

Basic information	
<p><b>2020/0060(COD)</b></p> <p>COD - Ordinary legislative procedure (ex-codecision procedure) Regulation</p>	Procedure completed
<p>Medical devices</p> <p>Amending Regulation 2017/745 <a href="#">2012/0266(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> <p><b>Legislative priorities</b></p> <p><a href="#">The EU's response to the Covid-19 pandemic</a></p>	

Documentation gateway				
<b>European Parliament</b>				
Document type	Committee	Reference	Date	Summary
Text adopted by Parliament, 1st reading/single reading		<a href="#">T9-0053/2020</a>	17/04/2020	<a href="#">Summary</a>
<b>Council of the EU</b>				
Document type	Reference	Date	Summary	
Draft final act	<a href="#">00010/2020/LEX</a>	23/04/2020		
<b>European Commission</b>				
Document type	Reference	Date	Summary	
Legislative proposal	<a href="#">COM(2020)0144</a> 	03/04/2020	<a href="#">Summary</a>	
Commission response to text adopted in plenary	<a href="#">SP(2020)159</a>	13/05/2020		