

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>	

Key events			
14/10/2021	Legislative proposal published	COM(2021)0627	Summary
18/10/2021	Committee referral announced in Parliament, 1st reading		
27/10/2021	Decision by committee, without report		
13/12/2021	Debate in Parliament		
15/12/2021	Decision by Parliament, 1st reading	T9-0498/2021	Summary
20/12/2021	Act adopted by Council after Parliament's 1st reading		
25/01/2022	Final act signed		
28/01/2022	Final act published in Official Journal		