
STATUTORY INSTRUMENTS

2023 No. 235

AGRICULTURE, ENGLAND
FOOD, ENGLAND

The Genetically Modified Food and Feed (Authorisations and Modifications of Authorisations) (England) Regulations 2023

Made - - - - 28th February 2023
Laid before Parliament 2nd March 2023
Coming into force - - 26th April 2023

The Secretary of State makes these Regulations in exercise of the powers conferred by Articles 7(3), 19(3) and 35(3) of Regulation (EC) No. 1829/2003 of the European Parliament and of the Council on genetically modified food and feed⁽¹⁾.

As required by Article 9 of Regulation (EC) No. 178/2002 of the European Parliament and of the Council laying down the general principles and requirements of food law, establishing the European Food Safety Authority and laying down procedures in matters of food safety⁽²⁾, there has been open and transparent public consultation during the preparation and evaluation of these Regulations.

PART 1

Introduction

Citation, commencement, extent and application

1. These Regulations—

- (a) may be cited as the Genetically Modified Food and Feed (Authorisations and Modifications of Authorisations) (England) Regulations 2023;
- (b) come into force on 26th April 2023;

(1) EUR 2003/1829, amended by S.I. 2019/705 and 2022/377. There are other amendments, but these are not relevant for this instrument. The terms “appropriate authority” and “prescribe” are defined in Article 2 of EUR 2003/1829. With respect to food, Article 7(3) applies in relation to modifications, and renewals of authorisations as read with Articles 9(2) and 11 respectively. With respect to feed, Article 19(3) applies in relation to modifications, and renewals of authorisations as read with Articles 21(2) and 23 respectively.

(2) EUR 2002/178; relevant amending instrument S.I. 2019/641. There are other amendments, but these are not relevant for this instrument.

- (c) extend to England and Wales; and
- (d) apply in relation to England only.

Interpretation

2. In these Regulations—

“Regulation 1829/2003” means Regulation (EC) No. 1829/2003 of the European Parliament and of the Council on genetically modified food and feed;

“Regulation 1830/2003” means Regulation (EC) No. 1830/2003 of the European Parliament and of the Council 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(3);

“Decision 2009/770” means Commission Decision 2009/770/EC 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 the placing on the market, pursuant to Directive 2001/18/EC of the European Parliament and of the Council(4).

PART 2

Authorisations

Authorisation of the placing on the market of products containing, consisting of, or produced from genetically modified organisms

3.—(1) Schedules 1 to 7, which contain authorisations for products containing, consisting of, or produced from genetically modified organisms, have effect.

(2) Schedule 8, which contains an authorisation for products containing or consisting of genetically modified organisms, has effect.

PART 3

Modifications of existing authorisations

Amendment of Commission Decision 2011/891/EU

4.—(1) Commission Decision 2011/891/EU authorising the placing on the market of products containing, consisting of, or produced from genetically modified cotton 281-24-236×3006-210-23 (DAS-24236-5×DAS-21Ø23-5) pursuant to Regulation (EC) No 1829/2003 of the European Parliament and of the Council(5) is amended as follows.

(2) In Article 6 (authorisation holder), for the text substitute—

“The authorisation holder is Corteva Agriscience LLC, United States of America, represented in Great Britain by Corteva Agriscience UK Limited.”.

(3) In Article 8 (addressee), for the text substitute—

(3) EUR 2003/1830; relevant amending instruments are S.I. 2019/90, 2019/778 and 2020/1421.

(4) EUDN 2009/770, amended by S.I. 2019/90.

(5) EUDN 2011/891, amended by S.I. 2019/705.

“This Decision is addressed to Corteva Agriscience LLC, 9330 Zionsville Road, Indianapolis, Indiana, 46268-1054, United States of America, represented in Great Britain by Corteva Agriscience UK Limited, Cpc2 Capital Park, Fulbourn, Cambridge, England, CB21 5XE.”.

(4) In the Annex for point (a) (applicant and authorisation holder) substitute—

“(a) **Authorisation holder**

(1) The authorisation holder is Corteva Agriscience LLC, 9330 Zionsville Road, Indianapolis, Indiana, 46268-1054, United States of America.

(2) The authorisation holder is represented in Great Britain by Corteva Agriscience UK Limited, Cpc2 Capital Park, Fulbourn, Cambridge, England, CB21 5XE.”.

Amendment of Commission Implementing Decision 2012/84/EU

5.—(1) Commission Implementing [Decision 2012/84/EU](#) authorising the placing on the market of products containing, consisting of, or produced from genetically modified soybean 356043 (DP-356043-5) pursuant to Regulation (EC) [No 1829/2003](#) of the European Parliament and of the Council(6) is amended as follows.

(2) In Article 6 (authorisation holder), for the text substitute—

“The authorisation holder is Corteva Agriscience LLC, United States of America, represented in Great Britain by Corteva Agriscience UK Limited.”.

(3) In Article 8 (addressee), for the text substitute—

“This Decision is addressed to Corteva Agriscience LLC, 9330 Zionsville Road, Indianapolis, Indiana, 46268-1054, United States of America, represented in Great Britain by Corteva Agriscience UK Limited, Cpc2 Capital Park, Fulbourn, Cambridge, England, CB21 5XE.”.

(4) In the Annex, for point (a) (applicant and authorisation holder) substitute—

“(a) **Authorisation holder**

(1) The authorisation holder is Corteva Agriscience LLC, 9330 Zionsville Road, Indianapolis, Indiana, 46268-1054, United States of America.

(2) The authorisation holder is represented in Great Britain by Corteva Agriscience UK Limited, Cpc2 Capital Park, Fulbourn, Cambridge, England, CB21 5XE.”.

Amendment of Commission Implementing Decision 2013/648/EU

6.—(1) Commission Implementing [Decision 2013/648/EU](#) authorising the placing on the market of products containing, consisting of, or produced from genetically modified maize MON89034 × 1507 × NK603 (MON-89034-3 × DAS-01507-1 × MON-00603-6) pursuant to Regulation (EC) [No 1829/2003](#) of the European Parliament and of the Council(7) is amended as follows.

(2) In Article 6 (authorisation holders), for paragraph 1, substitute—

“(1) The authorisations holders are:

(a) Corteva Agriscience LLC, United States of America, represented in Great Britain by Corteva Agriscience UK Limited, and

(b) Bayer CropScience LP, United States of America, represented in Great Britain by Bayer CropScience Limited.”.

