2025 Training Calendar

GMED NORTH AMERICA
LNE GROUP

VIRTUAL CLASSROOM TRAININGS	Jan.	Feb.	Mar.	April	May	June	July	Aug.	Sept.	Oct.	Nov.	Dec.
NEW! Biocompatibility for Medical Devices												16-18
The Clinical Evaluation Report (CER) Requirements Under the EU MDR 2017/745				1-2					24-25			
NEW! Cleanliness of Newly Manufactured Medical Devices					6-7							
Medical Device Electrical Safety							22-24					
European in Vitro Diagnostic Device Regulation (EU) 2017/746		● 11-12								● 28-29		
NEW! Technical Documentation per In Vitro Diagnostic Device Regulation (EU) 2017/746											● 18-20	
NEW! Medical Device Software Lifecycle per IEC 62304				22-24								
European Medical Device Regulation (EU) 2017/745	○ 28-30						● 15-16		○ 16-17			
Technical Documentation per Medical Device Regulation (EU) 2017/745		○ 25-27						● 12-14		○ 14-16		
Medical Device Single Audit Program (MDSAP) for Manufacturers									29 - 3			
Post Market Surveillance and Vigilance							29-30				4	
Risk Management Applied to Medical Devices		4-5				16-17						2-3

● Bundle offer available when color is the same.

GMED North America

gmedna-training@lne-gmed.com