Training Calendar

Courses

29 February and 01 March 2024 *Online*	**Labelling Requirements for Medical Devices** *Understanding the regulatory labelling requirements for medical devices in the context of the MDR 2017/745* *Medical Device / Regulatory*
06 and 07 March 2024 *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*
12 and 13 March 2024 *Online*	**Medical Devices Periodic Safety Update Report (PSUR)** *What's behind the MDCG guidelines* *Medical Device*
13 and 15 March 2024 *Online*	**The Next Generation of Medical Affairs: ChatGPT and Large Language Models (LLMs)** *Equip yourself with the skills and knowledge to succeed in the AI-powered medical affairs landscape* *Medical Affairs*
14, 19 e 21 marzo 2024 *Online*	**La Statistica Medica per Non Statistici** *I principi statistici dalla pianificazione della ricerca alla pubblicazione sulle riviste scientifiche* *Medical Affairs / Clinical Research / Statistics and Data Management*
20, 21 and 25 March 2024 *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*

20 March 2024 *Online*	**“Attributability” 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*
26 marzo 2024 *Online*	**Il Consenso quale Base Giuridica per il Trattamento Dati nella Ricerca Clinico-Scientifica** *Deep dive sul consenso, legittimo interesse e riutilizzo dei dati* *Medical Affairs / Regulatory / Clinical Research*
26 e 27 marzo 2024 *Online*	**Annex 1 e Contamination Control Strategy** *Approcci e applicazioni tecnico-operative per la conformità al nuovo standard* *GMP (Good Manufacturing Practices)*
03 e 05 aprile 2024 *Online*	**Le Sfide della Produzione di un Investigational Medicinal Product (IMP)** *Come declinare le richieste delle Good Manufacturing Practices (GMP) nel processo di produzione e controllo del farmaco sperimentale* *GMP (Good Manufacturing Practices)*
03, 04 and 05 April 2024 *Online*	**Clinical Study Protocols – Structure & Content** Medical Affairs / Medical Writing
09 e 11 aprile 2024 *Online*	**Il Responsabile del Servizio Scientifico** *Consapevolezza di un ruolo articolato* *Medical Affairs*
10 and 11 April 2024 *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*
10 aprile 2024 *Online*	**I Radiofarmaci** Medical Affairs / Regulatory / Clinical Research
10, 17 e 24 aprile 2024 *Online*	**Il Market Access per i Dispositivi Medici** *Dalla pianificazione strategica all’esecuzione dei piani di accesso al mercato* *Medical Device / Market Access*

16 e 18 aprile 2024 *Online*	**Normativa della Ricerca Clinica tra Presente e Futuro** *Pillole di regolatorio* *Clinical Research*
18 April 2024 *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) / Medical Affairs / Regulatory*
09 e 10 maggio 2024 *Online*	**Agile Project Management: Gestire l'Innovazione nel Contesto della Produzione Farmaceutica** *I fondamentali di Agile Project Management per affrontare progetti innovativi in realtà di produzione farmaceutica e nel rispetto delle GMP* *GMP (Good Manufacturing Practices)*
23 April 2024 *Online*	**Select the Ideal PMCF Strategy for a Medical Device** *An advanced training on post-market requirements under the MDR 2017/745* *Medical Device*
23 aprile 2024 *Online*	**Progettare un Patient Support Program (PSP): Configurazione del Processo e Strumenti** Medical Affairs
29 April 2024 *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*
07 e 09 maggio 2024 *Online*	**La Pubblicità del Farmaco** *Dall'informazione scientifica alla pubblicità al pubblico* *Regulatory*
08 maggio 2024 *Online*	**Progettare e Comunicare in Medical Affairs** Medical Affairs

08, 15, 22 May and 10, 17 October 2024 *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 using Real World Data (RWD) for regulatory and strategic purposes* *Evidence Generation / Medical Affairs / Clinical Research*
13, 15 and 16 May 2024 *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*
14, 16 e 21 maggio 2024 *Online*	**Patient Advocacy ed Engagement nell'Azienda Farmaceutica** *Ruolo e mission in un approccio inter-funzionale* *Medical Affairs*
15 e 16 maggio 2024 *Online*	**La Sorveglianza Post-Market dei Dispositivi Diagnostici in Vitro** *Come implementare le corrette procedure per la gestione post-vendita* *IVDs (In-Vitro Diagnostics)*
16 and 17 May 2024 *Online*	**Person Responsible for Regulatory Compliance (PRRC) - An MDR/IVDR Requirement** *Responsibilities & challenges for medical device companies within MDR & IVDR* *Medical Device / IVDs (In-Vitro Diagnostics) / Regulatory*
21, 24 and 28 May 2024 *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*
21 and 22 May 2024 *Online*	**Pharmacovigilance Agreements** *What pharmaceutical companies should consider when contracting with partners who may receive safety information relevant to their active substances* *Pharmacovigilance*
22 May 2024 *Online*	**Beyond Slide Decks** *How to use storytelling in your presentations* *Soft Skill*