(3) In Article 8 (addressees), for the text substitute—

“This Decision is addressed to:

(6) EUDN 2012/84, amended by [S.I. 2019/705](#).

(7) EUDN 2013/648, amended by [S.I. 2019/705](#).

- (a) Corteva Agriscience LLC, 9330 Zionsville Road, Indianapolis, Indiana, 46268-1054, United States of America, represented in Great Britain by Corteva Agriscience UK Limited, Cpc2 Capital Park, Fulbourn, Cambridge, England, CB21 5XE, and
 - (b) Bayer CropScience LP, 800 N. Lindbergh Boulevard, St. Louis, Missouri 63167, United States of America, represented in Great Britain by Bayer CropScience Limited, 230 Cambridge Science Park, Milton Road, Cambridge, England, CB4 0WB.”.
- (4) In the Annex, for point (a) (applicants and authorisation holders) substitute—
- “(a) **Authorisation holders**
- (1) Corteva Agriscience LLC, 9330 Zionsville Road, Indianapolis, Indiana, 46268-1054, United States of America, represented in Great Britain by Corteva Agriscience UK Limited, Cpc2 Capital Park, Fulbourn, Cambridge, England, CB21 5XE, and
 - (2) Bayer CropScience LP, 800 N. Lindbergh Boulevard, St. Louis, Missouri 63167, United States of America, represented in Great Britain by Bayer CropScience Limited, 230 Cambridge Science Park, Milton Road, Cambridge, England, CB4 0WB.”.

Amendment of Commission Implementing Decision 2013/650/EU

7.—(1) Commission Implementing [Decision 2013/650/EU](#) authorising the placing on the market of products containing, consisting of, or produced from genetically modified (GM) maize MON 89034 × 1507 × MON88017 × 59122 (MON-89034-3 × DAS-01507-1 × MON-88017-3 × DAS-59122-7), four related GM maizes combining three different single GM events (MON89034 × 1507 × MON88017 (MON-89034-3 × DAS-01507-1 × MON-88017-3), MON89034 × 1507 × 59122 (MON-89034-3 × DAS-01507-1 × DAS-59122-7), MON89034 × MON88017 × 59122 (MON-89034-3 × MON-88017-3 × DAS-59122-7), 1507 × MON 88017 × 59122 (DAS-01507-1 × MON-88017-3 × DAS-59122-7)) and four related GM maizes combining two different single GM events (MON89034 × 1507 (MON-89034-3 × DAS-01507-1), MON89034 × 59122 (MON-89034-3 × DAS-59122-7), 1507 × MON88017 (DAS-01507-1 × MON-88017-3), MON 88017 × 59122 (MON-88017-3 × DAS-59122-7)) pursuant to Regulation [\(EC\) No 1829/2003](#) of the European Parliament and of the Council⁽⁸⁾ is amended as follows.

- (2) In Article 6 (authorisation holders), for paragraph 1, substitute—
- “(1) The authorisation holders are:
- (a) Corteva Agriscience LLC, United States of America, represented in Great Britain by Corteva Agriscience UK Limited, and
 - (b) Bayer CropScience LP, United States of America, represented in Great Britain by Bayer CropScience Limited.”.

- (3) In Article 8 (addressees), for the text substitute—

“This Decision is addressed to:

- (a) Corteva Agriscience LLC, 9330 Zionsville Road, Indianapolis, Indiana, 46268-1054, United States of America, represented in Great Britain by Corteva Agriscience UK Limited, Cpc2 Capital Park, Fulbourn, Cambridge, England, CB21 5XE, and
 - (b) Bayer CropScience LP, 800 N. Lindbergh Boulevard, St. Louis, Missouri 63167, United States of America, represented in Great Britain by Bayer CropScience Limited, 230 Cambridge Science Park, Milton Road, Cambridge, England, CB4 0WB.”.
- (4) In the Annex, for point (a) (applicant and authorisation holders) substitute—

“(a) **Authorisation holders**

⁽⁸⁾ EUDN 2013/650, amended by [S.I. 2019/705](#).

(1) Corteva Agriscience LLC, 9330 Zionsville Road, Indianapolis, Indiana, 46268-1054, United States of America, represented in Great Britain by Corteva Agriscience UK Limited, Cpc2 Capital Park, Fulbourn, Cambridge, England, CB21 5XE, and

(2) Bayer CropScience LP, 800 N. Lindbergh Boulevard, St. Louis, Missouri 63167, United States of America. represented in Great Britain by Bayer CropScience Limited, 230 Cambridge Science Park, Milton Road, Cambridge, England, CB4 0WB.”.

Amendment of Commission Implementing Decision (EU) 2015/698

8.—(1) Commission Implementing Decision (EU) 2015/698 authorising the placing on the market of products containing, consisting of, or produced from genetically modified soybean 305423 (DP-3Ø5423-1) pursuant to Regulation (EC) No 1829/2003 of the European Parliament and of the Council⁽⁹⁾ is amended as follows.

(2) In Article 7 (authorisation holder), for the text substitute—

“The authorisation holder is Corteva Agriscience LLC, United States of America, represented in Great Britain by Corteva Agriscience UK Limited.”.

(3) In Article 9 (addressee), for the text substitute—

“This Decision is addressed to Corteva Agriscience LLC, 9330 Zionsville Road, Indianapolis, Indiana, 46268-1054, United States of America, represented in Great Britain by Corteva Agriscience UK Limited, Cpc2 Capital Park, Fulbourn, Cambridge, England, CB21 5XE.”.

(4) In the Annex, for point (a) (applicant and authorisation holder) substitute—

“(a) **Authorisation holder**

1. The authorisation holder is Corteva Agriscience LLC, 9330 Zionsville Road, Indianapolis, Indiana, 46268-1054, United States of America.

2. The authorisation holder is represented in Great Britain by Corteva Agriscience UK Limited, Cpc2 Capital Park, Fulbourn, Cambridge, England, CB21 5XE.”.

Amendment of Commission Implementing Decision (EU) 2016/1215

9.—(1) Commission Implementing Decision (EU) 2016/1215 authorising the placing on the market of products containing, consisting of, or produced from genetically modified soybean FG72 (MST- FGØ72-2) pursuant to Regulation (EC) No 1829/2003 of the European Parliament and of the Council⁽¹⁰⁾ is amended as follows.

