flessibilità nelle relazioni per comunicare, guidare e motivare colleghi e interlocutori, in presenza e da remoto* *Soft Skill*
30 settembre e 02 ottobre 2025 *Online*	**Cosmetici - Aspetti Tecnico-Regolatori e Panorama Normativo** Regulatory
01 e 03 ottobre 2025 *Online*	**Market Access: Conoscere e Anticipare le Sfide del Settore Sanità per Garantire un Accesso Omogeneo e Sostenibile ai Farmaci** Market Access
01 October 2025 *Online*	**Pharmacovigilance Documents - Focus on the Risk Management Plan (RMP)** Pharmacovigilance / Medical Writing
01 and 02 October 2025 *Online*	**Labelling Requirements for Medical Devices** *Understanding the regulatory labelling requirements for medical devices in the context of the MDR 2017/745* *Medical Device*

02, 07 e 09 ottobre 2025 *Online*	**Packaging Primario per Farmaci Parenterali: Combinare Aspetti Regolatori e Tecnici per una Somministrazione Sicura ed Efficace** GMP (Good Manufacturing Practices)
07 ottobre 2025 *Online*	**Il Futuro della Raccolta Dati nella Ricerca Clinica: Digital Endpoints e Digital Evidence** *Perché e come usare dispositivi digitali per la raccolta di dati del paziente nella ricerca clinica.* *Medical Affairs / Clinical Research*
07, 09, 14 e 16 ottobre 2025 *Online*	**Selezione e Convalida di Una Soluzione Cloud in Ambito GxP** *Rischi e opportunità nella scelta di una soluzione Cloud a supporto di processi GXP* *Clinical Research*
08 and 09 October 2025 *Online*	**State of the Art Section for Medical Devices – Unpacking the Tips and Tricks of a Complex Document** Medical Device / Medical Writing
08 October and 13 October 2025 *Online*	**Pharmacovigilance Documents – Focus on the Periodic Safety Update Report (PSUR)** Pharmacovigilance / Medical Writing
08 October 2025 *Online*	**Veterinary Pharmacovigilance System** Animal Health / Pharmacovigilance
09 and 10 October 2025 *Online*	**Combination Products under the EU Medical Devices Regulation (MDR)** *A targeted training to understand the regulatory pathway for device drug and drug device combinations in the European Union (EU). Updates from the MDR 2017/745 and EMA guidance* *Medical Device / Regulatory*
13 and 14 October 2025 *Online*	**Mastering Continuous Process Validation in Pharmaceutical Manufacturing** *A Comprehensive Workshop on the Life Cycle Approach to Process Validation* *GMP (Good Manufacturing Practices)*
14 e 21 ottobre 2025 *Online*	**Off-Label, Uso Compassionevole e Accesso Precoce (Early Access)** *Le normative e le loro applicazioni* *Medical Affairs / Regulatory / Clinical Research*

14, 21 October 2025 | 11, 18 November 2025 *Online*	**A Systematic Approach to Real World Evidence (RWE) Generation in Life Sciences - a 2 step intensive course** *How to manage the main methodological, regulatory and operational challenges in generating RWE for approval and access purposes* *Evidence Generation / Medical Affairs / Clinical Research*
15 and 16 October 2025 *Online*	**Medical Devices Periodic Safety Update Report (PSUR)** *What's behind the MDCG guidelines* *Medical Device*
15 e 22 ottobre 2025 *Online*	**Terapie Digitali (DTx): a che Punto Siamo?** *L'innovazione digitale per il futuro delle terapie* *Medical Affairs / Clinical Research*
15 e 16 ottobre 2025 *Online*	**Strumenti di Intelligenza Artificiale e Conformità all'AI Act per Professionisti del Medical Affair: Esercizi ed Esempi Pratici** *Come acquisire le competenze e le conoscenze necessarie per integrare l'intelligenza artificiale nel medical affair garantendo la conformità etica e normativa* *Medical Affairs / Medical Writing*
16 e 17 ottobre 2025 *Online*	**Gestione Efficace delle Deviazioni in Ambito GMP** *Processi, Strumenti e Soluzioni per il Miglioramento Continuo nella Produzione Farmaceutica* *GMP (Good Manufacturing Practices)*
16 October 2025 *Online*	**How to Create Effective Visuals for Better Communicating your Science** Medical Affairs / Medical Writing / Soft Skill
17 ottobre 2025 *Online*	**I Radiofarmaci** Medical Affairs / Regulatory / Clinical Research
20 October 2025 *Online*	**Cosmetovigilance in the EU** *An introduction to cosmetic safety and risk management* *Regulatory*
20, 23 e 27 ottobre 2025 *Online*	**Technology Transfer Farmaceutico** *Strumenti, fasi chiave e case studies per una gestione efficace* *GMP (Good Manufacturing Practices)*

