

Procedure file

Basic information	
<p>COD - Ordinary legislative procedure (ex-codecision procedure) Regulation</p> <p>2012/0266(COD)</p>	Procedure completed
<p>Medical devices</p> <p>Repealing Directives 90/385/EEC and 93/42/EEC Amending Directive 2001/83/EC 1999/0134(COD) Amending Regulation (EC) No 178/2002 2000/0286(COD) Amending Regulation (EC) No 1223/2009 2008/0035(COD) See also 2012/0267(COD) Amended by 2020/0060(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	2012/0266(COD)
Procedure type	COD - Ordinary legislative procedure (ex-codecision procedure)
Procedure subtype	Legislation
Legislative instrument	Regulation
	<p>Repealing Directives 90/385/EEC and 93/42/EEC Amending Directive 2001/83/EC 1999/0134(COD) Amending Regulation (EC) No 178/2002 2000/0286(COD) Amending Regulation (EC) No 1223/2009 2008/0035(COD) See also 2012/0267(COD) Amended by 2020/0060(COD)</p>
Legal basis	Treaty on the Functioning of the EU TFEU 114-p1; Treaty on the Functioning of the EU TFEU 168-p4
Mandatory consultation of other institutions	European Economic and Social Committee European Committee of the Regions
Stage reached in procedure	Procedure completed
Committee dossier	ENVI/8/06745