(2) In Article 6 (authorisation holder), for the text substitute—

“The authorisation holder is Syngenta Crop Protection AG, Switzerland represented in Great Britain by Syngenta Limited.”.

(3) In Article 8 (addressee), for the text substitute—

“This Decision is addressed to Syngenta Crop Protection AG, Rosentalstrasse 67, CH-4058 Basel, Switzerland, represented in Great Britain by Syngenta Limited, Jealott’s Hill International Research Centre, Bracknell, Berkshire, England, RG42 6EY.”.

(4) In the Annex, for point (a) (applicant and authorisation holder) substitute—

“(a) **Authorisation holder:**

(1) The authorisation holder is Syngenta Crop Protection AG, Rosentalstrasse 67, CH-4058 Basel, Switzerland.

⁽⁹⁾ EUDN 2015/698, amended by [S.I. 2019/705](#).

⁽¹⁰⁾ EUDN 2016/1215, amended by [S.I. 2019/705](#).

(2) The authorisation holder is represented in Great Britain by Syngenta Limited, Jealott's Hill International Research Centre, Bracknell, Berkshire, England, RG42 6EY.”.

Amendment of Commission Implementing Decision (EU) 2017/1211

10.—(1) Commission Implementing Decision (EU) 2017/1211 authorising the placing on the market of products containing, consisting of, or produced from genetically modified cotton 281-24-236 × 3006-210-23 × MON 88913 (DAS-24236-5 × DAS-21Ø23-5 × MON-88913-8) pursuant to Regulation (EC) No 1829/2003 of the European Parliament and of the Council⁽¹¹⁾ is amended as follows.

(2) In Article 6 (authorisation holder), for the text substitute—

“The authorisation holder is Corteva Agriscience LLC, United States of America, represented in Great Britain by Corteva Agriscience UK Limited.”.

(3) In Article 8 (addressee), for the text substitute—

“This Decision is addressed to Corteva Agriscience LLC, 9330 Zionsville Road, Indianapolis, Indiana, 46268-1054, United States of America, represented in Great Britain by Corteva Agriscience UK Limited, Cpc2 Capital Park, Fulbourn, Cambridge, England, CB21 5XE.”.

(4) In the Annex, for point (a) (applicant and authorisation holder) substitute—

“(a) **Authorisation holder:**

(1) The authorisation holder is Corteva Agriscience LLC, 9330 Zionsville Road, Indianapolis, Indiana, 46268-1054, United States of America.

(2) The authorisation holder is represented in Great Britain by Corteva Agriscience UK Limited, Cpc2 Capital Park, Fulbourn, Cambridge, England, CB21 5XE.”.

Amendment of Commission Implementing Decision (EU) 2017/1212

11.—(1) Commission Implementing Decision (EU) 2017/1212 authorising the placing on the market of products containing, consisting of, or produced from genetically modified maize DAS-40278-9, pursuant to Regulation (EC) No 1829/2003 of the European parliament and of the Council on genetically modified food and feed⁽¹²⁾ is amended as follows.

(2) In Article 6 (authorisation holder), for the text substitute—

“The authorisation holder is Corteva Agriscience LLC, United States of America, represented in Great Britain by Corteva Agriscience UK Limited.”.

(3) In Article 8 (addressee), for the text substitute—

“This Decision is addressed to Corteva Agriscience LLC, 9330 Zionsville Road, Indianapolis, Indiana, 46268-1054, United States of America, represented in Great Britain by Corteva Agriscience UK Limited, Cpc2 Capital Park, Fulbourn, Cambridge, England, CB21 5XE.”.

(4) In the Annex, for point (a) (authorisation holder) substitute—

“(a) **Authorisation holder:**

(1) The authorisation holder is Corteva Agriscience LLC, 9330 Zionsville Road, Indianapolis, Indiana, 46268-1054, United States of America.

(2) The authorisation holder is represented in Great Britain by Corteva Agriscience UK Limited, Cpc2 Capital Park, Fulbourn, Cambridge, England, CB21 5XE.”.

⁽¹¹⁾ EUDN 2017/1211, amended by S.I. 2019/705.

⁽¹²⁾ EUDN 2017/1212, amended by S.I. 2019/705.

Amendment of Commission Implementing Decision (EU) 2017/2448

12.—(1) Commission Implementing Decision (EU) 2017/2448 authorising the placing on the market of products containing, consisting of, or produced from genetically modified soybean 305423 × 40-3-2 (DP-305423-1 × MON-04032-6) pursuant to Regulation (EC) No 1829/2003 of the European Parliament and of the Council on genetically modified food and feed⁽¹³⁾ is amended as follows.

(2) In Article 7 (authorisation holder), for the text substitute—

“The authorisation holder is Corteva Agriscience LLC, United States of America, represented in Great Britain by Corteva Agriscience UK Limited.”.

(3) In Article 9 (addressee), for the text substitute—

“This Decision is addressed to Corteva Agriscience LLC, 9330 Zionsville Road, Indianapolis, Indiana, 46268-1054, United States of America, represented in Great Britain by Corteva Agriscience UK Limited, Cpc2 Capital Park, Fulbourn, Cambridge, England, CB21 5XE.”.

(4) In the Annex, for point (a) (applicant and authorisation holder) substitute—

“(a) **Authorisation holder:**

(1) The authorisation holder is Corteva Agriscience LLC, 9330 Zionsville Road, Indianapolis, Indiana, 46268-1054, United States of America.

(2) The authorisation holder is represented in Great Britain by Corteva Agriscience UK Limited, Cpc2 Capital Park, Fulbourn, Cambridge, England, CB21 5XE.”.

Amendment of Commission Implementing Decision (EU) 2017/2449

13.—(1) Commission Implementing Decision (EU) 2017/2449 authorising the placing on the market of products containing, consisting of, or produced from genetically modified soybean DAS-68416-4, pursuant to Regulation (EC) No 1829/2003 of the European Parliament and of the Council on genetically modified food and feed⁽¹⁴⁾ is amended as follows.

(2) In Article 7 (authorisation holder), for the text substitute—

“The authorisation holder is Corteva Agriscience LLC, United States of America, represented in Great Britain by Corteva Agriscience UK Limited.”.

