

# Procedure file

Basic information	
<p>COD - Ordinary legislative procedure (ex-codecision procedure) Regulation <a href="#">2008/0257(COD)</a></p>	Procedure completed
<p>Medicinal products for human use: pharmacovigilance of products</p> <p>Amending Regulation (EC) No 726/2004, Community procedures <a href="#">2001/0252(COD)</a></p> <p>Amending Regulation (EC) No 1394/2007 <a href="#">2005/0227(COD)</a></p> <p>See also <a href="#">2008/0260(COD)</a></p> <p>Subject</p> <p>4.20.04 Pharmaceutical products and industry</p> <p>4.20.05 Health legislation and policy</p> <p>4.60.08 Safety of products and services, product liability</p>	

Documentation gateway					
Legislative proposal		<a href="#">COM(2008)0664</a>	10/12/2008	EC	Summary
Document attached to the procedure		<a href="#">SEC(2008)2670</a>	10/12/2008	EC	
Document attached to the procedure		<a href="#">SEC(2008)2671</a>	10/12/2008	EC	
Document attached to the procedure		JOC_2009/C/229/04 <a href="#">OJ C 229 23.09.2009, p. 0019</a>	22/04/2009	EDPS	Summary
Economic and Social Committee: opinion, report		<a href="#">CES1023/2009</a>	10/06/2009	ESC	
Committee draft report		<a href="#">PE430.928</a>	17/12/2009	EP	
Committee opinion	IMCO	<a href="#">PE431.040</a>	24/02/2010	EP	
Amendments tabled in committee		<a href="#">PE438.413</a>	01/03/2010	EP	
Committee opinion	ITRE	<a href="#">PE430.771</a>	16/04/2010	EP	
Committee report tabled for plenary, 1st reading/single reading		<a href="#">A7-0153/2010</a>	10/05/2010	EP	
Text adopted by Parliament, 1st reading/single reading		<a href="#">T7-0331/2010</a>	22/09/2010	EP	Summary
Commission response to text adopted in plenary		<a href="#">SP(2010)7193</a>	13/10/2010	EC	
Draft final act		<a href="#">00046/2010/LEX</a>	15/12/2010	CSL	