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Directorate F – Health, Consumers and Reference Materials (Geel/Ispra)
European Union Reference Laboratory for Feed Additives

JRC F.5/CvH/SB/AS/Ares

**Evaluation Report on the Analytical Methods submitted
in connection with the Application for Authorisation of a
Feed Additive according to Regulation (EC) No 1831/2003**

L-methionine
produced by fermentation with
Corynebacterium glutamicum KCCM80184 and
Escherichia coli K12 KCCM80096
(*FAD-2018-0085; CRL/180051*)

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in connection with the Application for Authorisation of a
Feed Additive according to Regulation (EC) No 1831/2003**

Dossier related to: **FAD-2018-0085 - CRL/180051**

Name of Product: ***L-methionine produced by fermentation
with *Corynebacterium glutamicum*
KCCM80184 and *Escherichia coli* K12
KCCM80096***

Active Agent: **L-methionine**

Rapporteur Laboratory: **European Union Reference Laboratory for
Feed Additives (EURL-FA)
JRC Geel, Belgium**

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Date: **07/05/2019**

EXECUTIVE SUMMARY

In the current application authorisation is sought under Article 4(1) for *L-methionine produced by fermentation with Corynebacterium glutamicum KCCM80184 and Escherichia coli K12 KCCM80096*, under the category/functional group 3(c) 'nutritional additives/'amino acids, their salts and analogues', according to Annex I of Regulation (EC) No 1831/2003. Authorisation is sought for all animal species.

According to the Applicant, *L-methionine* has a minimum purity (mass fraction) of 98.5 %. The *feed additive* is intended to be added directly into *feedingstuffs* (or through *premixtures*) and *water* for drinking. However, the Applicant did not propose any minimum or maximum content of *L-methionine* in *feedingstuffs*.

For the quantification of *L-methionine* in the *feed additive* the Applicant submitted the method based on titration described in the corresponding European Pharmacopoeia monograph. For the quantification of *methionine* in the *feed additive* the EURL identified instead the ring-trial validated method EN ISO 17180:2013 based on ion-exchange chromatography coupled to visible or fluorescence detection (IEC-VIS/FLD). This standard method does not distinguish between the salts of amino acids and it cannot differentiate between enantiomers. It applies for products containing more than 10 % of amino acid. The following performance characteristics were reported: a relative standard deviation for repeatability (RSD_r) ranging from 0.5 to 1.6 % and a relative standard deviation for reproducibility (RSD_R) ranging from 1.5 to 2.6 %. In addition, the EURL identified the "L-methionine monograph" of the Food Chemical Codex (FCC) for the identification of *L-methionine* in the *feed additive*.

For the quantification of *L-methionine* in *premixtures* and *feedingstuffs* the Applicant submitted the ring-trial validated Community method (Commission Regulation (EC) No 152/2009) based on IEC coupled to photometric detection (IEC-VIS). This method, designed only for the analysis of amino acids in *premixtures* and *feedingstuffs*, does not distinguish between the salts and the amino acid enantiomers. The Community method was ring-trial validated using four different matrices. This method was further ring-trial validated by twenty-three laboratories, resulting in the EN ISO 13903:2005 method. The following performance characteristics were reported for the quantification of total *methionine*: RSD_r ranging from 1.1 to 5.6 % and RSD_R ranging from 6.9 to 13 %.

The Applicant did not provide experimental data to determine *L-methionine* in *water*. Nevertheless, for the quantification of *methionine* in *water*, as concluded in the previous EURL reports and further specified in the corresponding legislation, the EURL recommended the Community method for official control.

In the frame of this authorisation the EURL recommends for official control (i) the "L-methionine monograph" of the Food Chemical Codex (FCC) based on infrared absorption for the identification of *L-methionine* in the *feed additive*; (ii) the ring-trial validated method EN ISO 17180:2013 based on ion-exchange chromatography coupled to visible or fluorescence detection (IEC-VIS/FLD) to quantify free *methionine* in the *feed additive* and *premixtures* (containing more than 10 % *methionine*); and (iii) the Community method based on IEC-VIS for the quantification of *methionine* in *premixtures*, *feedingstuffs* and *water*.

Further testing or validation of the methods to be performed through the consortium of National Reference Laboratories as specified by Article 10 (Commission Regulation (EC) No 378/2005), as last amended by Regulation (EU) 2015/1761) is not considered necessary.