(3) In Article 9 (addressee), for the text substitute—

“This Decision is addressed to Corteva Agriscience LLC, 9330 Zionsville Road, Indianapolis, Indiana, 46268-1054, United States of America, represented in Great Britain by Corteva Agriscience UK Limited, Cpc2 Capital Park, Fulbourn, Cambridge, England, CB21 5XE.”.

(4) In the Annex, for point (a) (authorisation holder) substitute—

“(a) **Authorisation holder:**

(1) The authorisation holder is Corteva Agriscience LLC, 9330 Zionsville Road, Indianapolis, Indiana, 46268-1054, United States of America.

(2) The authorisation holder is represented in Great Britain by Corteva Agriscience UK Limited, Cpc2 Capital Park, Fulbourn, Cambridge, England, CB21 5XE.”.

Amendment of Commission Implementing Decision (EU) 2017/2450

14.—(1) Commission Implementing Decision (EU) 2017/2450 authorising the placing on the market of products containing, consisting of, or produced from genetically modified soybean

⁽¹³⁾ EUDN 2017/2448, amended by [S.I. 2019/705](#).

⁽¹⁴⁾ EUDN 2017/2449, amended by [S.I. 2019/705](#).

DAS-44406-6, pursuant to Regulation (EC) No 1829/2003 of the European Parliament and of the Council on genetically modified food and feed(15) is amended as follows.

(2) In Article 7 (authorisation holder)—

- (a) in the heading, for “holder” substitute “holders”;
- (b) for the text substitute—

“The authorisation holders are Corteva Agriscience LLC, United States of America and M.S. Technologies LLC, United States of America, both represented in Great Britain by Corteva Agriscience UK Limited.”.

(3) In Article 9 (addressee)—

- (a) in the heading, for “Addressee” substitute “Addressees”;
- (b) for the text substitute—

“This Decision is addressed to Corteva Agriscience LLC, 9330 Zionsville Road, Indianapolis, Indiana, 46268-1054, United States of America and to M.S. Technologies LLC, 103 Avenue D, West Point, IA 52656, United States of America, represented in Great Britain by Corteva Agriscience UK Limited, Cpc2 Capital Park, Fulbourn, Cambridge, England, CB21 5XE.”.

(4) In the Annex, for point (a) (authorisation holder) substitute—

“(a) **Authorisation holders:**

(1) The authorisation holders are—

Corteva Agriscience LLC, 9330 Zionsville Road, Indianapolis, Indiana, 46268-1054, United States of America; and

M.S. Technologies LLC, 103 Avenue D, West Point, IA 52656, United States of America.

(2) Both authorisation holders are represented in Great Britain by Corteva Agriscience UK Limited, Cpc2 Capital Park, Fulbourn, Cambridge, England, CB21 5XE.”.

Amendment of Commission Implementing Decision (EU) 2017/2452

15.—(1) Commission Implementing Decision (EU) 2017/2452 renewing the authorisation for the placing on the market of products containing, consisting of, or produced from genetically modified maize 1507 (DAS-Ø15Ø7-1) pursuant to Regulation (EC) No 1829/2003 of the European Parliament and of the Council(16) is amended as follows.

(2) In Article 7 (authorisation holders)—

- (a) in the heading, for “holders” substitute “holder”; and
- (b) for paragraphs 1 and 2 substitute—

“The authorisation holder is Corteva Agriscience LLC, United States of America, represented in Great Britain by Corteva Agriscience UK Limited.”.

(3) In Article 9 (addressee), for the text substitute—

“This Decision is addressed to Corteva Agriscience LLC, 9330 Zionsville Road, Indianapolis, Indiana, 46268-1054, United States of America, represented in Great Britain by Corteva Agriscience UK Limited, Cpc2 Capital Park, Fulbourn, Cambridge, England, CB21 5XE.”.

(4) In the Annex, for point (a) (applicants and authorisation holders) substitute—

“(a) **Authorisation holder:**

(15) EUDN 2017/2450, amended by S.I. 2019/705.

(16) EUDN 2017/2452, amended by S.I. 2019/705.

(1) The authorisation holder is Corteva Agriscience LLC, 9330 Zionsville Road, Indianapolis, Indiana, 46268-1054, United States of America.

(2) The authorisation holder is represented in Great Britain by Corteva Agriscience UK Limited, Cpc2 Capital Park, Fulbourn, Cambridge, England, CB21 5XE.”.

Amendment of Commission Implementing Decision (EU) 2018/1109

16.—(1) Commission Implementing Decision (EU) 2018/1109 renewing the authorisation for the placing on the market of products containing, consisting of, or produced from genetically modified maize 59122 (DAS-59122-7) pursuant to Regulation (EC) No 1829/2003 of the European Parliament and of the Council⁽¹⁷⁾ is amended as follows.

(2) In Article 7 (authorisation holders)—

(a) in the heading, for “holders” substitute “holder”; and

(b) for paragraphs 1 and 2 substitute—

“The authorisation holder is Corteva Agriscience LLC, United States of America, represented in Great Britain by Corteva Agriscience UK Limited.”.

(3) In Article 9 (addressee), for paragraphs 1 and 2 substitute—

“This Decision is addressed to Corteva Agriscience LLC, 9330 Zionsville Road, Indianapolis, Indiana, 46268-1054, United States of America, represented in Great Britain by Corteva Agriscience UK Limited, Cpc2 Capital Park, Fulbourn, Cambridge, England, CB21 5XE.”.

(4) In the Annex, for point (a) (applicants and authorisation holders) substitute—

“(a) **Authorisation holder:**

(1) The authorisation holder is Corteva Agriscience LLC, 9330 Zionsville Road, Indianapolis, Indiana, 46268-1054, United States of America.

(2) The authorisation holder is represented in Great Britain by Corteva Agriscience UK Limited, Cpc2 Capital Park, Fulbourn, Cambridge, England, CB21 5XE.”.

Amendment of Commission Implementing Decision (EU) 2018/1110

17.—(1) Commission Implementing Decision (EU) 2018/1110 authorising the placing on the market of products containing, consisting of, or produced from genetically modified maize 1507 × 59122 × MON 810 × NK603, and genetically modified maize combining two or three of the single events 1507, 59122, MON 810 and NK603, and repealing Decisions 2009/815/EC, 2010/428/EU and 2010/432/EU⁽¹⁸⁾ is amended as follows.

