


Basic information	
<p>2024/0021(COD)</p> <p>COD - Ordinary legislative procedure (ex-codecision procedure) Regulation</p>	<p>Procedure completed</p>
<p>Gradual roll-out of Eudamed, information obligation in case of interruption of supply and transitional provisions for certain in vitro diagnostic medical devices</p> <p>Amending Regulation 2017/745 2012/0266(COD) 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>	

Documentation gateway				
European Parliament				
Document type	Committee	Reference	Date	Summary
Text adopted by Parliament, 1st reading/single reading		T9-0368/2024	25/04/2024	Summary
Council of the EU				
Document type	Reference	Date	Summary	
Draft final act	00054/2024/LEX	13/06/2024		
European Commission				
Document type	Reference	Date	Summary	
Legislative proposal	COM(2024)0043 	23/01/2024	Summary	
National parliaments				
Document type	Parliament /Chamber	Reference	Date	Summary
Contribution	ES_PARLIAMENT	COM(2024)0043	09/04/2024	
Other institutions and bodies				
Institution/body	Document type	Reference	Date	Summary
EESC	Economic and Social Committee: opinion, report	CES0746/2024	20/03/2024	

