Changes to legislation: There are outstanding changes not yet made to Commission Implementing Decision (EU) 2015/2279. Any changes that have already been made to the legislation appear in the content and are referenced with annotations. (See end of Document for details)

Commission Implementing Decision (EU) 2015/2279 of 4 December 2015 authorising the placing on the market of products containing, consisting of, or produced from genetically modified maize NK603 × T25 (MON-ØØ6Ø3-6 × ACS-ZMØØ3-2) pursuant to Regulation (EC) No 1829/2003 of the European Parliament and of the Council (notified under document C(2015) 8581) (Only the Dutch and French texts are authentic) (Text with EEA relevance)

COMMISSION IMPLEMENTING DECISION (EU) 2015/2279

of 4 December 2015

authorising the placing on the market of products containing, consisting of, or produced from genetically modified maize NK603 × T25 (MON-ØØ6Ø3-6 × ACS-ZMØØ3-2) pursuant to Regulation (EC) No 1829/2003 of the European Parliament and of the Council

(notified under document C(2015) 8581)

(Only the Dutch and French texts are authentic)

(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 1829/2003 of the European Parliament and of the Council of 22 September 2003 on genetically modified food and feed⁽¹⁾, and in particular Articles 7(3) and 19(3) thereof,

Whereas:

- (1) On 17 May 2010, Monsanto Europe SA submitted to the competent authority of the Netherlands an application in accordance with Articles 5 and 17 of Regulation (EC) No 1829/2003 for the placing on the market of foods, food ingredients, and feed containing, consisting of, or produced from NK603 × T25 maize ('the application').
- (2) The application also covers the placing on the market of genetically modified maize NK603 × T25 in products consisting of it or containing it for other uses than food and feed as any other maize, with the exception of cultivation.
- (3) In accordance with Articles 5(5) and 17(5) of Regulation (EC) No 1829/2003, the application includes the data and information required by Annexes III and IV to Directive 2001/18/EC of the European Parliament and of the Council⁽²⁾ and information and conclusions about the risk assessment carried out in accordance with the principles set out in Annex II to that Directive. It also includes a monitoring plan for environmental effects conforming with Annex VII to Directive 2001/18/EC.
- (4) On 15 July 2015, the European Food Safety Authority ('EFSA') gave a favourable opinion in accordance with Articles 6 and 18 of Regulation (EC) No 1829/2003.

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It concluded that genetically modified maize NK603 \times T25, as described in the application, is as safe as its non-genetically modified comparator and other non-genetically modified maize varieties with respect to potential effects on human and animal health and the environment in the context of its scope⁽³⁾.

- (5) In its opinion, EFSA considered all the specific questions and concerns raised by the Member States in the context of the consultation of the national competent authorities as provided for by Articles 6(4) and 18(4) of Regulation (EC) No 1829/2003.
- (6) In its opinion, EFSA also concluded that the monitoring plan for environmental effects, consisting of a general surveillance plan, submitted by the applicant is in line with the intended uses of the products.
- (7) Taking into account these considerations, authorisation should be granted to the products containing, consisting of, or produced from genetically modified maize NK603 × T25.
- (8) A unique identifier should be assigned to each genetically modified organism (hereinafter 'GMO') as provided for in Commission Regulation (EC) No 65/2004⁽⁴⁾.
- (9) On the basis of the EFSA opinion, no specific labelling requirements, other than those provided for in Articles 13(1) and 25(2) of Regulation (EC) No 1829/2003, appear to be necessary for food, food ingredients and feed containing, consisting of, or produced from genetically modified maize NK603 × T25. However, in order to ensure the use of those products within the limits of the authorisation granted by this Decision, the labelling of the products containing or consisting of NK603 × T25 maize, with the exception of food products, should be complemented by a clear indication that the products in question are not intended for cultivation.
- (10) Regulation (EC) No 1830/2003 of the European Parliament and of the Council⁽⁵⁾ lays down labelling requirements in Article 4(6) for products containing or consisting of GMOs. Traceability requirements for those products are laid down in paragraphs 1 to 5 of Article 4 and traceability requirements for food and feed produced from GMOs are laid down in Article 5 of that Regulation.
- (11) The authorisation holder should submit annual reports on the implementation and the results of the activities set out in the monitoring plan for environmental effects. Those results should be presented in accordance with Commission Decision 2009/770/EC⁽⁶⁾. The EFSA opinion does not justify either the imposition of specific conditions or restrictions for the placing on the market and/or the use and handling of the food and feed, including post-market monitoring requirements or specific conditions for the protection of particular ecosystems/environment and/or geographical areas, as provided for in Articles 6(5)(e) and 18(5)(e) of Regulation (EC) No 1829/2003.
- (12) All relevant information on the authorisation of the products should be entered in the Community register of genetically modified food and feed provided for in Regulation (EC) No 1829/2003.
- (13) This Decision is to be notified through the Biosafety Clearing-House to the Parties to the Cartagena Protocol on Biosafety to the Convention on Biological Diversity, pursuant to

