


Basic information	
<p>2023/0005(COD)</p> <p>COD - Ordinary legislative procedure (ex-codecision procedure) Regulation</p>	Procedure completed
<p>Transitional provisions for certain medical devices and in vitro diagnostic medical devices</p> <p>Amending Regulation 2017/745 2012/0266(COD) Amending Regulation 2017/746 2012/0267(COD)</p> <p>Subject</p> <p>2.10.03 Standardisation, EC/EU standards and trade mark, certification, compliance 3.40.11 Precision engineering, optics, photography, medical 4.20.05 Health legislation and policy 4.60.08 Safety of products and services, product liability</p>	

Documentation gateway				
European Parliament				
Document type	Committee	Reference	Date	Summary
Text adopted by Parliament, 1st reading/single reading		T9-0052/2023	16/02/2023	Summary
Council of the EU				
Document type	Reference	Date	Summary	
Draft final act	00001/2023/LEX	15/03/2023		
European Commission				
Document type	Reference	Date	Summary	
Legislative proposal	COM(2023)0010 	06/01/2023	Summary	
Other institutions and bodies				
Institution/body	Document type	Reference	Date	Summary
ESC	Economic and Social Committee: opinion, report	CES0203/2023	24/01/2023	