

Basic information

1997/0197(COD)

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

Medicinal products for human use: implementation of good clinical practice in the conduct of clinical trials

Repealed by [2012/0192\(COD\)](#)

Amended by [2004/0217\(COD\)](#)

Amended by [2021/0431\(COD\)](#)

Subject

4.20.02.06 Clinical practice and experiments

4.20.04 Pharmaceutical products and industry

Procedure completed

Documentation gateway




European Parliament

Document type	Committee	Reference	Date	Summary
Committee report tabled for plenary, 1st reading/single reading		A4-0407/1998 OJ C 379 07.12.1998, p. 0005	29/10/1998	
Text adopted by Parliament, 1st reading/single reading		T4-0648/1998 OJ C 379 07.12.1998, p. 0017-0034	17/11/1998	Summary
Committee recommendation tabled for plenary, 2nd reading		A5-0349/2000 OJ C 232 17.08.2001, p. 0010	21/11/2000	
Text adopted by Parliament, 2nd reading		T5-0548/2000 OJ C 232 17.08.2001, p. 0035-0052	12/12/2000	Summary

Council of the EU

Document type	Reference	Date	Summary
Council position	08878/1/2000 OJ C 300 20.10.2000, p. 0032	20/07/2000	Summary

European Commission

Document type	Reference	Date	Summary
Legislative proposal	COM(1997)0369  OJ C 306 08.10.1997, p. 0009	03/09/1997	Summary
Modified legislative proposal	COM(1999)0193  OJ C 161 08.06.1999, p. 0005	26/04/1999	Summary
Commission communication on Council's position	SEC(2000)1293 	26/07/2000	Summary

Other institutions and bodies

Institution/body	Document type	Reference	Date	Summary
EESC	Economic and Social Committee: opinion, report	CES0099/1998 OJ C 095 30.03.1998, p. 0001	28/01/1998	Summary