



Basic information	
<p>2018/0161(COD)</p> <p>COD - Ordinary legislative procedure (ex-codecision procedure) Regulation</p>	Procedure completed
<p>Supplementary protection certificate for medicinal products</p> <p>Amending Regulation (EC) No 469/2009 2008/0126(COD)</p> <p>Subject</p> <p>3.50.01.05 Research specific areas 3.50.16 Industrial property, European patent, Community patent, design and pattern 4.20.04 Pharmaceutical products and industry 6.20.05 Multilateral and plurilateral economic and trade agreements and relations</p>	

Key events			
Date	Event	Reference	Summary
28/05/2018	Legislative proposal published	COM(2018)0317 	Summary
02/07/2018	Committee referral announced in Parliament, 1st reading		
23/01/2019	Vote in committee, 1st reading		
23/01/2019	Committee decision to open interinstitutional negotiations with report adopted in committee		
29/01/2019	Committee report tabled for plenary, 1st reading	A8-0039/2019	Summary
30/01/2019	Committee decision to enter into interinstitutional negotiations announced in plenary (Rule 71)		
11/02/2019	Committee decision to enter into interinstitutional negotiations confirmed by plenary (Rule 71)		
26/02/2019	Approval in committee of the text agreed at 1st reading interinstitutional negotiations	GEDA/A/(2019)002691 PE637.374	
16/04/2019	Debate in Parliament	CRE link	
17/04/2019	Decision by Parliament, 1st reading	T8-0401/2019	Summary
17/04/2019	Results of vote in Parliament		
14/05/2019	Act adopted by Council after Parliament's 1st reading		
20/05/2019	Final act signed		
20/05/2019	End of procedure in Parliament		
11/06/2019	Final act published in Official Journal		