Fiche de procédure

Basic information

CNS - Consultation procedure Regulation	1999/0072(CNS)	Procedure completed
Foodstuffs of animal origin: maximum residue limits of veterinary medicinal products (amend. Regulation (EEC) No 2377/90) Subject 3.10.08 Animal health requirements, veterinary legislation and pharmacy 4.60.04.04 Food safety		

Key players

European Parliament	Committee responsible	Rapporteur	Appointed
	ENVI Environment, Public Health and Consumer Protection		
	Committee for opinion	Rapporteur for opinion	Appointed
	AGRI Agriculture and Rural Development		
Council of the European Union	Council configuration	Meeting	Date
	Agriculture and Fisheries	2404	21/01/2002
	Agriculture and Fisheries	2190	15/06/1999
	Agriculture and Fisheries	2178	17/05/1999

Key events

17/03/1999	Legislative proposal published	COM(1999)0130	Summary
12/04/1999	Committee referral announced in Parliament		
21/04/1999	Vote in committee		
04/05/1999	Decision by Parliament	T4-0353/1999	Summary
17/05/1999	Resolution/conclusions adopted by Council		
15/06/1999	Act adopted by Council after consultation of Parliament		
15/06/1999	End of procedure in Parliament		
23/06/1999	Final act published in Official Journal		

Technical information

Procedure reference	1999/0072(CNS)

Procedure type	CNS - Consultation procedure
Procedure subtype	Legislation
Legislative instrument	Regulation
Legal basis	EC Treaty (after Amsterdam) EC 037; Rules of Procedure EP 52-p1
Stage reached in procedure	Procedure completed
Committee dossier	ENVI/4/10878

Documentation gateway

Legislative proposal		COM(1999)0130 OJ C 131 12.05.1999, p. 0014	18/03/1999	EC	Summary
Economic and Social Committee: opinion, report		CES0454/1999 OJ C 169 16.06.1999, p. 0029	28/04/1999	ESC	
Text adopted by Parliament, 1st reading/single reading		T4-0353/1999 OJ C 279 01.10.1999, p. 0020-0054	04/05/1999	EP	Summary
Implementing legislative act		32002R0868 OJ L 037 25.05.2002, p. 0006	24/05/2002	EU	
Implementing legislative act		32002R0869 OJ L 037 25.05.2002, p. 0010	24/05/2002	EU	

Additional information

European Commission	EUR-Lex

Final act

Regulation 1999/1308
OJ L 156 23.06.1999, p. 0001 Summary

Foodstuffs of animal origin: maximum residue limits of veterinary medicinal products (amend. Regulation (EEC) No 2377/90)

PURPOSE: to amend regulation 2377/90 laying down a Community procedure for the establishment of maximum residue limits of veterinary medicinal products in foodstuffs of animal origin. CONTENT: the Commission proposes adapting regulation 2377/90/EEC by assigning to the European Agency for the assessment of medicines the task of dealing with requests for the establishment, modification and extension of maximum residue limits and by aligning the decision-making process with that of centralised procedure (role of the agenct secretariat, deadlines, appeals). It is also necessary to adapt the deadline for submission of projects and measures to the regulatory committee in order to allow the Community to fulfil its obligations resulting from the agreement on the application of sanitary and phytosanitary measures, concluded during Uruguay round negotiations and approved by the Community.?

Foodstuffs of animal origin: maximum residue limits of veterinary medicinal products (amend. Regulation (EEC) No 2377/90)

Under consultation procedure without report, the European Parliament approved the Commission proposal for a Council regulation amending regulation 2377/90/EEC laying down a Community procedure for the establishment of maximum residue limits of veterinary medicinal products in foodstuffs of animal origin.?

Foodstuffs of animal origin: maximum residue limits of veterinary medicinal products (amend. Regulation (EEC) No 2377/90)

PURPOSE: to amend Regulation 2377/90/EEC establishing a Community procedure for fixing the maximum limits of residues of veterinary medicines in foodstuffs of animal origin. COMMUNITY MEASURE: Council Regulation 1308/1999/EC. CONTENT: The purpose of the

modification introduced by the Council is to amend Regulation 2377/90/EEC laying down a Community procedure for the establishment of maximum residue limits of veterinary medicinal products in foodstuffs of animal origin by according the European Medicines Evaluation Agency Medicines the task of dealing with requests for the establishment, modification and extension of maximum residue limits and by aligning the decision-making process regarding the authorisation and surveillance of veterinary medicines with that laid down by Regulation 2309/93/EEC. ENTRY INTO FORCE: 26/06/1999.?