(2) In Article 7 (authorisation holder), for the text substitute—

“The authorisation holder is Corteva Agriscience LLC, United States of America, represented in Great Britain by Corteva Agriscience UK Limited.”.

(3) In Article 10 (addressee), for the text substitute—

“This Decision is addressed to Corteva Agriscience LLC, 9330 Zionsville Road, Indianapolis, Indiana, 46268-1054, United States of America, represented in Great Britain by Corteva Agriscience UK Limited, Cpc2 Capital Park, Fulbourn, Cambridge, England, CB21 5XE.”.

(4) In the Annex, for point (a) (applicant and authorisation holder) for the text substitute—

“(a) **Authorisation holder:**

⁽¹⁷⁾ EUDN 2018/1109, amended by S.I. 2019/705.

⁽¹⁸⁾ EUDN 2018/1110, amended by S.I. 2019/705.

(1) The authorisation holder is Corteva Agriscience LLC, 9330 Zionsville Road, Indianapolis, Indiana, 46268-1054, United States of America.

(2) The authorisation holder is represented in Great Britain by Corteva Agriscience UK Limited, Cpc2 Capital Park, Fulbourn, Cambridge, England, CB21 5XE.”.

Amendment of Commission Implementing Decision (EU) 2019/1304

18.—(1) Commission Implementing Decision (EU) 2019/1304 authorising the placing on the market of products containing, consisting of or produced from genetically modified maize 4114 (DP-ØØ4114-3), pursuant to Regulation (EC) No 1829/2003 of the European Parliament and of the Council(**19**) is amended as follows.

(2) In Article 7 (authorisation holder), for the text substitute—

“The authorisation holder is Corteva Agriscience LLC, United States of America, represented in Great Britain by Corteva Agriscience UK Limited.”.

(3) In Article 9 (addressee), for the text substitute—

“This Decision is addressed to Corteva Agriscience LLC, 9330 Zionsville Road, Indianapolis, Indiana, 46268-1054, United States of America, represented in Great Britain by Corteva Agriscience UK Limited, Cpc2 Capital Park, Fulbourn, Cambridge, England, CB21 5XE.”.

(4) In the Annex, for point (a) (applicant and authorisation holder) substitute—

“(a) **Authorisation holder:**

(1) The authorisation holder is Corteva Agriscience LLC, 9330 Zionsville Road, Indianapolis, Indiana, 46268-1054, United States of America.

(2) The authorisation holder is represented in Great Britain by Corteva Agriscience UK Limited, Cpc2 Capital Park, Fulbourn, Cambridge, England, CB21 5XE.”.

Amendment of Commission Implementing Decision (EU) 2019/1306

19.—(1) Commission Implementing Decision (EU) 2019/1306 renewing the authorisation for the placing on the market of products containing, consisting of or produced from genetically modified maize 1507 × NK603 (DAS-Ø15Ø7-1 × MON-ØØ6Ø3-6) pursuant to Regulation (EC) No 1829/2003 of the European Parliament and of the Council(**20**) is amended as follows.

(2) In Article 7 (authorisation holders)—

(a) in the heading, for “holders” substitute “holder”;

(b) for paragraphs 1 and 2 substitute—

“The authorisation holder is Corteva Agriscience LLC, United States of America, represented in Great Britain by Corteva Agriscience UK Limited.”.

(3) In Article 9 (addressees)—

(a) in the heading, for “Addressees” substitute “Addressee”;

(b) for the text substitute—

“This Decision is addressed to Corteva Agriscience LLC, 9330 Zionsville Road, Indianapolis, Indiana, 46268-1054, United States of America, represented in Great Britain by Corteva Agriscience UK Limited, Cpc2 Capital Park, Fulbourn, Cambridge, England, CB21 5XE.”.

(4) In the Annex, for point (a) (applicants and authorisation holders) substitute—

(19) EUDN 2019/1304, amended by S.I. 2019/705.

(20) EUDN 2019/1306, amended by S.I. 2019/705.

“(a) **Authorisation holder:**

(1) The authorisation holder is Corteva Agriscience LLC, 9330 Zionsville Road, Indianapolis, Indiana, 46268-1054, United States of America.

(2) The authorisation holder is represented in Great Britain by Corteva Agriscience UK Limited, Cpc2 Capital Park, Fulbourn, Cambridge, England, CB21 5XE.”.

Amendment of Commission Implementing Decision (EU) 2019/2085

20.—(1) Commission Implementing Decision (EU) 2019/2085 authorising the placing on the market of products containing, consisting of or produced from genetically modified maize MON 89034 × 1507 × NK603 × DAS-40278-9 and sub-combinations MON 89034 × NK603 × DAS-40278-9, 1507 × NK603 × DAS-40278-9 and NK603 × DAS-40278-9 pursuant to Regulation (EC) No 1829/2003 of the European Parliament and of the Council⁽²¹⁾ is amended as follows.

(2) In Article 7 (authorisation holder), for the text substitute—

“The authorisation holder is Corteva Agriscience LLC, United States of America, represented in Great Britain by Corteva Agriscience UK Limited.”.

(3) In Article 9 (addressee), for the text substitute—

“This Decision is addressed to Corteva Agriscience LLC, 9330 Zionsville Road, Indianapolis, Indiana, 46268-1054, United States of America, represented in Great Britain by Corteva Agriscience UK Limited, Cpc2 Capital Park, Fulbourn, Cambridge, England, CB21 5XE.”.

(4) In the Annex, for point (a) (applicant and authorisation holder) substitute—

“(a) **Authorisation holder:**

(1) The authorisation holder is Corteva Agriscience LLC, 9330 Zionsville Road, Indianapolis, Indiana, 46268-1054, United States of America.

(2) The authorisation holder is represented in Great Britain by Corteva Agriscience UK Limited, Cpc2 Capital Park, Fulbourn, Cambridge, England, CB21 5XE.”.

Amendment of Commission Implementing Decision (EU) 2019/2086

21.—(1) Commission Implementing Decision (EU) 2019/2086 authorising the placing on the market of products containing, consisting of or produced from genetically modified maize MON 89034 × 1507 × MON 88017 × 59122 × DAS-40278-9 and genetically modified maize combining two, three or four of the single events MON 89034, 1507, MON 88017, 59122 and DAS-40278-9 pursuant to Regulation (EC) No 1829/2003 of the European Parliament and of the Council⁽²²⁾ is amended as follows.

