

Basic information	
<p>2012/0267(COD)</p> <p>COD - Ordinary legislative procedure (ex-codecision procedure) Regulation</p>	Procedure completed
<p>In vitro diagnostic medical devices</p> <p>Repealing Directive 98/79/EC 1995/0013(COD) Amended by 2021/0323(COD) Amended by 2023/0005(COD) Amended by 2024/0021(COD) See also 2012/0266(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>	

Technical information	
Procedure reference	2012/0267(COD)
Procedure type	COD - Ordinary legislative procedure (ex-codecision procedure)
Procedure subtype	Legislation
Legislative instrument	Regulation
	Repealing Directive 98/79/EC 1995/0013(COD) Amended by 2021/0323(COD) Amended by 2023/0005(COD) Amended by 2024/0021(COD) See also 2012/0266(COD)
Legal basis	Treaty on the Functioning of the EU TFEU 168-p4 Treaty on the Functioning of the EU TFEU 114-p1
Other legal basis	Rules of Procedure EP 165
Mandatory consultation of other institutions	European Economic and Social Committee European Committee of the Regions
Stage reached in procedure	Procedure completed
Committee dossier	ENVI/8/06746