Basic information	
2025/0404(COD) COD - Ordinary legislative procedure (ex-codecision procedure) Regulation	Awaiting committee decision
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 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) **Subject** 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	

Technical information	
Procedure reference	2025/0404(COD)
Procedure type	COD - Ordinary legislative procedure (ex-codecision procedure)
Procedure subtype	Legislation
Legislative instrument	Regulation
Amendments and repeals	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)
Legal basis	Rules of Procedure EP 58 Treaty on the Functioning of the European Union TFEU 168-p4 Treaty on the Functioning of the European Union TFEU 114
Mandatory consultation of other institutions	European Economic and Social Committee European Committee of the Regions
Stage reached in procedure	Awaiting committee decision
Committee dossier	SANT/10/04839