(2) In Article 7 (authorisation holder), for the text substitute—

“The authorisation holder is Corteva Agriscience LLC, United States of America, represented in Great Britain by Corteva Agriscience UK Limited.”.

(3) In Article 9 (addressee), for the text substitute—

“This Decision is addressed to Corteva Agriscience LLC, 9330 Zionsville Road, Indianapolis, Indiana, 46268-1054, United States of America, represented in Great Britain by Corteva Agriscience UK Limited, Cpc2 Capital Park, Fulbourn, Cambridge, England, CB21 5XE.”.

(4) In the Annex, for point (a) (applicant and authorisation holder) substitute—

“(a) **Authorisation holder:**

⁽²¹⁾ EUDN 2019/2085, amended by [S.I. 2019/705](#).

⁽²²⁾ EUDN 2019/2086, amended by [S.I. 2019/705](#).

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(1) The authorisation holder is Corteva Agriscience LLC, 9330 Zionsville Road, Indianapolis, Indiana, 46268-1054, United States of America.

(2) The authorisation holder is represented in Great Britain by Corteva Agriscience UK Limited, Cpc2 Capital Park, Fulbourn, Cambridge, England, CB21 5XE.”.

PART 4

Revocations

Revocation of Commission Decision 2010/429/EU

22. Commission [Decision 2010/429/EU](#) authorising the placing on the market of products containing, consisting of, or produced from genetically modified maize MON 88017 × MON 810 (MON-88Ø17-3 × MON-ØØ81Ø-6) pursuant to Regulation (EC) No 1829/2003 of the European Parliament and of the Council(**23**) is revoked.

28th February 2023

Neil O'Brien
Parliamentary Under-Secretary of State,
Department of Health and Social Care

(23) EUDN 2010/429, amended by [S.I. 2019/705](#). See Schedule 7 of these Regulations for the renewal of the authorisation contained in EUDN 2010/429.

SCHEDULES

SCHEDULE 1

Regulation 3

Authorisation of the placing on the market of products containing, consisting of, or produced from genetically modified soybean DAS-81419-2 × DAS-44406-6

Genetically modified organism and unique identifier

1. For the purposes of Articles 7(3) and 19(3) of Regulation 1829/2003, the unique identifier DAS-81419-2 × DAS-44406-6 is specified for genetically modified soybean DAS-81419-2 × DAS-44406-6.

Authorisation

2. The following products are authorised for the purposes of Articles 4(2) and 16(2) of Regulation 1829/2003, in accordance with the conditions set out in this Schedule—

- (a) food and food ingredients containing, consisting of, or produced from genetically modified soybean DAS-81419-2 × DAS-44406-6;
- (b) feed containing, consisting of, or produced from genetically modified soybean DAS-81419-2 × DAS-44406-6; and
- (c) products containing or consisting of genetically modified soybean DAS-81419-2 × DAS-44406-6 for uses other than those in sub-paragraphs (a) and (b), with the exception of cultivation.

Labelling

3.—(1) For the purposes of the labelling requirements laid down in Articles 13(1) and 25(2) of Regulation 1829/2003, and in Article 4(6) of Regulation 1830/2003, the ‘name of the organism’ is ‘soybean’.

(2) The words ‘not for cultivation’ must appear on the label of, and in documents accompanying, products containing or consisting of genetically modified soybean DAS-81419-2 × DAS-44406-6, with the exception of food and food ingredients.

Method for detection

4.—(1) For the purposes of Articles 7(3) and 19(3) of Regulation 1829/2003, the methods specified in sub-paragraph (2) are to be used for the detection of genetically modified soybean DAS-81419-2 × DAS-44406-6.

(2) The methods are set out in—

- (a) for DAS-81419-2, the document entitled “Event-specific Method for the Quantification of Soybean DAS-81419-2 by Real-time PCR”, reference “EURL-VL-03/13 VP” and dated 13 March 2015;
- (b) for DAS-44406-6 the document entitled “Event-specific Method for the Quantification of Soybean DAS-44406-6 by Real-time PCR”, reference “EURL-VL-01/12 VP” and dated 17 March 2015.

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(3) The method of DNA extraction for use in the detection methods specified in sub-paragraph (2) is set out in the document entitled “Report on the In-house Validation of a DNA Extraction Method from Soybean Grains and Validated Method”, reference “EURL-VL-11/10XP” and dated 13 May 2014.

(4) For the purposes of Articles 7(3) and 19(3) of Regulation 1829/2003, the following reference materials are accessible via the Joint Research Centre of the European Commission⁽²⁴⁾—

- (a) “ERM®-BF437” (for DAS-81419-2);
- (b) “ERM®-BF436” (for DAS-44406-6).

Monitoring plan for environmental effects

5.—(1) The authorisation holder must ensure that the monitoring plan for environmental effects, which accompanied the application for authorisation of the genetically modified soybean DAS-81419-2 × DAS-44406-6, reference number “RP1133” submitted to the Food Safety Authority⁽²⁵⁾ on 8 June 2021, is implemented.

(2) The authorisation holder must submit to the Food Safety Authority annual reports on the implementation of, and the results of the activities set out in, the monitoring plan in accordance with the format set out in Annex 2 of Decision 2009/770.

Authorisation holder

6.—(1) The name and address of the authorisation holder is Corteva Agriscience LLC, 9330 Zionsville Road, Indianapolis, Indiana, 46268-1054, United States of America.

(2) The authorisation holder is represented in Great Britain by Corteva Agriscience UK Limited, Cpc2 Capital Park, Fulbourn, Cambridge, England, CB21 5XE.

SCHEDULE 2

Regulation 3

Authorisation of the placing on the market of products containing,
consisting of, or produced from genetically modified soybean DAS-81419-2

Genetically modified organism and unique identifier

1. For the purposes of Articles 7(3) and 19(3) of Regulation 1829/2003, the unique identifier DAS-81419-2 is specified for genetically modified soybean DAS-81419-2.

Authorisation

2. The following products are authorised for the purposes of Articles 4(2) and 16(2) of Regulation 1829/2003, in accordance with the conditions set out in this Schedule—

- (a) food and food ingredients containing, consisting of or produced from genetically modified soybean DAS-81419-2;
- (b) feed containing, consisting of, or produced from genetically modified soybean DAS-81419-2; and

⁽²⁴⁾ <https://crm.jrc.ec.europa.eu/>

⁽²⁵⁾ “Food Safety Authority” is defined in Article 2(17) of 1829/2003, as regards England, as the Food Standards Agency.