KEYWORDS

L-methionine produced by fermentation with Corynebacterium glutamicum KCCM80184 and Escherichia coli K12 KCCM80096, nutritional additives, amino acids, all animal species and categories

1. BACKGROUND

In the current application authorisation is sought under Article 4(1) (authorisation of a new feed additive) for *L-methionine produced by fermentation with Corynebacterium glutamicum KCCM80184 and Escherichia coli K12 KCCM80096*, under the category/functional group 3(c) 'nutritional additives'/amino acids, their salts and analogues', according to Annex I of Regulation (EC) No 1831/2003. Authorisation is sought for all animal species [1-2]. In various forms and analogues, *methionine* produced synthetically or by fermentation with different microorganisms is already authorised as nutritional *feed additive* [3-4].

According to the Applicant, the white crystalline powdered *L-methionine* has a minimum purity (mass fraction) of 98.5 % [1,5]. The *feed additive* is produced by fermentation with genetically modified strains of *Corynebacterium glutamicum* and *Escherichia coli K12* [6]. The production strains are deposited in the "Korean Culture Centre of Microorganisms" (KCCM) with the accession number 80814 for *Corynebacterium glutamicum* and 80096 for *Escherichia coli K12* [7].

The *feed additive* is intended to be added directly into *feedingstuffs* (or through *premixtures*) and *water* for drinking [8]. However, the Applicant did not propose any minimum or maximum content of *L-methionine* in *feedingstuffs* [1,8].

Note: The EURL has previously evaluated the analytical methods for the determination of *methionine* in the frame of two dossiers [9-10].

2. TERMS OF REFERENCE

In accordance with Article 5 of Regulation (EC) No 378/2005, as last amended by Regulation (EU) 2015/1761, on detailed rules for the implementation of Regulation (EC) No 1831/2003 of the European Parliament and of the Council as regards the duties and the tasks of the European Union Reference Laboratory concerning applications for authorisations of feed additives, the EURL is requested to submit a full evaluation report to the European Food Safety Authority for each application or group of applications. For this particular dossier, the methods of analysis submitted in connection with *L-methionine* and their suitability to be used for official controls in the frame of the authorisation were evaluated.

3. EVALUATION

Description of the analytical methods for the determination of the active substance in the feed additive, premixtures, feedingstuffs and when appropriate water (section 2.6.1 of the dossier - Annex II of Commission Regulation (EC) No 429/2008)

For the quantification of *L-methionine* in the *feed additive* the Applicant submitted the method based on titration described in the corresponding European Pharmacopoeia monograph [11,12].

However, the EURL found the ring-trial validated method EN ISO 17180:2013 - "Animal feeding stuffs – Determination of lysine, methionine and threonine in commercial amino acid products and premixtures" [13]. This standard method is based on the experimental protocol described in the Community method for the analysis of free amino acids (including *methionine*) [14]. It does not distinguish between the salts of amino acids and cannot differentiate between enantiomers. It applies for products containing more than 10 % of amino acid.

Free *methionine* is extracted with diluted hydrochloric acid and further diluted with sodium citrate buffer. After addition of norleucine as internal standard, the amino acids are separated by high performance liquid chromatography (HPLC) with an ion-exchange column (IEC). Free *methionine* is quantified either after post-column derivatisation with ninhydrine and Visible (VIS) detection at 440 nm and 570 nm or by fluorescence detection (FLD) after post-column reaction with ortho-phthaldialdehyde with a detector excitation wavelength at 330 nm and emission at 460 nm. The performance characteristics reported for the quantification of free *methionine* are listed in Table 1.

Based on the performance characteristics available, the EURL recommends for official control the EN ISO 17180:2013 method for the quantification of free *methionine* in the *feed additive* and *premixtures* (containing more than 10 % *methionine*).

For the quantification of *L-methionine* in *premixtures* and *feedingstuffs* the Applicant submitted the ring-trial validated Community method mentioned above [11,14]. This method was designed for the quantification of free (synthetic and natural) and total (peptide-bound and free) amino acids in *premixtures* and *feedingstuffs*, using an amino acid analyser or IEC coupled with post-column derivatisation and VIS detection. It does not distinguish between the salts of amino acids and cannot differentiate between enantiomers.

