

Basic information	
<p>2025/0404(COD)</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 2012/0266(COD) Amending Regulation 2017/746 2012/0267(COD) Amending Regulation 2022/123 2020/0321(COD) Amending Regulation 2024/1689 2021/0106(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>	

Key players			
European Parliament	Committee responsible	Rapporteur	Appointed
	SANT Public Health	SCHENK Oliver (EPP)	13/02/2026
		Shadow rapporteur WÖLKEN Tiemo (S&D) CASTILLO Laurent (Pfe) RAZZA Ruggero (ECR) GLÜCK Andreas (Renew) MARINO Ignazio Roberto (Greens/EFA) PALMISANO Valentina (The Left) ANDERSON Christine (ESN)	
	Committee for opinion	Rapporteur for opinion	Appointed
	IMCO Internal Market and Consumer Protection	GUZENINA Maria (S&D)	10/03/2026
	Committee for budgetary assessment	Rapporteur for budgetary assessment	Appointed

	BUDG Budgets		
Council of the European Union			
European Commission	Commission DG	Commissioner	
	Health and Food Safety	VÁRHELYI Olivér	
European Economic and Social Committee			
European Committee of the Regions			