- (c) products containing or consisting of genetically modified soybean DAS-81419-2 for uses other than those provided for in sub-paragraphs (a) and (b), with the exception of cultivation.

Labelling

3.—(1) For the purposes of the labelling requirements laid down in Articles 13(1) and 25(2) of Regulation 1829/2003, and in Article 4(6) of Regulation 1830/2003, the ‘name of the organism’ is ‘soybean’.

(2) The words ‘not for cultivation’ must appear on the label of, and in documents accompanying, the products containing or consisting of genetically modified soybean DAS-81419-2, with the exception of food and food ingredients.

Method for detection

4.—(1) For the purposes of Articles 7(3) and 19(3) of Regulation 1829/2003, the method specified in sub-paragraph (2) is to be used for the detection of genetically modified soybean DAS-81419-2.

(2) The method is set out in the document entitled “Event-specific Method for the Quantification of Soybean DAS-81419-2 by Real-time PCR”, reference “EURL-VL-03/13 VP” and dated 13 March 2015.

(3) The method of DNA extraction for use in the detection method specified in sub-paragraph (2) is set out in the document entitled “Report on the In-house Validation of a DNA Extraction Method from Soybean Grains and Validated Method”, reference “EURL-VL-11/10XP” and dated 13 May 2014.

(4) For the purposes of Articles 7(3) and 19(3) of Regulation 1829/2003, the reference material “ERM®-BF437” is accessible via the Joint Research Centre of the European Commission.

Monitoring plan for environmental effects

5.—(1) The authorisation holder must ensure that the monitoring plan for environmental effects, which accompanied the application for authorisation of genetically modified soybean DAS-81419-2, reference number “RP1134” submitted to the Food Safety Authority on 8 June 2021, is implemented.

(2) The authorisation holder must submit to the Food Safety Authority annual reports on the implementation of, and the results of the activities set out in, the monitoring plan in accordance with the format set out in Annex 2 of Decision 2009/770.

Authorisation holder

6.—(1) The name and address of the authorisation holder is Corteva Agriscience LLC, 9330 Zionsville Road, Indianapolis, Indiana, 46268-1054, United States of America.

(2) The authorisation holder is represented in Great Britain by Corteva Agriscience UK Limited, Cpc2 Capital Park, Fulbourn, Cambridge, England, CB21 5XE.

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SCHEDULE 3

Regulation 3

Authorisation of the placing on the market of products containing, consisting of, or produced from genetically modified soybean SYHT0H2

Genetically modified organism and unique identifier

1. For the purposes of Articles 7(3) and 19(3) of Regulation 1829/2003, the unique identifier SYN-ØØØH2-5 is specified for genetically modified soybean SYHT0H2.

Authorisation

2. The following products are authorised for the purposes of Articles 4(2) and 16(2) of Regulation 1829/2003, in accordance with the conditions set out in this Schedule—

- (a) food and food ingredients containing, consisting of, or produced from genetically modified soybean SYN-ØØØH2-5;
- (b) feed containing, consisting of, or produced from genetically modified soybean SYN-ØØØH2-5; and
- (c) products containing or consisting of genetically modified soybean SYN-ØØØH2-5 for uses other than those provided for in sub-paragraphs (a) and (b), with the exception of cultivation.

Labelling

3.—(1) For the purposes of the labelling requirements laid down in Articles 13(1) and 25(2) of Regulation 1829/2003, and in Article 4(6) of Regulation 1830/2003, the ‘name of the organism’ is ‘soybean’.

(2) The words ‘not for cultivation’ must appear on the label of, and in documents accompanying, products containing or consisting of genetically modified soybean SYN-ØØØH2-5, with the exception of food and food ingredients.

Method for detection

4.—(1) For the purposes of Articles 7(3) and 19(3) of Regulation 1829/2003, the method specified in sub-paragraph (2) is to be used for the detection of genetically modified soybean SYN-ØØØH2-5.

(2) The method is set out in the document entitled “Event-specific Method for the Quantification of Soybean SYHT0H2 by Real-time PCR”, reference “EURL-VL-04/12VP” and dated 3 August 2016.

(3) The method of DNA extraction for use in the detection method specified in sub-paragraph (2) is set out in the document entitled “Report on the Validation of a DNA Extraction Method for Soybean Seeds”, reference “CRLVL04/07XP” and dated 22 January 2009.

(4) For the purposes of Articles 7(3) and 19(3) of Regulation 1829/2003, the reference material “AOCS 0112-A” is accessible via the American Oil Chemists’ Society (AOCS)(26).

Monitoring plan for environmental effects

5.—(1) The authorisation holder must ensure that the monitoring plan for environmental effects, which accompanied the application for authorisation of genetically modified soybean SYN-

(26) <https://www.aocs.org/crm>

ØØØH2-5, reference number “RP1138” submitted to the Food Safety Authority on 10 June 2021, is implemented.

(2) The authorisation holder must submit to the Food Safety Authority annual reports on the implementation of, and the results of the activities set out in, the monitoring plan in accordance with the format set out in Annex 2 of Decision 2009/770.

Authorisation holder

6.—(1) The name and address of the authorisation holder is Syngenta Crop Protection AG, Rosentalstrasse 67, CH-0458 Basel, Switzerland.

(2) The authorisation holder is represented in Great Britain by Syngenta Limited, Jealott’s Hill International Research Centre, Bracknell, Berkshire, England, RG42 6EY.

