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Brussels, XXX  
SANTE/3274304/2025  
[...] (2025) XXX draft

**COMMISSION IMPLEMENTING REGULATION (EU) .../...**

**of XXX**

**concerning the authorisation of a preparation of chromium chelate of DL-methionine as a feed additive for salmonids (holder of authorisation: Zinpro Animal Nutrition Europe, Inc.)**

(Text with EEA relevance)

# COMMISSION IMPLEMENTING REGULATION (EU) .../...

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**concerning the authorisation of a preparation of chromium chelate of DL-methionine as a feed additive for salmonids (holder of authorisation: Zinpro Animal Nutrition Europe, Inc.)**

(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<sup>1</sup>, 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 an authorisation.
- (2) In accordance with Article 7 of Regulation (EC) No 1831/2003, an application was submitted for the authorisation of a preparation of chromium chelate of DL-methionine. That application was accompanied by the particulars and documents required under Article 7(3) of Regulation (EC) No 1831/2003.
- (3) The application concerns the authorisation a preparation of preparation of chromium chelate of DL-methionine as a feed additive for salmonids, requesting that additive to be classified in the category ‘zootechnical additives’ and in the functional group ‘other zootechnical additives’.
- (4) The European Food Safety Authority (‘the Authority’) concluded in its opinions of 28 February 2025<sup>2</sup> that, under the proposed conditions of use, the preparation of preparation of chromium chelate of DL-methionine (‘the preparation’) is safe for salmonids, consumers and the environment. It also concluded that the preparation is not a skin or eye irritant but is considered a skin and respiratory sensitiser, while inhalation and dermal exposure is considered a risk. The Authority further concluded that the preparation may have the potential to be efficacious in improving the performance of salmonids. It did not consider that there is a need for specific requirements of post-market monitoring. In accordance with Article 5(4), point (a), of Commission Regulation (EC) No 378/2005<sup>3</sup>, the Reference Laboratory set up by Regulation (EC) No 1831/2003 considered that the conclusions and recommendations

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<sup>1</sup> OJ L 268, 18.10.2003, p. 29, ELI: <http://data.europa.eu/eli/reg/2003/1831/oj>.

<sup>2</sup> EFSA Journal. 2025;23:e9310. <https://doi.org/10.2903/j.efsa.2025.9310>

<sup>3</sup> Commission Regulation (EC) No 378/2005 of 4 March 2005 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 tasks of the Community Reference Laboratory concerning applications for authorisations of feed additives. OJ L 59, 5.3.2005, p. 8, ELI: <http://data.europa.eu/eli/reg/2005/378/oj>

reached in the previous assessment concerning the same additive are valid and applicable for the current application.

- (5) In view of the above, the Commission considers that the preparation of chromium chelate of DL-methionine satisfies the conditions provided for in Article 5 of Regulation (EC) No 1831/2003. Accordingly, the use of that preparation should be authorised. In addition, the Commission considers that appropriate protective measures should be taken to prevent adverse effects on the health of the users of the additive.
- (6) 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*  
**Authorisation**

The preparation specified in the Annex, belonging to the additive category ‘zootechnical additives’ and to the functional group ‘other zootechnical additives’, is authorised as an additive in animal nutrition, subject to the conditions laid down in that Annex.

*Article 2*  
**Entry into force**

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,

*For the Commission*  
*The President*  
Ursula VON DER LEYEN