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COMMISSION REGULATION (EC) No 2430/1999

of 16 November 1999

linking the authorisation of certain additives belonging to the group of coccidiostats and other medicinal substances in feedingstuffs to persons responsible for putting them into circulation

(Text with EEA relevance)

(OJ L 296, 17.11.1999, p. 3)

Amended by:

		Official Journal		
	No	page	date	
►M1	Council Regulation (EC) No 1756/2002 of 23 September 2002	L 265	1	3.10.2002
►M2	Commission Regulation (EC) No 2037/2005 of 14 December 2005	L 328	21	15.12.2005

▼B**COMMISSION REGULATION (EC) No 2430/1999****of 16 November 1999**

linking the authorisation of certain additives belonging to the group of coccidiostats and other medicinal substances in feedingstuffs to persons responsible for putting them into circulation

(Text with EEA relevance)

THE COMMISSION OF THE EUROPEAN COMMUNITIES,

Having regard to the Treaty establishing the European Community,

Having regard to Council Directive 70/524/EEC of 23 November 1970 concerning additives in feedingstuffs⁽¹⁾, as last amended by Commission Regulation (EC) No 1636/1999⁽²⁾, and in particular Article 9h(3)(b) and Article 9i(3)(b) thereof,

Whereas:

- (1) because of the risk for human and animal health posed by the circulation in the Community of poor copies of zootechnical additives, Directive 70/524/EEC, as amended by Council Directive 96/51/EC⁽³⁾, provides for the linking of the authorisation of certain classes of additives to the person responsible for putting them into circulation;
- (2) in particular Article 9h of Directive 70/524/EEC provides for the replacement of the provisional authorisations of additives included in Annex I after 31 December 1987 and belonging to the group of coccidiostats and other medicinal substances and transferred to Chapter II of Annex B by authorisations linked to the person responsible for putting them into circulation for a period of 10 years;
- (3) in particular Article 9i of Directive 70/524/EEC provides for the replacement of the provisional authorisations of additives included in Annex II before 1 April 1998 and belonging to the group of coccidiostats and other medicinal substances and transferred to Chapter III of Annex B by provisional authorisations linked to the person responsible for putting them into circulation;
- (4) the additives listed in the Annexes to this Regulation were the subject of new applications for authorisation by the person responsible for the dossier on the basis of which the former authorisations were given or by their successors. The applications relating to those additives were accompanied by the required monographs and identification notes;
- (5) the linking of the authorisation to a person responsible for putting the additive into circulation is based on a purely administrative procedure and did not entail a fresh assessment of the additives. Although the authorisations are given for a specified period they may be withdrawn at any time in accordance with Article 9m and Article 11 of Directive 70/524/EEC. In particular, authorisations of additives may be withdrawn as a result of the re-evaluation carried out under Article 9g of Directive 70/524/EEC;
- (6) the measures provided for in this Regulation are in accordance with the opinion of the Standing Committee for Feedingstuffs,

⁽¹⁾ OJ L 270, 14.12.1970, p. 1.

⁽²⁾ OJ L 194, 27.7.1999, p. 17.

⁽³⁾ OJ L 235, 17.9.1996, p. 39.

▼B

HAS ADOPTED THIS REGULATION:

Article 1

The provisional authorisations of the additives listed in Annex I to this Regulation are replaced by authorisations granted to the person responsible for putting the additive in circulation, inserted in the second column of Annex I.

Article 2

The provisional authorisations of the additives listed in Annex II to this Regulation are replaced by provisional authorisations granted to the person responsible for putting the additive into circulation, inserted in the second column of Annex II.

Article 3

This Regulation shall enter into force on the seventh day of its publication in the *Official Journal of the European Communities*.

This Regulation shall be binding in its entirety and directly applicable in all Member States.

