

Basic information	
<p><b>2021/0323(COD)</b></p> <p>COD - Ordinary legislative procedure (ex-codecision procedure) Regulation</p>	Procedure completed
<p>Transitional provisions for certain in vitro diagnostic medical devices and deferred application of requirements for in-house devices</p> <p>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> <p><b>Legislative priorities</b></p> <p><a href="#">The EU's response to the Covid-19 pandemic</a></p>	

Technical information	
<b>Procedure reference</b>	2021/0323(COD)
<b>Procedure type</b>	COD - Ordinary legislative procedure (ex-codecision procedure)
<b>Procedure subtype</b>	Legislation
<b>Legislative instrument</b>	Regulation
<b>Amendments and repeals</b>	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 European Union TFEU 114 Treaty on the Functioning of the European Union TFEU 168-p4
<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/9/07442