

Basic information	
<p><b>2012/0266(COD)</b></p> <p>COD - Ordinary legislative procedure (ex-codecision procedure) Regulation</p>	Procedure completed
<p>Medical devices</p> <p>Amending Directive 2001/83/EC <a href="#">1999/0134(COD)</a> Amending Regulation (EC) No 178/2002 <a href="#">2000/0286(COD)</a> Amending Regulation (EC) No 1223/2009 <a href="#">2008/0035(COD)</a> See also <a href="#">2012/0267(COD)</a> Amended by <a href="#">2020/0060(COD)</a> Amended by <a href="#">2022/0417(COD)</a> Amended by <a href="#">2023/0005(COD)</a> Amended by <a href="#">2024/0021(COD)</a></p> <p><b>Subject</b></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>	

Delegated acts	
Reference	Subject
<a href="#">2023/2796(DEA)</a>	Examination of delegated act
<a href="#">2022/2986(DEA)</a>	Examination of delegated act