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ANNEX I

Registration number of additive	Name and registration number of person responsible for putting additive into circulation	Additive (trade name)	Composition, chemical formula, description	Species or category of animal	Maximum age	Minimum content mg of active substance/kg of complete feedingstuff	Maximum content mg of active substance/kg of complete feedingstuff	Other provisions	Period of authorisation	►M2 Maximum Residue Limits (MRLs) ▼
E 758	Roche Vitamins Europe Ltd	Robenidine hydrochloride 66 g/kg (Cycostat 66 G)	Additive composition: Robenidine lydrochloride: 66 g/kg Lignosulfonate: 40 g/kg Calcium sulfate dihydrate: 894 g/kg Active substance: Robenidine lydrochloride, $C_{15}H_{13}Cl_2N_5HCl$, 1,3-bis[(p-chlorobenzylidene)amino]guanidine hydrochloride, CAS number: 25875-50-7 Related impurities: N,N,N',N'-Tetris[(p-Cl-benzylidene)amino]guanidine: ≤ 1 % Bis-[4-Cl-benzylidene]hydrazine: ≤ 1 %	Rabbits for breeding purposes	—	50	66	Use prohibited at least five days before slaughter	30.9.2009	Regulation (EEC) No 2377/90
▼M2 Coccidiostats and histomonostats										
E 763	Alpharma (Belgium) BVBA	Lasalocid A sodium 15 g/100 g (Avatec 15 % cc)	Additive composition: Lasalocid A sodium: 15 g/100 g Corn cob meal: 80,95 g/100 g Lecitin: 2 g/100 g Soya oil: 2 g/100 g Ferric oxide: 0,05 g/100 g Active substance:	Turkeys	12 weeks	90	125	Use prohibited at least five days before slaughter. Indicate in the instructions for use: 'Dangerous for equine species' 'This feedingstuff contains an ionophore: simultaneous use with certain medicinal'	30.9.2009	

▼M2

Registration number of additive	Name and registration number of person responsible for putting additive into circulation	Additive (trade name)	Composition, chemical formula, description	Species or category of animal	Maximum age	Minimum content mg of active substance/kg of complete feedingstuff	Maximum content mg of active substance/kg of complete feedingstuff	Other provisions	Period of authorisation	►M2 Maximum Residue Limits (MRLs) ▼
			Lasalocid A sodium, $C_{34}H_{51}O_8Na$, CAS number: 25999-20-6, sodium salt of 6-[(3R, 4S, 5S, 7R)-7-[(2S, 3S, 5S)- 5-ethyl-5-[(2R, 5R, 6S)-5-ethyl-5- hydroxy-6-methyltetrahydro-2H- pyran2-yl]-tetrahydro-3-methyl-2- furyl]-4-hydroxy-3,5-dimethyl-6- oxononyl]-2,3-cresotic acid, produced by <i>Streptomyces lasaliensis</i> subsp. <i>lasaliensis</i> (ATCC 31180) Related impurities: Lasalocid sodium B-E; ≤ 10 %					substances can be contraindicated'.		
			Lasalocid A sodium 15 g/ 100 g (Avatec 150 G)	Turkeys	12 weeks	90	125	Use prohibited at least five days before slaughter. Indicate in the instructions for use: 'Dangerous for equine species' 'This feedingstuff contains an ionophore: simultaneous use with certain medicinal substances can be contraindicated'.	30.9.2009	Regulation (EEC) No 2377/90

