


Basic information	
<p><b>2023/0005(COD)</b></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 <a href="#">2012/0266(COD)</a> Amending Regulation 2017/746 <a href="#">2012/0267(COD)</a></p> <p><b>Subject</b></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				
<b>European Parliament</b>				
Document type	Committee	Reference	Date	Summary
Text adopted by Parliament, 1st reading/single reading		<a href="#">T9-0052/2023</a>	16/02/2023	<a href="#">Summary</a>
<b>Council of the EU</b>				
Document type	Reference	Date	Summary	
Draft final act	<a href="#">00001/2023/LEX</a>	15/03/2023		
<b>European Commission</b>				
Document type	Reference	Date	Summary	
Legislative proposal	<a href="#">COM(2023)0010</a> 	06/01/2023	<a href="#">Summary</a>	
<b>Other institutions and bodies</b>				
Institution/body	Document type	Reference	Date	Summary
ESC	Economic and Social Committee: opinion, report	<a href="#">CES0203/2023</a>	24/01/2023	