The free amino acids are extracted with diluted hydrochloric acid. Co-extracted nitrogenous macromolecules are precipitated with sulfosalicylic acid and removed by filtration. The solution is filtered and adjusted to pH 2.2. The amino acids are separated by IEC and determined by post-column derivatisation with ninhydrin and photometric detection at 570 nm. The procedure chosen for the determination of the total amino acids depends on the amino acid under investigation. *Methionine* must be oxidised to methionine sulfone prior to hydrolysis. Oxidation is performed at 0 °C with a performic acid/phenol mixture. The excess of oxidation reagent is decomposed with sodium disulfite. The oxidised sample is hydrolysed with hydrochloric acid (6 mol/l) for 23 hours. The hydrolysate is adjusted to pH 2.2. The amino acids are separated by IEC and determined by post-column derivatisation with ninhydrin and photometric detection at 570 nm. *Methionine* is determined as methionine sulfone in hydrolysates of oxidised sample and calculated as *methionine* by introducing the molar mass of the amino acid in the equation used for the calculation of results.

The Community method was ring-trial validated using four different matrices. This method was further ring-trial validated by twenty-three laboratories, resulting in the EN ISO 13903:2005 method [15]. The performance characteristics reported for the quantification of total *methionine* are listed in Table 1. Furthermore, the following limits of quantification were reported for free *methionine* and total *methionine*: 0.04 and 0.25 g/kg *feedingstuffs*, respectively [15].

Based on the performance characteristics available, the EURL recommends for official control the ring-trial validated Community method based on IEC-VIS to quantify *methionine* in *premixtures* and *feedingstuffs*.

The Applicant did not provide experimental data to determine *L-methionine* in *water* [11]. Nevertheless, even if the determination of *methionine* in *water* is not explicitly stated in the scope, as concluded in the previous EURL reports and further specified in the corresponding legislation, the EURL recommends the Community method for official control for the quantification of *methionine* in *water* [3,4-9,10].

Table 1: Method performance characteristics obtained in the frame of ring-trial validation studies (EN ISO 17180:2013 [13], Community method [14] and EN ISO 13903:2005 [15]) for the determination of total *L-methionine* in the *feed additive, premixtures* and *feedingstuffs*.

Ring-Trial	Matrix	<i>methionine</i> content g/kg	RSD _r %	RSD _R %
[13]	Premix 2	270	1.6	1.9
	Premix 3	110	1.2	1.8
	Premix 4	140	0.9	1.9
	Premix 5	320	0.5	2.6
	Premix 6	307	1.1	2.5
	Premix 7	90	0.9	1.5
	Premix 8	89	1.2	1.9
	Premix 9	190	1.0	1.7
	Premix 10	184	0.7	1.5
	DL-methionine	936	0.9	1.5
[14]	Mixed pig feed	3.3	3.4	7
	Broiler compound	5.1	3.1	10.9
	Protein concentrate	12	2.2	13
	Premix	90.2	2.4	6.9
[15]	Poultry meal	11.7	2.1	12
	Broiler finisher feed	5.3	1.1	7.6
	Broiler starter feed	6.2	2.1	10.2
	Corn	1.8	5.6	11.7
	Fishmeal	16.1	1.9	9.7

RSD_r, RSD_R - relative standard deviation for *repeatability* and *reproducibility*, respectively

Methods of analysis for the determination of the residues of the additive in food (section 2.6.2 of the dossier - Annex II of Commission Regulation (EC) No 429/2008)

The evaluation of corresponding methods of analysis is not relevant for the present application.

Identification/Characterisation of the feed additive (section 2.6.3 of the dossier - Annex II of Commission Regulation (EC) No 429/2008)

The EURL found the "L-methionine monograph" of the Food Chemical Codex (FCC) where identification is based on infrared absorption [16].

The EURL recommends the Food Chemical Codex for the identification of *L-methionine* in the *feed additive*.

Further testing or validation of the methods to be performed through the consortium of National Reference Laboratories as specified by Article 10 (Commission Regulation (EC) No 378/2005, as last amended by Regulation (EU) 2015/1761) is not considered necessary.

4. CONCLUSIONS AND RECOMMENDATIONS

In the frame of this authorisation the EURL recommends for official control (i) the "L-methionine monograph" of the Food Chemical Codex (FCC) based on infrared absorption for the identification of *L-methionine* in the *feed additive*; (ii) the ring-trial validated method EN ISO 17180:2013 based on ion-exchange chromatography coupled to visible or fluorescence detection (IEC-VIS/FLD) to quantify free *methionine* in the *feed additive* and *premixtures* (containing more than 10 % *methionine*); and (iii) the Community method based on IEC-VIS for the quantification of *methionine* in *premixtures*, *feedingstuffs* and *water*.