▼M2	Registration number of additive	Name and registration number of person responsible for putting additive into circulation	Additive (trade name)	Composition, chemical formula, description	Species or category of animal	Maximum age	Minimum content mg of active substance/kg of complete feedingstuff	Other provisions	Period of authorisation	►M2 Maximum Residue Limits (MRLs) ▼
▼B	E 764	Hoechst Roussel Vet GmbH	Halofuginone hydrobromide 6 g/kg (Stenorol)	Additive composition: Halofuginone hydrobromide: 6 g/kg Gelatine: 13,2 g/kg Starch: 19,2 g/kg Sugar: 21,6 g/kg Calcium carbonate: 940 g/kg Active substance: Halofuginone hydrobromide, $C_{16}H_{17}BrClN_3O_3$, HBr DL-trans-7-bromo-6-chloro-3-(3-(3-hydroxy-2-piperidy)acetyl)- quinazolin-4(3H)-one hydrobromide, CAS number: 64924-67-0 Related impurities: Cis-isomer of halofuginone: < 1,5 %	Chickens for laying	16 weeks	2	3	—	30.9.2009
▼M1	—	—	—	—	—	—	—	—	—	—
▼B	E 770	Roche Vitamins Europe Ltd	Maduramicin ammonium alpha 1 g/100 g (Cygro 1 %)	Additive composition: Maduramicin ammonium alpha: 1 g/100 g Benzyl alcohol: 5 g/100 g	Chickens for fattening	—	5	5	Use prohibited at least five days before slaughter	30.9.2009

Registration number of additive	Name and registration number of person responsible for putting additive into circulation	Additive (trade name)	Composition, chemical formula, description	Species or category of animal	Maximum age	Minimum content mg of active substance/kg of complete feedingstuff	Maximum content	Other provisions	Period of authorisation	►M2 Maximum Residue Limits (MRLs) ▼
			Corn cob grits qs 100 g Active substance: Maduramicin ammonium alpha, C ₄₇ H ₈₃ O ₁₇ N, CAS number: 84878-61-5, ammonium salt of a polyether monocarboxylic acid produced by <i>Actinomadura yumaensis</i> (ATCC 31585) (NRRL 12515) Related impurities: Maduramicin ammonium beta: < 10 %					Indicate in the instructions for use: ‘Dangerous for equines?’ ‘This feedingstuff contains an ionophore: simultaneous use with certain substances (e.g. tiamulin) can be contraindicate’		
E 771	Janssen Animal Health B.V.B.A	Diclazuril 0,5 g/ 100 g (Clinacox 0,5 % Premix)	Diclazuril 0,5 g/ 100 g (Clinacox 0,2 % Premix)	Chickens for fattening	—	1	1	Use prohibited as least five days before slaughter	30.9.2009	

Registration number of additive	Name and registration number of person responsible for putting additive into circulation	Additive (trade name)	Composition, chemical formula, description CAS number: 101831-37-2	Species or category of animal	Maximum age	Minimum content mg of active substance/kg of complete feedingstuff	Other provisions	►M2 Maximum Residue Limits (MRLs) ▼	
								Period of authorisation	
E 772	Eli Lilly and Company Ltd	Narasin 80 g/kg — Nicarbazin 80 g/kg (Maxiban G160)	(±)-4-chlorophenyl[2,6-dichloro-4-(2,3,4,5-tetrahydro-3,5-dioxo-1,2,4-triazin-2-yl)phenyl]acetonitrile, Degradation compound (R064318): ≤ 0,2 % Related impurities: Other related impurities (R066891, R06896, R068610, R070156, R068584, R070016): ≤ 0,5 % individually Total impurities: ≤ 1,5 %	Chickens for fattening	—	80	100	Use prohibited at least five days before slaughter	30.9.2009
				Vermiculite: 0-20 g/kg Microtracer F-Red: 11 g/kg Corn cob grits or rice hulls qs 1 kg				Indicate in the instructions for use: ‘Dangerous for equines’ ‘This feedingstuff contains an ionophore: simultaneous use with certain medicinal substances (e.g.	

<u>B</u>	Name and registration number of person responsible for putting additive into circulation	Additive (trade name)	Composition, chemical formula, description	Species or category of animal	Maximum age	Minimum content	Maximum content	Other provisions	Period of authorisation	<u>M2</u> Maximum Residue Limits (MRLs) ▶
						mg of active substance/kg of complete feedingstuff	mg of active substance/kg of complete feedingstuff			
			(b) Nicarbazin, C ₁₉ H ₁₈ N ₆ O ₆ CAS number: 330-95-0 Equimolecular complex of 1,3-bis (4-nitrophenyl) urea and 4,6-dimethylpyrimidin-2-ol, in granular form Related impurities: p-nitroaniline: ≤ 1 %					tiamulin) can be contraindicated,		