21 and 22 October 2025 *Online*	**Initiating the Development of Artificial Intelligence (AI) Medical Devices** *Developing AI medical devices considering the latest industry expectations* *Medical Device*
22, 23 and 24 October 2025 *Online*	**Clinical Study Protocols – Structure & Content** Medical Device / Medical Affairs / Medical Writing / Clinical Research
22 October 2025 *Online*	**Pharmacovigilance Documents – Focus on Addendum to the Clinical Overview (AddCO) and Referrals** Pharmacovigilance / Medical Writing
23 ottobre 2025 *Online*	**Dal Riconoscimento alla Raccolta degli Insights** *Come il Medical Affairs può fare la differenza* *Medical Affairs*
28, 31 ottobre e 04 novembre 2025 *Online*	**Il Market Access per i Dispositivi Medici** *Dalla pianificazione strategica all'esecuzione dei piani di accesso al mercato* *Medical Device / Market Access*
28 and 29 October 2025 *Online*	**Ensuring Excellence in (Bio)pharmaceutical Cleaning Validation** *A Practical Guide to Contamination Control through Effective Cleaning Strategies and Continuous Process Verification* *GMP (Good Manufacturing Practices)*
28 October 2025 *Online*	**Veterinary Signal Detection** *Requirements and methodology* *Animal Health / Pharmacovigilance*
29 and 30 October 2025 *Online*	**Medical Reading** *The critical evaluation of scientific publications* *Medical Device / Medical Affairs / Clinical Research*
03, 04 and 05 November 2025 *Online*	**Ethylene Oxide Sterilization for Medical Devices: Principles, Validation, and Regulatory Compliance** *Mastering MDR/IVDR Requirements, Contamination Control, and Lifecycle Monitoring for Sterile Medical Devices* *Medical Device / GMP (Good Manufacturing Practices) / Regulatory*

03, 05, 10 e 12 novembre 2025 *Online*	**Safety Management e Farmacovigilanza** *Aspetti normativi, clinici e metodologici della farmacovigilanza, gestione della safety nello sviluppo clinico del farmaco* *Pharmacovigilance / Clinical Research*
04, 06 e 11 novembre 2025 *Online*	**Integratori Alimentari - Aspetti Tecnico-Regolatori e Panorama Normativo** Regulatory
05 and 07 November 2025 *Online*	**Beyond PubMed** *Additional approaches and sources for cost-effective literature monitoring* *Pharmacovigilance / Medical Affairs / Clinical Research*
05, 12 and 19 November 2025 *Online*	**All you Need to Know to Understand Statistics if You are not a Statistician** *Statistical principles from research planning to publication in scientific journals* *Medical Device / Medical Affairs / Medical Writing / Clinical Research*
06 e 07 novembre 2025 *Milano*	**La Gestione dell'Advisory Board nei Progetti Life Science** *Modelli e strumenti* *Medical Affairs / Soft Skill*
06 novembre 2025 *Online*	**Progettare e Comunicare in Medical Affairs** Evidence Generation / Medical Affairs
10 and 11 November 2025 *Online*	**Pre-Market Submission of Medical Devices to the US FDA: Navigating the Pre-Sub and 510(k) Process** *A roadmap of regulatory pathways and tools for EU professionals* *Medical Device / Regulatory*
10, 14 e 17 novembre 2025 *Online*	**Audit to Computer Systems** *Be ready!* *Clinical Research*
11 e 13 novembre 2025 *Online*	**La Vigilanza Post Market per i Dispositivi Medici secondo MDR e FDA** *Dalla gestione del reclamo alla sua notifica come incidente grave alle Autorità Competenti in Europa o come evento avverso in USA. La gestione delle field action in Europa.* *Medical Device / Regulatory*

