

# Procedure file

Basic information	
COD - Ordinary legislative procedure (ex-codecision procedure) Regulation  Medicinal products for human use: implementation of good clinical practice in the conduct of clinical trials  Amended by <a href="#">2004/0217(COD)</a> Repealed by <a href="#">2012/0192(COD)</a> Amended by <a href="#">2021/0431(COD)</a>  Subject 4.20.02.06 Clinical practice and experiments 4.20.04 Pharmaceutical products and industry	Procedure completed

Documentation gateway					
Legislative proposal		<a href="#">COM(1997)0369</a> <a href="#">OJ C 306 08.10.1997, p. 0009</a>	03/09/1997	EC	Summary
Economic and Social Committee: opinion, report		<a href="#">CES0099/1998</a> <a href="#">OJ C 095 30.03.1998, p. 0001</a>	28/01/1998	ESC	Summary
Committee report tabled for plenary, 1st reading/single reading		<a href="#">A4-0407/1998</a> <a href="#">OJ C 379 07.12.1998, p. 0005</a>	29/10/1998	EP	
Text adopted by Parliament, 1st reading/single reading		T4-0648/1998 <a href="#">OJ C 379 07.12.1998, p. 0017-0034</a>	17/11/1998	EP	Summary
Modified legislative proposal		COM(1999)0193 <a href="#">OJ C 161 08.06.1999, p. 0005</a>	26/04/1999	EC	Summary
Council position		<a href="#">08878/1/2000</a> <a href="#">OJ C 300 20.10.2000, p. 0032</a>	20/07/2000	CSL	Summary
Commission communication on Council's position		SEC(2000)1293	26/07/2000	EC	Summary
Committee recommendation tabled for plenary, 2nd reading		<a href="#">A5-0349/2000</a> <a href="#">OJ C 232 17.08.2001, p. 0010</a>	21/11/2000	EP	
Text adopted by Parliament, 2nd reading		<a href="#">T5-0548/2000</a> <a href="#">OJ C 232 17.08.2001, p. 0035-0052</a>	12/12/2000	EP	Summary