22 maggio 2024 *Online*	**Etichettatura degli Integratori Alimentari - Applichiamo la Normativa ad Esempi Reali** Regulatory
22 maggio 2024 *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*
23 e 24 maggio 2024 *Milano*	**La Gestione dell'Advisory Board nei Progetti Life Science** *Modelli e strumenti* *Medical Affairs / Soft Skill*
27 May 2024 *Online*	**Oral Presentations** *Skills to help you survive or even shine* *Soft Skill*
28 e 30 maggio 2024 *Online*	**Il Responsabile del Servizio Scientifico** *Consapevolezza di un ruolo articolato* *Medical Affairs*
04 e 06 giugno 2024 *Online*	**General Data Protection Regulation (GDPR), Ricerca Scientifica e Norme Locali** *L'impatto del GDPR sugli studi clinici a livello locale ed internazionale* *Clinical Research*
04, 06 e 11 giugno 2024 *Online*	**Signal Detection e Signal Management** *Ricerca, identificazione, interpretazione e gestione dei segnali di sicurezza* *Pharmacovigilance*
04 June 2024 *Online*	**How to Become a Successfully Published Author - Practical steps to make the publication process as smooth and as successful as possible** Medical Device / Medical Affairs / Medical Writing / Clinical Research
05 e 07 giugno 2024 *Online*	**La Gestione del Case Processing in Farmacovigilanza** *Gestione delle segnalazioni di sospette reazioni avverse e aspetti di qualità nel case processing* *Pharmacovigilance*

05 e 06 giugno 2024 *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*
06, 07, 12 e 14 giugno 2024 *Online*	**Il Sistema di Qualità Applicato alla Farmacovigilanza** *Approfondire il ruolo centrale svolto dall'Assicurazione della Qualità nelle attività di Farmacovigilanza* *Pharmacovigilance*
10 e 17 giugno 2024 *Online*	**Hospital Meeting: come Pianificare e Condurre una Riunione di Successo** *Corso teorico pratico di alta formazione diretto all'apprendimento delle basi di pianificazione e conduzione ottimale di un hospital meeting* *Medical Affairs*
10, 14 e 17 giugno 2024 *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 giugno 2024 *Online*	**Le Good Calibration Practices (GCalP) negli Impianti Automatici di Produzione** *Come fidarsi dei dati che produciamo?* *GMP (Good Manufacturing Practices)*
12 e 13 giugno 2024 *Online*	**Il Patient Support Program (PSP) e la Transizione Digitale: Opportunità e Rischi** Medical Affairs
13 June 2024 *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)*
14 and 21 June 2024 *Online*	**The PSMF (Pharmacovigilance System Master File): from GVPs to Inspections** *A practical, hands-on guide* *Pharmacovigilance*
18 and 20 June 2024 *Online*	**Fundamentals of European Cosmetics Regulatory Affairs** Regulatory

18 and 20 June 2024 *Online*	**Pharmacovigilance System: Audit & Inspection Readiness** Pharmacovigilance
18, 24 and 27 June 2024 *Online*	**A Practical Guide to Innovative Trial Design** Clinical Research / Statistics and Data Management
19 and 20 June 2024 *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*
24 and 26 June 2024 *Online*	**Biologics and Biosimilars Manufacturing** *A practical guide to demystify and address the most common challenges* *GMP (Good Manufacturing Practices)*
25 e 27 giugno, 2 e 4 luglio 2024 *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*
25 and 26 June 2024 *Online*	**Reporting Requirements in Veterinary Pharmacovigilance** Animal Health / Pharmacovigilance
26 giugno 2024 *Online*	**Integratori Alimentari e Advertising: Applichiamo la Teoria alla Pratica** Regulatory
26 and 27 June 2024 *Online*	**Good Distribution Practices of Medical Devices** *Ensuring Compliance* *Medical Device / Regulatory / Clinical Research*
28 giugno 2024 *Online*	**Qualifica dei Vendor nella Ricerca Clinica - Approfondimenti** *Esempi e aspetti pratici* *Clinical Research*
02 July 2024 *Online*	**Statistical Process Control for Pharmaceutical Manufacturing** *Tips for non-statisticians to navigate statistical methods and make data-driven decisions* *GMP (Good Manufacturing Practices)*

