

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>	

Additional information		
Source	Document	Date
EP Research Service	<a href="#">Briefing</a>	03/04/2024