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- Articles 9(1) and 15(2)(c) of Regulation (EC) No 1946/2003 of the European Parliament and of the Council⁽⁷⁾.
- (14) The Standing Committee on Plants, Animals, Food and Feed has not delivered an opinion within the time-limit laid down by its Chairman. An implementing act was deemed to be necessary and the chair submitted the draft implementing act to the appeal committee for further deliberation. The appeal committee did not deliver an opinion,

HAS ADOPTED THIS DECISION:

Article 1

Genetically modified organism and unique identifier

Genetically modified maize (*Zea mays* L.) NK603 \times T25, as specified in point (b) of the Annex to this Decision, is assigned the unique identifier MON- $\emptyset\emptyset6\emptyset3$ -6 \times ACS-ZM $\emptyset\emptyset3$ -2, as provided for in Regulation (EC) No 65/2004.

Article 2

Authorisation

The following products are authorised for the purposes of Articles 4(2) and 16(2) of Regulation (EC) No 1829/2003 in accordance with the conditions set out in this Decision:

- (a) foods and food ingredients containing, consisting of, or produced from MON-ØØ6Ø3-6 × ACS-ZMØØ3-2 maize;
- (b) feed containing, consisting of, or produced from MON-ØØ6Ø3-6 × ACS-ZMØØ3-2 maize;
- (c) MON- $\emptyset\emptyset6\emptyset3$ -6 × ACS-ZM $\emptyset\emptyset3$ -2 maize in products containing it or consisting of it for any other use than those provided in points (a) and (b), with the exception of cultivation.

Article 3

Labelling

- 1 For the purposes of the labelling requirements laid down in Articles 13(1) and 25(2) of Regulation (EC) No 1829/2003 and in Article 4(6) of Regulation (EC) No 1830/2003, the 'name of the organism' shall be 'maize'.
- The words 'not for cultivation' shall appear on the label of and in the documents accompanying products containing or consisting of MON- $\emptyset\emptyset6\emptyset3-6 \times ACS-ZM\emptyset\emptyset3-2$ maize, with the exception of products referred to in point (a) of Article 2.

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Article 4

Monitoring for environmental effects

- 1 The authorisation holder shall ensure that the monitoring plan for environmental effects, as set out in point (h) of the Annex, is put in place and implemented.
- 2 The authorisation holder shall submit to the Commission annual reports on the implementation and the results of the activities set out in the monitoring plan in accordance with Decision 2009/770/EC.

Article 5

Community register

The information set out in the Annex to this Decision shall be entered in the Community register of genetically modified food and feed, as provided for in Article 28 of Regulation (EC) No 1829/2003.

I^{F1}Article 6

Authorisation holder

The authorisation holder shall be Bayer Agriculture BVBA, Belgium, representing Monsanto Company, United States of America.]