02 and 03 July 2024 *online*	**Labelling Requirements for Medical Devices** *Understanding the regulatory labelling requirements for medical devices in the context of the MDR 2017/745* *Medical Device*
16 September 2024 *Online*	**Quality by Design** *How to understand and control process and product variables to ensure quality from the very beginning* *GMP (Good Manufacturing Practices)*
16, 19 and 23 September 2024 *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*
17, 18 and 19 September 2024 *Online*	**Tips and Tricks to Improve your Technical/Scientific Writing** *Learn the basic techniques to effectively write technical/scientific documents* *Medical Writing*
17 and 18 September 2024 *Online*	**Introduction to Pharmacovigilance** *A short but comprehensive guide to the basis of drug safety* *Pharmacovigilance*
17 September 2024 *Online*	**Good Distribution Practices (GDP) for Veterinary Medicinal Products** Animal Health / Regulatory
18 e 19 settembre 2024 *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)*
18 September 2024 *Online*	**Pharmacovigilance Documents - Basic Concepts and Definitions for Pharmacovigilance Writing** Pharmacovigilance / Medical Writing

19 e 20 settembre 2024 *Online*	**La Conduzione di uno Studio Clinico con un Dispositivo Medico-Diagnostico in Vitro** *Dai requisiti regolatori per la sottomissione alla gestione dello studio* *IVDs (In-Vitro Diagnostics) / Clinical Research*
20 settembre 2024 *Online*	**ICH Q9 Quality Risk Management** *Strumenti per la gestione del rischio nell'industria farmaceutica* *GMP (Good Manufacturing Practices)*
23 and 26 September 2024 *Online*	**Searching the Medical Literature** *Best resources and tips for finding medical information* *Medical Device / Pharmacovigilance / Market Access / Medical Affairs / Medical Writing / Clinical Research*
23 e 24 settembre 2024 *Online*	**La Biocompatibilità e la Caratterizzazione Chimica: Dispositivi Medici Sicuri** *Requisiti normativi, gestione del rischio e valutazione dei dati* *Medical Device / Regulatory*
24, 26 settembre e 01 ottobre 2024 *Online*	**Patient Support Program (PSP) e Patient Solution: Compiere una Scelta Strategica a Supporto del Paziente** Market Access / Medical Affairs / Regulatory
24 e 26 settembre 2024 *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*
25 e 27 settembre 2024 *Online*	**Buone Pratiche per la Gestione dei Documenti Cartacei e dei Dati Elettronici in Ambito GxP** *Strumenti per la conformità alle Good Documentation Practice (GDP) e ai requisiti di Data Integrity* *Pharmacovigilance / GMP (Good Manufacturing Practices) / Clinical Research*
25 e 26 settembre 2024 *Online*	**ISO 14155/2020 - Come Svolgere uno Studio Clinico con Dispositivi Medici** *Implementazione, Ottemperanza e Chiavi per il Successo* *Medical Device*
25 September 2024 *Online*	**Pharmacovigilance Documents - Focus on Signal Management and Development Safety Update Reports (DSUR)** Pharmacovigilance / Medical Writing

27 settembre 2024 *Online*	**Codice Deontologico di Farmindustria e le Linee Guida per la Certificazione delle Attività Scientifiche: come applicarle in azienda** Medical Affairs / Regulatory
01 and 02 October 2024 *Online*	**State of the Art Section for Medical Devices – Unpacking the Tips and Tricks of a Complex Document** Medical Device
01 e 03 ottobre 2024 *Online*	**Cosmetici - Aspetti Tecnico-Regolatori e Panorama Normativo** Regulatory
01 and 02 October 2024 *Online*	**Process Validation** *Strategies to implement a continuous process verification during the product life cycle* *GMP (Good Manufacturing Practices)*
2 October 2024 *Online*	**Pharmacovigilance Documents - Focus on the Risk Management Plan (RMP)** Pharmacovigilance / Medical Writing
02 October 2024 *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*
08 e 10 ottobre 2024 *Online*	**Gestione della Proprietà Industriale - Il Brevetto per Tutelare le Invenzioni** Regulatory
09 ottobre 2024 *Online*	**Integratori Alimentari ed Advertising** *Caratteristiche di una Corretta Comunicazione Commerciale e Vincoli Normativi* *Regulatory*
09 and 10 October 2024 *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*

