

Basic information

2008/0257(COD)

COD - Ordinary legislative procedure (ex-codecision procedure)
Regulation

Medicinal products for human use: pharmacovigilance of products

Amending Regulation (EC) No 726/2004, Community procedures [2001/0252\(COD\)](#)

Amending Regulation (EC) No 1394/2007 [2005/0227\(COD\)](#)

See also [2008/0260\(COD\)](#)

Subject

4.20.04 Pharmaceutical products and industry

4.20.05 Health legislation and policy

4.60.08 Safety of products and services, product liability

Procedure completed

Documentation gateway




European Parliament

Document type	Committee	Reference	Date	Summary
Committee draft report		PE430.928	17/12/2009	
Committee opinion	IMCO	PE431.040	24/02/2010	
Amendments tabled in committee		PE438.413	01/03/2010	
Committee opinion	ITRE	PE430.771	16/04/2010	
Committee report tabled for plenary, 1st reading/single reading		A7-0153/2010	10/05/2010	
Text adopted by Parliament, 1st reading/single reading		T7-0331/2010	22/09/2010	Summary

Council of the EU

Document type	Reference	Date	Summary
Draft final act	00046/2010/LEX	15/12/2010	

European Commission

Document type	Reference	Date	Summary
Legislative proposal	COM(2008)0664 	10/12/2008	Summary
Document attached to the procedure	SEC(2008)2670 	10/12/2008	
Document attached to the procedure	SEC(2008)2671 	10/12/2008	
Commission response to text adopted in plenary	SP(2010)7193	13/10/2010	

Other institutions and bodies

Institution/body	Document type	Reference	Date	Summary
EDPS	Document attached to the procedure	JOC_2009/C/229/04 OJ C 229 23.09.2009, p. 0019	22/04/2009	Summary
EESC	Economic and Social Committee: opinion, report	CES1023/2009	10/06/2009	