

Procedure file

Basic information	
<p>COD - Ordinary legislative procedure (ex-codecision procedure) Regulation</p> <p>2021/0323(COD)</p>	Procedure completed
<p>Transitional provisions for certain in vitro diagnostic medical devices and deferred application of requirements for in-house devices</p> <p>Amending Regulation 2017/746 2012/0267(COD)</p> <p>Subject</p> <p>2.10.03 Standardisation, EC/EU standards and trade mark, certification, compliance</p> <p>3.40.11 Precision engineering, optics, photography, medical</p> <p>4.20.05 Health legislation and policy</p> <p>4.60.08 Safety of products and services, product liability</p>	

Technical information	
Procedure reference	2021/0323(COD)
Procedure type	COD - Ordinary legislative procedure (ex-codecision procedure)
Procedure subtype	Legislation
Legislative instrument	Regulation
	Amending Regulation 2017/746 2012/0267(COD)
Legal basis	Treaty on the Functioning of the EU TFEU 114; Treaty on the Functioning of the EU TFEU 168-p4; Rules of Procedure EP 163
Mandatory consultation of other institutions	European Economic and Social Committee European Committee of the Regions
Stage reached in procedure	Procedure completed
Committee dossier	ENVI/9/07442