09 October and 16 October 2024 *Online*	**Pharmacovigilance Documents – Focus on the Periodic Safety Update Report (PSUR)** Pharmacovigilance / Medical Writing
10 and 11 October 2024 *Online*	**US FDA 101 - Fundamentals of Pre-market Submissions to CDRH** *Understanding US FDA pre-market submission pathways from an EU perspective – and why engaging the Agency through Pre-Submission Meetings is a valuable tool for Sponsors* *Medical Device / Regulatory*
15 and 16 October 2024 *Online*	**Medical Devices Periodic Safety Update Report (PSUR)** *What's behind the MDCG guidelines* *Medical Device*
15 and 17 October 2024 *Online*	**The Basics of Regulatory Affairs for Cosmetic Products in US and Canada** Regulatory
15 October 2024 *Online*	**Introduction to Aseptic Process Simulation (APS)** *Fundamentals to comprehend Aseptic Process Simulation* *GMP (Good Manufacturing Practices)*
16 e 23 ottobre 2024 *Online*	**Off-Label, Uso Compassionevole e Accesso Precoce (Early Access)** *Le normative e le loro applicazioni* *Medical Affairs / Regulatory / Clinical Research*
17 October 2024 *Online*	**How to Create Effective Visuals for Better Communicating your Science** Pharmacovigilance / Medical Affairs / Medical Writing / Soft Skill
21 e 24 ottobre 2024 *Online*	**Terapie Digitali (DTx): a che Punto Siamo?** *L'innovazione digitale per il futuro delle terapie* *Medical Affairs / Clinical Research*
22 October 2024 *Online*	**What You Need to Know about Medical Device Software and Never Dared to Ask** *More than just apps – An introduction to Medical Device Software* *Medical Device*
22 and 23 October 2024 *Online*	**Veterinary Pharmacovigilance System** Animal Health / Pharmacovigilance

22 e 29 ottobre 2024 *Online*	**Terapie Geniche e Terapie Cellulari** *Dalla Ricerca all'Impiego nel Real World* *Regulatory*
23 and 24 October 2024 *Online*	**Clinical Study Reports – a 360° Perspective** *Planning and Authoring CSRs in Accordance with Public Disclosure Requirements* *Medical Writing / Clinical Research*
23 October 2024 *Online*	**Pharmacovigilance Documents – Focus on Addendum to the Clinical Overview (AddCO) and Referrals** Pharmacovigilance / Medical Writing
24 ottobre 2024 *Online*	**La Compliance ai Requisiti GMP: Ruolo e Funzioni dell'Assicurazione Qualità** *Standard GMP e processi di qualità: teoria ed esempi pratici* *GMP (Good Manufacturing Practices)*
28 and 29 October 2024 *Online*	**Initiating the Development of Artificial Intelligence (AI) Medical Devices** *Developing AI medical devices considering the latest industry expectations* *Medical Device*
29 October 2024 *Online*	**Writing Science for Lay Audiences** *Ways to better get your message across* *Medical Device / Medical Affairs / Medical Writing / Clinical Research*
29 and 31 October 2024 *Online*	**Cleaning Validation** *GMP requirements and technical methods to ensure clean production systems and small materials* *GMP (Good Manufacturing Practices)*
30 October and 6 November 2024 *Online*	**Advanced Therapy Medicinal Product (ATMP): a Roadmap from Classification to Regulation and Manufacturing** GMP (Good Manufacturing Practices) / Regulatory
30 October 2024 *Online*	**Veterinary Marketing Authorisation Application (MAA) in the EU** *Clinical and Safety documentation* *Animal Health / Regulatory*