Recommended text for the register entry (analytical method)

For the identification of *L-methionine* in the *feed additive*:

- Food Chemical Codex "L-methionine monograph"

For the quantification of *methionine* in the *feed additive* and *premixtures* (containing more than 10 % *methionine*):

- Ion-exchange chromatography coupled with post-column derivatisation and optical detection (IEC-VIS/FLD) – EN ISO 17180

For the quantification of *methionine* in *premixtures* and *feedingstuffs*:

- Ion-exchange chromatography coupled with post-column derivatisation and photometric detection (IEC-VIS), Commission Regulation (EC) No 152/2009 (Annex III, F)

For the quantification of *methionine* in *water*:

- ion-exchange chromatography coupled with post-column derivatisation and photometric detection (IEC-VIS)

5. DOCUMENTATION AND SAMPLES PROVIDED TO EURL

In accordance with the requirements of Regulation (EC) No 1831/2003, reference samples of *L-methionine produced by fermentation with Corynebacterium glutamicum KCCM80184 and Escherichia coli K12 KCCM80096* have been sent to the European Union Reference Laboratory for Feed Additives. The dossier has been made available to the EURL by EFSA.

6. REFERENCES

- [1] *Application, Proposal of Registry Entry – Annex A
- [2] *Application, Reference SANTE/E5: Forw. Appl. 1831/0078-2018 & Annex I – submission number 1543244172061-2322
- [3] Commission Implementing Regulation (EU) No 469/2013 of 22 May 2013 concerning the authorisation of DL-methionine, DL-methionine sodium salt, hydroxy analogue of methionine, calcium salt of hydroxy analogue of methionine, isopropyl ester of hydroxy analogue of methionine, DL-methionine protected with copolymer vinylpyridine/styrene and DL-methionine protected with ethylcellulose as feed additives, O.J. L 136/1, 23.5.2013
- [4] Commission Implementing Regulation (EU) No 852/2014 of 5 August 2014 concerning the authorisation of L-methionine as a feed additive for all animal species, O.J. L 233/22, 6.8.2014
- [5] *Technical dossier, Section II: II.2.1.1. Chemical substances
- [6] *Technical dossier, Section II: II.2.1.2. Micro-organisms
- [7] *Technical dossier, Section II: II.1.1. Name of the additive
- [8] *Technical dossier, Section II: II.5.1. Proposed mode of use in animal nutrition
- [9] FAD-2012-0016, L-methionine, Ref. JRC.D.5/SFB/CvH/SB/ag/ARES(2012)1459970, <https://ec.europa.eu/jrc/sites/jrcsh/files/FinRep-FAD-2012-0016-L-Methionine.pdf>
- [10] FAD-2010-0023, DL-Methionine; DL-Methionine sodium salt; DL-Methionine protected with ethylcellulose; DL-Methionine protected with copolymer vinylpyridine/styrene; Hydroxy analogue of Methionine; Calcium salt of hydroxy analogue of Methionine; Isopropyl ester of the hydroxylated analogue of Methionine, Ref. JRC.D.5/FSQ/CvH/SB/ag/ARES(2012)240861, <https://ec.europa.eu/jrc/sites/jrcsh/files/FinRep-FAD-2010-0023.pdf>
- [11] *Technical dossier, Section II: II.6.1. Methods of analysis for the active substance
- [12] European Pharmacopoeia 7.2, 01/2008:1027, p. 2379
- [13] EN ISO 17180:2013 - Animal feeding stuffs – Determination of lysine, methionine and threonine in commercial amino acid products and premixtures
- [14] Commission Regulation (EC) No 152/2009 of 27 January 2009 laying down the methods of sampling and analysis for the official control of feed, O.J. L 54, 26.02.2009
- [15] EN ISO 13903:2005- Animal feeding stuffs – Determination of amino acids content
- [16] Food Chemical Codex monograph "L-methionine", FCC 7 (2010), p. 642-643

*Refers to Dossier no: FAD-2018-0085

7. RAPPORTEUR LABORATORY & NATIONAL REFERENCE LABORATORIES

The Rapporteur Laboratory for this evaluation is the European Union Reference Laboratory for Feed Additives, JRC, Geel, Belgium. This report is in accordance with the opinion of the consortium of National Reference Laboratories as referred to in Article 6(2) of Commission Regulation (EC) No 378/2005, as last amended by Regulation (EU) 2015/1761.

8. ACKNOWLEDGEMENTS

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