Procedure file

Basic information

COD - Ordinary legislative procedure (ex-codecision procedure) Regulation	2008/0126(COD)
Supplementary protection certificate for medicinal products. Codification Amended by 2018/0161(COD) Subject 3.50.01.05 Research specific areas 3.50.16 Industrial property, European patent, Community patent, design and pattern 4.20.04 Pharmaceutical products and industry	Procedure completed

Key players

European Parliament	Committee responsible	Rapporteur	Appointed
	JURI Legal Affairs		25/06/2008
		ALDE WALLIS Diana	
Council of the European Union	Council configuration	Meeting	Date
	Justice and Home Affairs (JHA)	2936	06/04/2009
European Commission	Commission DG	Commissioner	
	Financial Stability, Financial Services and Capital Markets Union	MCCREEVY Charlie	

Key events

17/06/2008	Legislative proposal published	COM(2008)0369	Summary
19/06/2008	Committee referral announced in Parliament, 1st reading		
07/10/2008	Vote in committee, 1st reading		Summary
09/10/2008	Committee report tabled for plenary, 1st reading	A6-0385/2008	
21/10/2008	Results of vote in Parliament		
21/10/2008	Decision by Parliament, 1st reading	T6-0482/2008	Summary
06/04/2009	Act adopted by Council after Parliament's 1st reading		
06/05/2009	Final act signed		
06/05/2009	End of procedure in Parliament		
16/06/2009	Final act published in Official Journal		

Technical information

Procedure reference	2008/0126(COD)
Procedure type	COD - Ordinary legislative procedure (ex-codecision procedure)
Procedure subtype	Codification
Legislative instrument	Regulation
	Amended by 2018/0161(COD)
Legal basis	EC Treaty (after Amsterdam) EC 095
Stage reached in procedure	Procedure completed
Committee dossier	JURI/6/64375

Documentation gateway

Legislative proposal		COM(2008)0369	17/06/2008	EC	Summary
Economic and Social Committee: opinion, report		CES1510/2008	17/09/2008	ESC	
Committee report tabled for plenary, 1st reading/single reading		A6-0385/2008	09/10/2008	EP	
Text adopted by Parliament, 1st reading/single reading		T6-0482/2008	21/10/2008	EP	Summary
Draft final act		03602/2009/LEX	06/05/2009	CSL	

Additional information

National parliaments	IPEX
European Commission	EUR-Lex

Final act

Regulation 2009/469
OJ L 152 16.06.2009, p. 0001 Summary

Supplementary protection certificate for medicinal products. Codification

PURPOSE: to codify Council Regulation (EEC) No 1768/92 concerning the creation of a supplementary protection certificate for medicinal products.

PROPOSED ACT: Regulation of the European Parliament and of the Council.

CONTENT: the purpose of this proposal is to undertake a codification of Council Regulation (EEC) No 1768/92 concerning the creation of a supplementary protection certificate for medicinal products. The new Regulation will supersede the various acts incorporated into it. This proposal fully preserves the content of the acts being codified and hence does no more than bring them together with only such formal amendments as are required by the codification exercise itself.

The codification proposal was drawn up on the basis of a preliminary consolidation, in all official languages, of Regulation (EEC) No 1768/92 and the instruments amending it, carried out by the Office for Official Publications of the European Communities, by means of a data-processing system. Where the Articles have been given new numbers, the correlation between the old and the new numbers is shown in a table set out in Annex II to the codified Regulation.

Supplementary protection certificate for medicinal products. Codification

The Legal Affairs Committee adopted a report drafted by Diana WALLIS (ALDE, UK) and approved the proposal on the codification of Council Regulation (EEC) No 1768/92 concerning the creation of a supplementary protection certificate for medicinal products. The Commission?s proposal was approved as adapted to the recommendations of the Consultative Working Party of the Legal Services of the European

Parliament, the Council and the Commission.

Supplementary protection certificate for medicinal products. Codification

The European Parliament adopted, by 654 votes to 14 with 8 abstentions, under 1st reading of the codecision procedure, a legislative resolution approving the proposal for a regulation of the European Parliament and of the Council concerning the supplementary protection certificate for medicinal products (codified version). The report had been tabled for consideration in plenary by Diana WALLIS (ALDE, UK) on behalf of the Legal Affairs Committee. The Commission proposal was approved as adapted to the recommendations of the Consultative Working Party of the Legal Services of the European Parliament, the Council and the Commission.

Supplementary protection certificate for medicinal products. Codification

PURPOSE: to codify Council Regulation (EEC) No 1768/92 concerning the creation of a supplementary protection certificate for medicinal products.

LEGISLATIVE ACT: Regulation (EC) No 469/2009 of the European Parliament and of the Council concerning the supplementary protection certificate for medicinal products

CONTENT: the aim of this Regulation is to codify Council Regulation (EEC) No 1768/92of 18 June 1992 concerning the creation of a supplementary protection certificate for medicinal products. The new Regulation will supersede the various acts incorporated into it. It fully preserves the content of the acts being codified and hence does no more than bring them together with only such formal amendments as are required by the codification exercise itself.

Regulation (EEC) No 1768/92, as amended by the acts listed in Annex I, is repealed.

ENTRY INTO FORCE: 6 July 2009.