31 ottobre 2024 *Online*	**La Ricerca della Letteratura Scientifica: come Sfruttare le Risorse Gratuite in Rete** Medical Device / Pharmacovigilance / Medical Affairs / Medical Writing / Clinical Research
05 and 07 November 2024 *Online*	**Marketing Authorization Application in EU, US and UK** *Regulatory framework for a strategic plan until the entry in the market* *Regulatory*
05 and 07 November 2024 *Online*	**Beyond PubMed** *Additional approaches and sources for cost-effective literature monitoring* *Pharmacovigilance / Medical Affairs / Clinical Research*
05, 07, 12 e 14 novembre 2024 *Online*	**Safety Management e Farmacovigilanza** *Aspetti normativi, clinici e metodologici della farmacovigilanza, gestione della safety nello sviluppo clinico del farmaco* *Pharmacovigilance / Clinical Research*
06, 08 e 13 novembre 2024 *Online*	**Integratori Alimentari - Aspetti Tecnico-Regolatori e Panorama Normativo** Regulatory
06, 13 and 20 November 2024 *Online*	**All You Need to Know to Understand Statistic if You are not a Statistician** *Statistical principles from research planning to publication in scientific journals* *Medical Affairs / Medical Writing / Clinical Research*
11, 15 e 18 novembre 2024 *Online*	**Audit to Computer Systems** *Be ready!* *Clinical Research*
12 e 14 novembre 2024 *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 2024 *Online*	**Sterilization validation** *How to be sure of achieving sterility in pharmaceutical processes* *GMP (Good Manufacturing Practices)*

13 e 15 novembre 2024 *Online*	**Condurre uno Studio Clinico Adempiendo alla Normativa GDPR: un Ponte fra Good Clinical Practices (GCP) e General Data Protection Regulation (GDPR)** Clinical Research
19 and 21 November 2024 *.Online*	**Mastering Benefit Risk Assessment in Pharmacovigilance** Pharmacovigilance
19, 21, 26 and 28 November 2024 *Online*	**Computer System Validation (CSV) and Assurance (CSA)** *Insights into CSV fundamentals and different approaches for CSV and CSA* *GMP (Good Manufacturing Practices)*
19 and 21 November 2024 *Online*	**Pharmacovigilance Quality Management System (QMS)** Pharmacovigilance
20 and 25 November 2024 *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*
20 November 2024 *Online*	**Decoding Electronic Product Information (ePI)** *An in-depth exploration from paper to ePI* *Regulatory*
26 and 27 November 2024 *Online*	**Medical Reading** *The critical evaluation of scientific publications* *Medical Device / Medical Affairs / Clinical Research*
27 November 2024 *Online*	**Veterinary Marketing Authorisation Variation (MAV) in the EU** Animal Health / Regulatory
28 November 2024 *Online*	**The “Global” Qualified Person Responsible for Pharmacovigilance (QPPV) Workshop** *Evolution of the role, challenges and opportunities in a global environment* *Pharmacovigilance*
03 December 2024 *Online*	**Veterinary Quality Management** Animal Health / Pharmacovigilance

10 and 12 December 2024 *Online*	**Introduction to Veterinary Pharmacovigilance** Animal Health / Pharmacovigilance

Training Paths

Dal 12 marzo al 18 giugno 2024 *Online*	**Percorso Formativo GMP Project Management** ***Il ruolo del Project Manager nella gestione di progetti nei siti produttivi farmaceutici secondo la ISO 21502*** *GMP (Good Manufacturing Practices) / Soft Skill*
From 18 September to 23 October 2024 *Online*	**Pharmacovigilance Documents in the Life Cycle of a Medicinal Product** *From Papers and Patients to Health Authorities* *Pharmacovigilance / Medical Writing*
dal 27 settembre al 23 novembre 2024 *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*
Dal 04 ottobre al 13 dicembre 2024 *6 week end (online) e 1 project work (online)*	**Medical Affairs: Prepararsi alle Sfide Future di un Ruolo Complesso** *Corso di alta specializzazione per lo sviluppo delle competenze tecnico-scientifiche peculiari del Medical Advisor (MA) e del Medical Science Liaison (MSL)* *Medical Affairs*

Conferences

17 September 2024 *Frankfurt am Main*	**German Pharmacovigilance Day** *Pharmacovigilance: The Many Facets of one EU Framework*
07 and 08 October 2024 *Copenhagen*	**MedDev Day** *Medical Devices Regulation Update: Exploring the Thriving Regulatory Landscape*

23 Ottobre 2024 *Bergamo*	**Italian Pharmacovigilance Day** *La Farmacovigilanza al Servizio del Paziente: Nuove Sfide, Esempi di Successo e Strategie di Eccellenza*
13 November 2024 *Copenhagen*	**Nordic Pharmacovigilance Day** *Harmonizing Perspectives to Support Implementation of Common Objectives in Pharmacovigilance: Insights from the Nordic Region*