




Basic information	
<b>2008/0260(COD)</b> COD - Ordinary legislative procedure (ex-codecision procedure) Directive	Procedure completed
Medicinal products for human use: pharmacovigilance of products  Amending Directive 2001/83/EC, Community code <a href="#">1999/0134(COD)</a> See also <a href="#">2008/0257(COD)</a>  <b>Subject</b>  4.20.04 Pharmaceutical products and industry 4.60.08 Safety of products and services, product liability	

Documentation gateway				
<b>European Parliament</b>				
Document type	Committee	Reference	Date	Summary
Committee draft report		<a href="#">PE430.927</a>	17/12/2009	
Committee opinion	<a href="#">IMCO</a>	<a href="#">PE431.039</a>	05/03/2010	
Amendments tabled in committee		<a href="#">PE438.412</a>	15/03/2010	
Committee opinion	<a href="#">ITRE</a>	<a href="#">PE430.773</a>	15/04/2010	
Committee report tabled for plenary, 1st reading/single reading		<a href="#">A7-0159/2010</a>	02/06/2010	
Text adopted by Parliament, 1st reading/single reading		<a href="#">T7-0332/2010</a>	22/09/2010	<a href="#">Summary</a>
<b>Council of the EU</b>				
Document type	Reference	Date	Summary	
Draft final act	<a href="#">00047/2010/LEX</a>	15/12/2010		
<b>European Commission</b>				
Document type	Reference	Date	Summary	
Legislative proposal	<a href="#">COM(2008)0665</a> 	10/12/2008	<a href="#">Summary</a>	
Document attached to the procedure	<a href="#">SEC(2008)2670</a> 	10/12/2008		
Document attached to the procedure	<a href="#">SEC(2008)2671</a> 	10/12/2008		
Commission response to text adopted in plenary	<a href="#">SP(2010)7193</a>	13/10/2010		
<b>Other institutions and bodies</b>				
Institution/body	Document type	Reference	Date	Summary

EDPS	Document attached to the procedure	JOC_2009/C/229/04 <a href="#">OJ C 229 23.09.2009, p. 0019</a>	22/04/2009	<a href="#">Summary</a>
EESC	Economic and Social Committee: opinion, report	<a href="#">CES1024/2009</a>	10/06/2009	