2027 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		2-3				15-16		31 - 1				
European In Vitro Diagnostic Device Regulation (EU) 2017/746		9-10		27-28		22-23				19-20		
Risk management applied to Medical Devices		23-24				29-30					9-10	
Systematic literature review for medical devices			2						14			
Post Market Surveillance and Vigilance			9				13			26		
The Clinical Evaluation Report (CER) requirements under the EU MDR 2017/745				20				10				7
Technical Documentation per European Medical Device Regulation (EU) 2017/745			16-17				20-21		28-29			
Technical Documentation per European In Vitro Diagnostic Device Regulation (EU) 2017/746			30-31		18-19		27-28				16-17	
Medical Device Electrical Safety								3-4				
Medical device software lifecycle per IEC 62304					4-5					12-13		
Biocompatibility of medical devices		10-11			11-12			17-18			30 -1	
Cleanliness of newly manufactured medical devices					25-26							
Radiation sterilization of medical devices								24-25				
Medical Device Single Audit Program for manufacturers				6-7					21-22			
ISO 13485 requirements				13-14		8-9					2-3	

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