12 and 14 November 2025 *Online*	**The PSMF (Pharmacovigilance System Master File): from GVPs to Inspections** *A practical, hands-on guide* *Pharmacovigilance*
12 e 19 novembre 2025 *Online*	**Terapie Geniche e Terapie Cellulari** *Dalla Ricerca all'Impiego nel Real World* *Regulatory*
13 November 2025 *Online*	**Veterinary Pharmacovigilance Quality Management** Animal Health / Pharmacovigilance
13 and 18 November 2025 *Online*	**Advanced Therapy Medicinal Product (ATMP): a Roadmap from Classification to Regulation and Manufacturing** GMP (Good Manufacturing Practices) / Regulatory
17 and 18 November 2025 *Online*	**Hands-On Lab on AI Tools and Compliance with AI Act for Medical Affairs Professionals** *Equip yourself with the skills and knowledge to integrate AI into Medical Affairs while ensuring ethical and regulatory compliance.* *Medical Affairs / Medical Writing*
20 and 25 November 2025 *Online*	**Protocol Writing and Communication of Real World Evidence** *International methodological standards for writing and publishing observational study protocols* *Evidence Generation / Medical Affairs / Medical Writing / Clinical Research*
26 November 2025 *Online*	**Decoding Electronic Product Information (ePI)** *An in-depth exploration from paper to ePI* *Regulatory*
26 and 27 November 2025 *Online*	**Introduction to Veterinary Pharmacovigilance** Animal Health / Pharmacovigilance
Data TBD *Online*	**Organizzare Hospital Meeting di Successo: Strategie e Competenze Essenziali** *Corso teorico-pratico per acquisire le competenze chiave nella pianificazione e gestione ottimale di un hospital meeting* *Medical Affairs*

Training Paths

Dal 21 febbraio al 16 aprile 2025 *Online e 1 modulo in presenza a Milano*	**Medical Affairs Essentials** *Corso di alta specializzazione per lo sviluppo delle competenze tecnico-scientifiche del Medical Advisor (MA) e del Medical Science Liaison (MSL)* *Medical Affairs*
From 17 September to 22 October 2025 *Online*	**Pharmacovigilance Documents in the Life Cycle of a Medicinal Product** *From Papers and Patients to Health Authorities* *Pharmacovigilance / Medical Writing*
dal 26 settembre al 22 novembre 2025 *Online*	**Clinical Quality Assurance: un Ruolo Chiave nella Ricerca Clinica** *Percorso di alta specializzazione per lo sviluppo delle competenze peculiari del Clinical Quality Assurance (QA)* *Clinical Research*

Conferences

06 June 2025 *Frankfurt am Main*	**German Pharmacovigilance Day** *Shaping Pharmacovigilance for the Challenges of Tomorrow*
01 ottobre 2025 *Verona*	**Italian Pharmacovigilance Day** *Innovazione ed Efficienza in Farmacovigilanza: Verso un Modello Sostenibile*
13 and 14 October 2025 *Budapest*	**MedDev Day** *Shifting Mindsets: The Future of Global Medical Device Regulation - How Regulatory Changes Are Reshaping Global Business Strategies*
27 and 28 October 2025 *Bergamo*	**European Statistical Forum** *Advancing Clinical Trials: Master Protocols for Faster Drug Development*
11 November 2025 *Copenhagen*	**Nordic Pharmacovigilance Day** *Advancing Pharmacovigilance Together*