If you attend EPDP workshops and successfully complete the assignments before and after the workshop, you can gain EPDP credits. Credits can be accumulated towards an EMWA certificate, which shows your commitment to learning and professional development.

Thursday 9th November
17:00 19:00 Opening session and Networking Drinks

Friday 10th November
10:00 13:30 Writing in Plain Language for Peer Reviewed Publications
14:00 17:30 AI in Medical Writing
17:45 19:30 Freelance Business Forum

Monday 13th November
Morning
09:15 12:15 Introduction to Medical Writing

Afternoon
MSF10a 13:30 16:30 An Introduction to Vaccines
MCF33 13:30 16:30 Introduction to Enhanced Content in Publications

Tuesday 14th November
LWF13+14 09:15 12:45 Editing and Proofreading Essentials (Double Workshop)

Wednesday 15th November
Morning
LWF13+14 09:15 12:45 Editing and Proofreading Essentials (Double Workshop)

Afternoon
DDA19 13:30 17:00 Pharmacokinetic and Pharmacodynamic Modelling: an Overview for Medical Writers
LWA11a 13:30 16:30 Tense
Thursday 16th November

Morning
DDF33+34b 09:15 12:45 Clinical Study Reports - Mastering the Essential Skills (Double Workshop)

Afternoon
MCA5 13:10 17:00 Writing for the Public
DDF17b 13:30 17:00 Ethical Issues in Health Care

Friday 17th November

Morning
MDF2a 09:15 12:45 Going from Pharma to Medical Devices
DDF33+34d 09:15 12:45 Clinical Study Reports - Mastering the Essential Skills (Double Workshop in 2 parts)

Afternoon
MDA3 What Medical Writers should know about Medical Device Software

Saturday 18th November

Morning
DDF50 09:15 12:45 A Beginner’s Guide to Key Clinical Documents in the EU Drug Development Process
MCF32 09:15 12:45 How to handle advisory board meetings

Afternoon
LWA12 13:30 17:00 Master Class: Taxonomic Analysis of Medical Writing
MCF12a 13:30 16:30 Grant Writing

Monday 20th November

Morning
MCF22 09:15 12:15 Developing Effective Oral Presentations
DDA26 09:15 12:45 Post-submission Pharmacovigilance Writing: Interactions with Authorities and Impact on RMPs and PSURs

Afternoon
PTA11 13:30 17:00 Strategies for Improving Document Quality

Tuesday 21st November

Morning
MDF9 09:15 12:45 Writing the State for the Art of medical devices according to the EU Medical Devices Regulation
MCA4 09:15 12:45 Manuscript Writing: from Good to Excellent

Wednesday 22nd November

Morning
DDF38a 09:15 12:45 CORE Reference - Clarity and Openness in Reporting: E3-based
MCA28 09:15 12:45 Publication Planning

Afternoon
DDF36 13:30 17:00 An Introduction to Clinical Trial Disclosure – the Regulatory Requirements, Industry Commitments, and Protection of Data Privacy and Company Confidential Information

Thursday 23rd November

Afternoon
DDF41 13:30 17:00 Writing a Clinical Study Protocol

Friday 24th November

Morning
MCF17a 09:15 12:45 Using Writing Guidelines
MDA2b 09:15 12:45 How to Write a Clinical Evaluation Report