


Basic information	
<p>2021/0323(COD)</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 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> <p>Legislative priorities</p> <p>The EU's response to the Covid-19 pandemic</p>	

Documentation gateway				
European Parliament				
Document type	Committee	Reference	Date	Summary
Text adopted by Parliament, 1st reading/single reading		T9-0498/2021	15/12/2021	Summary
Council of the EU				
Document type	Reference	Date	Summary	
Draft final act	00079/2021/LEX	25/01/2022		
European Commission				
Document type	Reference	Date	Summary	
Legislative proposal	COM(2021)0627 	14/10/2021	Summary	
National parliaments				
Document type	Parliament /Chamber	Reference	Date	Summary
Contribution	ES_PARLIAMENT	COM(2021)0627	22/11/2021	
Other institutions and bodies				
Institution/body	Document type	Reference	Date	Summary
EESC	Economic and Social Committee: opinion, report	CES5475/2021	08/12/2021	

