





Basic information	
<p><b>2025/0404(COD)</b></p> <p>COD - Ordinary legislative procedure (ex-codecision procedure) Regulation</p>	Awaiting committee decision
<p>Simplifying and reducing the burden of the rules on medical devices and in vitro diagnostic medical devices, and support of the European Medicines Agency for the expert panels on medical devices and the list of Union harmonisation legislation</p> <p>Amending Regulation 2017/745 <a href="#">2012/0266(COD)</a> Amending Regulation 2017/746 <a href="#">2012/0267(COD)</a> Amending Regulation 2022/123 <a href="#">2020/0321(COD)</a> Amending Regulation 2024/1689 <a href="#">2021/0106(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>	

Documentation gateway				
<b>European Commission</b>				
Document type	Reference	Date	Summary	
Legislative proposal	<a href="#">COM(2025)1023</a> 	16/12/2025	<a href="#">Summary</a>	
Document attached to the procedure	<a href="#">SWD(2025)1050</a> 	17/12/2025		
Document attached to the procedure	<a href="#">SWD(2025)1051</a> 	17/12/2025		
Document attached to the procedure	<a href="#">SWD(2025)1052</a> 	17/12/2025		
<b>National parliaments</b>				
Document type	Parliament /Chamber	Reference	Date	Summary
Contribution	<a href="#">IT_CHAMBER</a>	<a href="#">COM(2025)1023</a>	27/04/2026	