2026 Training Calendar

GMED LNE GROUP NORTH AMERICA

VIRTUAL CLASSROOM TRAININGS	Jan.	Feb.	Mar.	April	May	June	July	Aug.	Sept.	Oct.	Nov.	Dec.
European Medical Device Regulation (EU) 2017/745	27-28					23-24			15-16			
European In Vitro Diagnostic Device Regulation (EU) 2017/746		17-18		8-9		2-3				27-28		
Risk management applied to Medical Devices		10-11				16-17						1-2
Systematic literature review for medical devices		25							23			
Post Market Surveillance and Vigilance			10				28				10	
The Clinical Evaluation Report (CER) requirements under the EU MDR 2017/745				1				4		21		
Clinical investigation of medical devices					20						18	
Technical Documentation per European Medical Device Regulation (EU) 2017/745			3-4				14-15			13-14		
Technical Documentation per European In Vitro Diagnostic Device Regulation (EU) 2017/746			10-11		26-27		14-15				24-25	
Medical Device Electrical Safety							21-22					
Medical device software lifecycle per IEC 62304				21-22								
Biocompatibility of medical devices				28-29				11-12				10-11
Cleanliness of newly manufactured medical devices					5-6							
Radiation sterilization of medical devices								25-26				
Medical Device Single Audit Program for manufacturers			30-31						29-30			
ISO 13485 requirements			3-4		12-13					6-7		

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