M1

ANNEX II

Registration number of additive	Name and registration number of person responsible for putting additive into circulation	Additive (trade name)	Composition, chemical formula, description additive into circulation	Species or category of animal	Maximum age	Minimum content mg of active substance/kg of complete feedingstuff	Maximum content mg of active substance/kg of complete feedingstuff	Other provisions	Period of authorisation
26	Hoechst Roussel Vet GmbH	Salinomycin sodium 120 g/kg (Sacox 120)	Additive composition: Salinomycin sodium ≥ 120 g/kg Silicium dioxide 10-100 g/kg Calcium carbonate: 350-700 g/kg Active substance: Salinomycin sodium, C ₂₂ H ₆₉ O ₁₁ Na, CAS number: 53003-10-4, sodium salt of a polyether monocarboxylic acid produced by fermentation of <i>Streptomyces albus</i> (DSM 12217) Related impurities: < 42 mg elaiophylin/kg salinomycin sodium < 40 g 17-epi-20-desoxy-salinomycin/kg salinomycin sodium	Rabbits for fattening	—	20	25	Use prohibited at least five days before slaughter Indicate in the instructions for use: ‘Dangerous for equines’ ‘This feedingstuff contains an ionophore: simultaneous use with certain medicinal substances (e.g. tiamulin) can be contraindicated’	30.9.2000 (¹)
27	Janssen Animal Health B.V.B.A.	Diclazuril 0,5 g/100 g (Climacox 0,5 % Premix)	Additive composition: Diclazuril: 0,5 g/100 g Soybean meal: 99,25 g/100 g Diclazuril 0,2 g/100 g (Climacox 0,2 % Premix) Polyvidone K 30: 0,2 g/100 g Sodium hydroxyde: 0,0538 g/100 g Diclazuril: 0,2 g/100 g Soybean meal: 39,7 g/100 g	Turkeys Chickens reared for laying	12 weeks 16 weeks	1 1	1 1	Use prohibited at least five days before slaughter —	30.9.2000 (¹) 30.9.2000 (²)

Registration number of additive	Name and registration number of person responsible for putting additive into circulation	Additive (trade name)	Composition, chemical formula, description	Species or category of animal	Maximum age	Minimum content	Maximum content	Other provisions	Period of authorisation
									mg of active substance/kg of complete feedingstuff
			Polyvidone K 30: 0,08 g/100 g Sodium hydroxide: 0,0215 g/100 g Wheat middlings: 60 g/100 g						
28	Roche Vitamins Europe Ltd	Maduramicin ammonium alpha 1 g/100 g (Cygro 1 %)	Additive composition: Maduramicin ammonium alpha: 1 g/100 g Benzyl alcohol: 5 g/100 g Corn cob grits qs 100 g	Turkeys	16 weeks	5	5	Use prohibited at least five days before slaughter Indicate in the instructions for use: 'Dangerous for equines' 'This feedingstuff contains an ionophore: simultaneous use with certain medicinal substances (e.g. tiamulin) can be contraindicated'	30.9.2000 (2)

Registration number of additive	Name and registration number of person responsible for putting additive into circulation	Additive (trade name)	Composition, chemical formula, description	Species or category of animal	Minimum content	Maximum content	Other provisions	Period of authorisation
					Maximum age	mg of active substance/kg of complete feedingstuff		
			ammonium salt of a polyether monocarboxylic acid produced by <i>Actinomadura yumaensis</i> (ATCC 31585) (NRRL 12515) Related impurities: Maduramicin ammonium beta: < 10 %					

(¹) First authorisation: Commission Directive 96/7/EC (OJ L 51, 1.3.1996, p. 45).

(²) First authorisation: Commission Directive 96/66/EC (OJ L 272, 25.10.1996, p. 32).

(³) First authorisation: Commission Directive 97/72/EC (OJ L 351, 23.12.1997, p. 55).