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COMMISSION IMPLEMENTING REGULATION (EU) 2020/1175

of 7 August 2020

concerning the authorisation of L-cysteine hydrochloride monohydrate produced by fermentation with Escherichia coli KCCM 80180 and Escherichia coli KCCM 80181 as a feed additive for all animal species

(Text with EEA relevance)

THE EUROPEAN COMMISSION,

Having regard to the Treaty on the Functioning of the European Union,

Having regard to Regulation (EC) No 1831/2003 of the European Parliament and of the Council of 22 September 2003 on additives for use in animal nutrition (¹), and in particular Article 9(2) thereof,

Whereas:

- (1) Regulation (EC) No 1831/2003 provides for the authorisation of additives for use in animal nutrition and for the grounds and procedures for granting such authorisation.
- (2) In accordance with Article 7 of Regulation (EC) No 1831/2003, an application was submitted for the authorisation of L-cysteine hydrochloride monohydrate produced by fermentation with *Escherichia coli* KCCM 80180 and *Escherichia coli* KCCM 80181. This application was accompanied by the particulars and documents required under Article 7(3) of Regulation (EC) No 1831/2003.
- (3) This application concerns the authorisation of L-cysteine hydrochloride monohydrate produced by fermentation with *Escherichia coli* KCCM 80180 and *Escherichia coli* KCCM 80181 as a feed additive for all animal species. The applicant requested this additive to be classified in the additive category 'sensory additives'.
- (4) The European Food Safety Authority ('the Authority') concluded in its opinion of 10 January 2020 (²) that, under the proposed conditions of use in feed L-cysteine hydrochloride monohydrate produced by fermentation with *Escherichia coli* KCCM 80180 and *Escherichia coli* KCCM 80181 does not have adverse effects on animal health, consumer safety or the environment. In its conclusions it also considered that the applicant proposes to label the additive with the hazard statement H335 (may cause respiratory irritation) under the Regulation (EC) No 1272/2008 (³). Therefore, the Commission considers that appropriate protective measures should be taken to prevent adverse effects on human health, in particular, as regards the users of the additive. The Authority further concluded that since the substance is used in food and its function in feed is the same as that in food, no further demonstration of efficacy in feed is necessary.
- (5) Restrictions and conditions should be provided for to allow for a better control. For L-cysteine hydrochloride monohydrate, recommended contents should be indicated on the label of the additive. Where such contents are exceeded, certain information should be indicated on the label of premixtures.
- (6) The Authority does not consider that there is a need for specific requirements of post-market monitoring. It also verified the report on the method of analysis of the feed additives in feed submitted by the Reference Laboratory set up by Regulation (EC) No 1831/2003.
- (7) The assessment of L-cysteine hydrochloride monohydrate produced by fermentation with *Escherichia coli* KCCM 80180 and *Escherichia coli* KCCM 80181 shows that the conditions for authorisation, as provided for in Article 5 of Regulation (EC) No 1831/2003, are satisfied. Accordingly, the use of this substance should be authorised as specified in the Annex to this Regulation.
- (8) The fact that L-cysteine hydrochloride monohydrate produced by fermentation with *Escherichia coli* KCCM 80180 and *Escherichia coli* KCCM 80181 is not authorised for use as a flavouring in water for drinking, does not preclude its use in compound feed, which is administered via water.

⁽¹⁾ OJ L 268, 18.10.2003, p. 29..

^{(&}lt;sup>2</sup>) EFSA Journal 2020;18(2):6003.

^{(&}lt;sup>3</sup>) Regulation (EC) No 1272/2008 of the European Parliament and of the Council of 16 December 2008 on classification, labelling and packaging of substances and mixtures, amending and repealing Directives 67/548/EEC and 1999/45/EC, and amending Regulation (EC) No 1907/2006 (OJ L 353, 31.12.2008, p. 1).

(9) The measures provided for in this Regulation are in accordance with the opinion of the Standing Committee on Plants, Animals, Food and Feed,

HAS ADOPTED THIS REGULATION:

Article 1

The substances specified in the Annex, belonging to the additive category 'sensory additives' and to the functional group 'flavouring compounds', are authorised as feed additives in animal nutrition subject to the conditions laid down in that Annex.

Article 2

This Regulation shall enter into force on the twentieth 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.

Done at Brussels, 7 August 2020.

For the Commission The President Ursula VON DER LEYEN

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Official Journal of the European Union

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Identification	Name of the		Composition showing formula	Species or	Manimum	Minimum content	Maximum content		End of period
number of the additive	holder of authorisation	Additive	Composition, chemical formula, description, analytical method	category of animal		mg of active substance/kg of complete feedingstuff with a moisture content of 12 %		Other provisions	of authorisation
Category: Se	ensory additi	ves. Functional gro	oup: Flavouring compounds						

hydrochloride minimum by such procedures monohydrate in premixtures:
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- ion-exchange chromatography coupled with post-column deri- vatisation and photometric de- tection (IEC-VIS), Commission Regulation (EC) No 152/2009 (Annex III, F)	and measures, the additive and premixtures shall be used with personal protective equipment, including breath- ing protection.
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