

Basic information	
<p>2013/0222(COD)</p> <p>COD - Ordinary legislative procedure (ex-codecision procedure) Regulation</p>	<p>Procedure completed</p>
<p>Medicinal products for human use: fees payable to the European Medicines Agency for the conduct of pharmacovigilance activities</p> <p>Repealed by 2022/0417(COD)</p> <p>Subject</p> <p>4.20.04 Pharmaceutical products and industry 8.40.08 Agencies and bodies of the EU</p>	

Documentation gateway				
European Parliament				
Document type	Committee	Reference	Date	Summary
Committee draft report		PE519.514	24/10/2013	
Amendments tabled in committee		PE523.004	11/11/2013	
Committee report tabled for plenary, 1st reading/single reading		A7-0476/2013	20/12/2013	Summary
Text adopted by Parliament, 1st reading/single reading		T7-0438/2014	16/04/2014	Summary
Council of the EU				
Document type		Reference	Date	Summary
Draft final act		00044/2014/LEX	15/05/2014	
European Commission				
Document type		Reference	Date	Summary
Legislative proposal		COM(2013)0472 	26/06/2013	Summary
Document attached to the procedure		SWD(2013)0234 	26/06/2013	
Document attached to the procedure		SWD(2013)0235 	26/06/2013	
Commission response to text adopted in plenary		SP(2014)471	09/07/2014	
Follow-up document		COM(2019)0439 	30/09/2019	Summary
National parliaments				
Document type	Parliament /Chamber	Reference	Date	Summary

Contribution	IT_SENATE	COM(2013)0472	20/08/2013	
Other institutions and bodies				
Institution/body	Document type	Reference	Date	Summary
EESC	Economic and Social Committee: opinion, report	CES5169/2013	16/10/2013	