


Basic information	
<p><b>2024/0021(COD)</b></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 <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
23/01/2024	Legislative proposal published	<a href="#">COM(2024)0043</a> 	<a href="#">Summary</a>
26/02/2024	Committee referral announced in Parliament, 1st reading		
25/04/2024	Decision by Parliament, 1st reading	<a href="#">T9-0368/2024</a>	<a href="#">Summary</a>
30/05/2024	Act adopted by Council after Parliament's 1st reading		
13/06/2024	Final act signed		
09/07/2024	Final act published in Official Journal		