Commission Implementing Regulation (EU) No 1196/2012 of 13 December 2012 amending Regulation (EU) No 9/2010 as regards the minimum content of a preparation of endo-1,4-beta-xylanase produced by Trichoderma reesei (ATCC PTA 5588) as a feed additive in feed for laying hens (holder of authorisation Danisco Animal Nutrition) (Text with EEA relevance)

COMMISSION IMPLEMENTING REGULATION (EU) No 1196/2012

of 13 December 2012

amending Regulation (EU) No 9/2010 as regards the minimum content of a preparation of endo-1,4-beta-xylanase produced by *Trichoderma reesei* (ATCC PTA 5588) as a feed additive in feed for laying hens (holder of authorisation Danisco Animal Nutrition)

(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 13(3) thereof,

Whereas:

- (1) The use of a preparation of endo-1,4-beta-xylanase (EC 3.2.1.8) produced by *Trichoderma reesei* (ATCC PTA 5588), belonging to the additive category of 'zootechnical additives', was authorised for 10 years as a feed additive for use on chickens for fattening, laying hens, ducks and turkeys for fattening by Commission Regulation (EU) No 9/2010⁽²⁾, for weaned piglets and pigs for fattening by Commission Implementing Regulation (EU) No 528/2011⁽³⁾ and for minor poultry species other than ducks by Commission Implementing Regulation (EU) No 1021/2012⁽⁴⁾.
- (2) In accordance with Article 13(3) of Regulation (EC) No 1831/2003, the holder of the authorisation has proposed changing the terms of the authorisation of the preparation concerned by reducing its minimum content from 2 500 U/kg to 625 U/kg as regards the use on laying hens. The application was accompanied by the relevant supporting data. The Commission forwarded that application to the European Food Safety Authority (hereinafter 'the Authority').
- (3) The Authority concluded in its opinion of 22 May 2012⁽⁵⁾ that, under the new proposed conditions of use, the preparation concerned is efficacious at the requested minimum dose of 625 U/kg. 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 additive in feed submitted by the Reference Laboratory set up by Regulation (EC) No 1831/2003.
- (4) The conditions provided for in Article 5 of Regulation (EC) No 1831/2003 are satisfied.

Status: Point in time view as at 31/01/2020. Changes to legislation: There are currently no known outstanding effects for the Commission Implementing Regulation (EU) No 1196/2012, Introductory Text. (See end of Document for details)

- (5) Regulation (EU) No 9/2010 should therefore be amended accordingly.
- (6) The measures provided for in this Regulation are in accordance with the opinion of the Standing Committee on the Food Chain and Animal Health,

HAS ADOPTED THIS REGULATION:

- (**1**) OJ L 268, 18.10.2003, p. 29.
- (2) OJ L 3, 7.1.2010, p. 10.
- (**3**) OJ L 143, 31.5.2011, p. 10.
- (4) OJ L 307, 7.11.2012, p. 68.
- (5) *EFSA Journal 2012*; 10(6):2739.

Status:

Point in time view as at 31/01/2020.

Changes to legislation:

There are currently no known outstanding effects for the Commission Implementing Regulation (EU) No 1196/2012, Introductory Text.