



Basic information	
<p><b>2023/0005(COD)</b></p> <p>COD - Ordinary legislative procedure (ex-codecision procedure) Regulation</p>	<p>Procedure completed</p>
<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>	

Key events			
Date	Event	Reference	Summary
06/01/2023	Legislative proposal published	<a href="#">COM(2023)0010</a> 	<a href="#">Summary</a>
26/01/2023	Committee referral announced in Parliament, 1st reading		
16/02/2023	Decision by Parliament, 1st reading	<a href="#">T9-0052/2023</a>	<a href="#">Summary</a>
16/02/2023	Results of vote in Parliament		
07/03/2023	Act adopted by Council after Parliament's 1st reading		
15/03/2023	Final act signed		
20/03/2023	Final act published in Official Journal		