This text is meant purely as a documentation tool and has no legal effect. The Union's institutions do not assume any liability for its contents. The authentic versions of the relevant acts, including their preambles, are those published in the Official Journal of the European Union and available in EUR-Lex. Those official texts are directly accessible through the links embedded in this document

►**B**

COMMISSION REGULATION (EC) No 971/2008

of 3 October 2008

concerning a new use of a coccidiostat as additive in feedingstuffs

(Text with EEA relevance)

(OJ L 265, 4.10.2008, p. 3)

Amended by:

		Official Journal		
		No	page	date
►**M1**	Commission Implementing Regulation (EU) No 160/2013 of 21 February 2013	L 49	50	22.2.2013
►**M2**	Commission Implementing Regulation (EU) 2019/138 of 29 January 2019	L 26	1	30.1.2019

Corrected by:

►**C1**	Corrigendum, OJ L 267, 8.10.2008, p. 32 (971/2008)

▼B

COMMISSION REGULATION (EC) No 971/2008

of 3 October 2008

concerning a new use of a coccidiostat as additive in feedingstuffs

(Text with EEA relevance)

Article 1

The preparation belonging to the group ‘Coccidiostats and other medicinal substances’, as specified in the Annex, is authorised for use for ten years as additive in animal nutrition under the conditions laid down in that Annex.

Article 2

This Regulation shall enter into force on the 20th day following that of its publication in the *Official Journal of the European Union*.

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

▼C1

ANNEX

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	End of period of authorisation	Maximum Residue Limits (MRLs) in the relevant foodstuffs of animal origin
						mg of active substance/kg of complete feedingstuff				
Coccidiostats and other medicinal substances										
E 771	►**M2** Elanco GmbH ◄	Diclazuril 0,5 g/100 g (Clinacox 0,5 % Premix)	*Additive composition:* Diclazuril: 0,5 g/100 g Soybean meal: 99,25 g/100 g Polyvidone K 30: 0,2 g/100 g Sodium hydroxide: 0,0538 g/100 g *Active substance:* Diclazuril $C_{17}H_9Cl_3N_4O_2$, (±)-4-chlorophenyl[2,6-dichloro-4-(2,3,4,5-tetrahydro-3,5-dioxo-1,2,4-triazin-2-yl) phenyl]acetonitrile, CAS number: 101831-37-2 *Related impurities:* Degradation compound (R064318): < 0,2 % Other related impurities (R066891, R066896, R068610, R070156, R068584, R070016): < 0,5 % individually Total impurities: < 1,5 %	Rabbits	—	1	1	Use prohibited at least one day before slaughter.	24 October 2018	2 500 µg diclazuril/kg of wet liver 1 000 µg diclazuril/kg of wet kidney 150 µg diclazuril/kg of wet muscle 300 µg diclazuril/kg of wet fat