Textual Amendments

F1 Substituted by Commission Implementing Decision (EU) 2019/1579 of 18 September 2019 amending Decisions 2008/933/EC, 2009/813/EC, 2009/814/EC and 2010/429/EU and Implementing Decisions 2012/82/EU, 2012/83/EU, 2012/347/EU, 2013/649/EU, (EU) 2015/683, (EU) 2015/684, (EU) 2015/685, (EU) 2015/686, (EU) 2015/687, (EU) 2015/688, (EU) 2015/689, (EU) 2015/693, (EU) 2015/695, (EU) 2015/696, (EU) 2015/700, (EU) 2015/701, (EU) 2015/2279, (EU) 2015/2281, (EU) 2016/1216, (EU) 2016/1217, (EU) 2017/1207, (EU) 2018/1111, (EU) 2018/2045 and (EU) 2018/2046 as regards the representative of the authorisation holder for placing on the market certain genetically modified food and feed in the Union (notified under document C(2019) 6520) (Only the Dutch text is authentic) (Text with EEA relevance).

Article 7

Validity

This Decision shall apply for a period of 10 years from the date of its notification.

Changes to legislation: There are outstanding changes not yet made to Commission Implementing Decision (EU) 2015/2279. Any changes that have already been made to the legislation appear in the content and are referenced with annotations. (See end of Document for details)

I^{F1}Article 8

Addressee

This Decision is addressed to Bayer Agriculture BVBA, Scheldelaan 460, 2040 Antwerp, Belgium.]

Textual Amendments

F1 Substituted by Commission Implementing Decision (EU) 2019/1579 of 18 September 2019 amending Decisions 2008/933/EC, 2009/813/EC, 2009/814/EC and 2010/429/EU and Implementing Decisions 2012/82/EU, 2012/83/EU, 2012/347/EU, 2013/649/EU, (EU) 2015/683, (EU) 2015/684, (EU) 2015/685, (EU) 2015/686, (EU) 2015/687, (EU) 2015/688, (EU) 2015/689, (EU) 2015/693, (EU) 2015/695, (EU) 2015/696, (EU) 2015/700, (EU) 2015/701, (EU) 2015/2279, (EU) 2015/2281, (EU) 2016/1216, (EU) 2016/1217, (EU) 2017/1207, (EU) 2018/1111, (EU) 2018/2045 and (EU) 2018/2046 as regards the representative of the authorisation holder for placing on the market certain genetically modified food and feed in the Union (notified under document C(2019) 6520) (Only the Dutch text is authentic) (Text with EEA relevance).

Changes to legislation: There are outstanding changes not yet made to Commission Implementing Decision (EU) 2015/2279. Any changes that have already been made to the legislation appear in the content and are referenced with annotations. (See end of Document for details)

ANNEX

(a) [F1Applicant and authorisation holder:

Name : Bayer Agriculture BVBA

Address : Scheldelaan 460, 2040 Antwerp, Belgium

On behalf of Monsanto Company — 800 N. Lindbergh Boulevard — St. Louis, Missouri 63167 — United States of America.]

(b) **Designation and specification of the products:**

- foods and food ingredients containing, consisting of, or produced from MON-ØØ6Ø3-6 × ACS-ZMØØ3-2 maize;
- (2) feed containing, consisting of, or produced from MON- $\emptyset\emptyset6\emptyset3-6 \times ACS-ZM\emptyset\emptyset3-2$ maize;
- (3) MON-ØØ6Ø3-6 × ACS-ZMØØ3-2 maize in products containing it or consisting of it for any other use than those provided in points (1) and (2), with the exception of cultivation.

The genetically modified MON- $\emptyset\emptyset6\emptyset3-6 \times$ ACS-ZM $\emptyset\emptyset3-2$ maize, as described in the application, expresses the CP4 EPSPS protein which confers tolerance to glyphosate herbicides and PAT protein which confers tolerance to glufosinate ammonium herbicides.

(c) Labelling:

- (1) For the purposes of the labelling requirements laid down in Articles 13(1) and 25(2) of Regulation (EC) No 1829/2003, and in Article 4(6) of Regulation (EC) No 1830/2003, the 'name of the organism' shall be 'maize';
- The words 'not for cultivation' shall appear on the label of and in the accompanying documents of the products containing or consisting of MON-ØØ6Ø3-6 × ACS-ZMØØ3-2 maize with the exception of products referred to in point (a) of Article 2.

