

Basic information	
<b>2023/0005(COD)</b> COD - Ordinary legislative procedure (ex-codecision procedure) Regulation	Procedure completed
Transitional provisions for certain medical devices and in vitro diagnostic medical devices  Amending Regulation 2017/745 <a href="#">2012/0266(COD)</a> Amending Regulation 2017/746 <a href="#">2012/0267(COD)</a>  <b>Subject</b>  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	

Technical information	
<b>Procedure reference</b>	2023/0005(COD)
<b>Procedure type</b>	COD - Ordinary legislative procedure (ex-codecision procedure)
<b>Procedure subtype</b>	Legislation
<b>Legislative instrument</b>	Regulation
	Amending Regulation 2017/745 <a href="#">2012/0266(COD)</a> Amending Regulation 2017/746 <a href="#">2012/0267(COD)</a>
<b>Legal basis</b>	Rules of Procedure EP 170 Treaty on the Functioning of the EU TFEU 168-p4 Treaty on the Functioning of the EU TFEU 114
<b>Mandatory consultation of other institutions</b>	<a href="#">European Economic and Social Committee</a> <a href="#">European Committee of the Regions</a>
<b>Stage reached in procedure</b>	Procedure completed
<b>Committee dossier</b>	ENVI/9/11070