(d) Method for detection:

- (1) Event-specific real-time quantitative PCR-based methods for MON-ØØ6Ø3-6 and ACS-ZMØØ3-2 maize; the detection methods are validated on the single-trait events and verified on genomic DNA extracted from seeds of MON-ØØ6Ø3-6 × ACS-ZMØØ3-2 maize;
- Validated by the EU Reference Laboratory established under Regulation (EC) No 1829/2003, published at http://gmo-crl.jrc.ec.europa.eu/statusofdossiers.aspx)
- (3) Reference material: ERM®-BF415 (for MON-ØØ6Ø3-6) accessible via the Joint Research Centre (JRC) of the European Commission, Institute for Reference Materials and Measurements (IRMM) at https://irmm.jrc.ec.europa.eu/rmcatalogue) and AOCS 0306-H6 and AOCS 0306-C2 (for ACS-ZMØØ3-2) accessible via the American Oil Chemists Society at http://www.aocs.org/LabServices/content.cfm?ItemNumber=19248)

(e) Unique identifier:

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$MON-\emptyset\emptyset6\emptyset3-6 \times ACS-ZM\emptyset\emptyset3-2$

(f) Information required under Annex II to the Cartagena Protocol on Biosafety to the Convention on Biological Diversity:

> (Biosafety Clearing-House, Record ID number: published in the Community register of genetically modified food and feed when notified).

Conditions or restrictions on the placing on the market, use or handling of the (g) products:

Not required.

(h) Monitoring plan for environmental effects:

> Monitoring plan for environmental effects conforming with Annex VII to Directive 2001/18/EC.

> (Link: plan published in the Community register of genetically modified food and feed)

Post-market monitoring requirements for the use of the food for human (i) consumption:

Not required.

Note: links to relevant documents may need to be modified over the time. Those modifications will be made available to the public via the updating of the Community register of genetically modified food and feed.

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- (1) OJ L 268, 18.10.2003, p. 1.
- (2) Directive 2001/18/EC of the European Parliament and of the Council of 12 March 2001 on the deliberate release into the environment of genetically modified organisms and repealing Council Directive 90/220/EEC (OJ L 106, 17.4.2001, p. 1).
- (3) EFSA GMO Panel (EFSA Panel on Genetically Modified Organisms), 2015. Scientific Opinion on application (EFSA-GMO-NL-2010-80) for the placing on the market of herbicide tolerant genetically modified maize NK603 × T25 for food and feed uses, import and processing under Regulation (EC) No 1829/2003 from Monsanto. *EFSA Journal*: 2015; 13(7):4165, 23 pp. doi:10.2903/j.efsa.2015.4165.
- (4) Commission Regulation (EC) No 65/2004 of 14 January 2004 establishing a system for the development and assignment of unique identifiers for genetically modified organisms (OJ L 10, 16.1.2004, p. 5).
- (5) Regulation (EC) No 1830/2003 of the European Parliament and of the Council of 22 September 2003 concerning the traceability and labelling of genetically modified organisms and the traceability of food and feed products produced from genetically modified organisms and amending Directive 2001/18/EC (OJ L 268, 18.10.2003, p. 24).
- (6) Commission Decision 2009/770/EC of 13 October 2009 establishing standard reporting formats for presenting the monitoring results of the deliberate release into the environment of genetically modified organisms, as or in products, for the purpose of placing on the market, pursuant to Directive 2001/18/EC of the European Parliament and of the Council (OJ L 275, 21.10.2009, p. 9).
- (7) Regulation (EC) No 1946/2003 of the European Parliament and of the Council of 15 July 2003 on transboundary movements of genetically modified organisms (OJ L 287, 5.11.2003, p. 1).

Status:

Point in time view as at 24/09/2019.

Changes to legislation:

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