SCHEDULE 4

Regulation 3

Authorisation of the placing on the market of products containing, consisting of, or produced from genetically modified maize MON 87427 × MON 87460 × MON 89034 × 1507 × MON 87411 × 59122 and its sub-combinations

Genetically modified organism and unique identifier

1. For the purposes of Articles 7(3) and 19(3) of Regulation 1829/2003, the following unique identifiers are specified for genetically modified maize MON 87427 × MON 87460 × MON 89034 × 1507 × MON 87411 × 59122 and its sub-combinations as follows—

- (a) MON-87427-7 × MON-87460-4 × MON-89034-3 × DAS-Ø15Ø7-1 × MON-87411-9 × DAS-59122-7 for genetically modified maize MON 87427 × MON 87460 × MON 89034 × 1507 × MON 87411 × 59122;
- (b) MON-87427-7 × MON-87460-4 × MON-89034-3 × DAS-Ø15Ø7-1 × MON-87411-9 for genetically modified maize MON 87427 × MON 87460 × MON 89034 × 1507 × MON 87411;
- (c) MON-87427-7 × MON-87460-4 × MON-89034-3 × DAS-Ø15Ø7-1 × DAS-59122-7 for genetically modified maize MON 87427 × MON 87460 × MON 89034 × 1507 × 59122;
- (d) MON-87427-7 × MON-87460-4 × MON-89034-3 × MON-87411-9 × DAS-59122-7 for genetically modified maize MON 87427 × MON 87460 × MON 89034 × MON 87411 × 59122;
- (e) MON-87427-7 × MON-87460-4 × DAS-Ø15Ø7-1 × MON-87411-9 × DAS-59122-7 for genetically modified maize MON 87427 × MON 87460 × 1507 × MON 87411 × 59122;
- (f) MON-87427-7 × MON-89034-3 × DAS-Ø15Ø7-1 × MON-87411-9 × DAS-59122-7 for genetically modified maize MON 87427 × MON 89034 × 1507 × MON 87411 × 59122;
- (g) MON-87460-4 × MON-89034-3 × DAS-Ø15Ø7-1 × MON-87411-9 × DAS-59122-7 for genetically modified maize MON 87460 × MON 89034 × 1507 × MON 87411 × 59122;
- (h) MON-87427-7 × MON-87460-4 × MON-89034-3 × DAS-Ø15Ø7-1 for genetically modified maize MON 87427 × MON 87460 × MON 89034 × 1507;
- (i) MON-87427-7 × MON-87460-4 × MON-89034-3 × MON-87411-9 for genetically modified maize MON 87427 × MON 87460 × MON 89034 × MON 87411;
- (j) MON-87427-7 × MON-87460-4 × MON-89034-3 × DAS-59122-7 for genetically modified maize MON 87427 × MON 87460 × MON 89034 × 59122;

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- (k) MON-87427-7 × MON-87460-4 × DAS-Ø15Ø7-1 × MON-87411-9 for genetically modified maize MON 87427 × MON 87460 × 1507 × MON 87411;
- (l) MON-87427-7 × MON-87460-4 × DAS-Ø15Ø7-1 × DAS-59122-7 for genetically modified maize MON 87427 × MON 87460 × 1507 × 59122;
- (m) MON-87427-7 × MON-87460-4 × MON-87411-9 × DAS-59122-7 for genetically modified maize MON 87427 × MON 87460 × MON 87411 × 59122;
- (n) MON-87427-7 × MON-89Ø34-3 × DAS-Ø15Ø7-1 × MON-87411-9 for genetically modified maize MON 87427 × MON 89034 × 1507 × MON 87411;
- (o) MON-87427-7 × MON-89Ø34-3 × MON-87411-9 × DAS-59122-7 for genetically modified maize MON 87427 × MON 89034 × MON 87411 × 59122;
- (p) MON-87427-7 × DAS-Ø15Ø7-1 × MON-87411-9 × DAS-59122-7 for genetically modified maize MON 87427 × 1507 × MON 87411 × 59122;
- (q) MON-87460-4 × MON-89Ø34-3 × DAS-Ø15Ø7-1 × MON-87411-9 for genetically modified maize MON 87460 × MON 89034 × 1507 × MON 87411;
- (r) MON-87460-4 × MON-89Ø34-3 × DAS-Ø15Ø7-1 × DAS-59122-7 for genetically modified maize MON 87460 × MON 89034 × 1507 × 59122;
- (s) MON-87460-4 × MON-89Ø34-3 × MON-87411-9 × DAS-59122-7 for genetically modified maize MON 87460 × MON 89034 × MON 87411 × 59122;
- (t) MON-87460-4 × DAS-Ø15Ø7-1 × MON-87411-9 × DAS-59122-7 for genetically modified maize MON 87460 × 1507 × MON 87411 × 59122;
- (u) MON-89Ø34-3 × DAS-Ø15Ø7-1 × MON-87411-9 × DAS-59122-7 for genetically modified maize MON 89034 × 1507 × MON 87411 × 59122;
- (v) MON-87427-7 × MON-87460-4 × DAS-Ø15Ø7-1 for genetically modified maize MON 87427 × MON 87460 × 1507;
- (w) MON-87427-7 × MON-87460-4 × MON-87411-9 for genetically modified maize MON 87427 × MON 87460 × MON 87411;
- (x) MON-87427-7 × MON-87460-4 × DAS-59122-7 for genetically modified maize MON 87427 × MON 87460 × 59122;
- (y) MON-87427-7 × DAS-Ø15Ø7-1 × MON-87411-9 for genetically modified maize MON 87427 × 1507 × MON 87411;
- (z) MON-87427-7 × MON-87411-9 × DAS-59122-7 for genetically modified maize MON 87427 × MON 87411 × 59122;
- (aa) MON-87460-4 × MON-89Ø34-3 × DAS-Ø15Ø7-1 for genetically modified maize MON 87460 × MON 89034 × 1507;
- (bb) MON-87460-4 × MON-89Ø34-3 × MON-87411-9 for genetically modified maize MON 87460 × MON 89034 × MON 87411;
- (cc) MON-87460-4 × MON-89Ø34-3 × DAS-59122-7 for genetically modified maize MON 87460 × MON 89034 × 59122;
- (dd) MON-87460-4 × DAS-Ø15Ø7-1 × MON-87411-9 for genetically modified maize MON 87460 × 1507 × MON 87411;
- (ee) MON-87460-4 × DAS-Ø15Ø7-1 × DAS-59122-7 for genetically modified maize MON 87460 × 1507 × 59122;
- (ff) MON-87460-4 × MON-87411-9 × DAS-59122-7 for genetically modified maize MON 87460 × MON 87411 × 59122;

- (gg) MON-89034-3 × DAS-Ø1507-1 × MON-87411-9 for genetically modified maize MON 89034 × 1507 × MON 87411;
- (hh) MON-89034-3 × MON-87411-9 × DAS-59122-7 for genetically modified maize MON 89034 × MON 87411 × 59122;
- (ii) DAS-Ø1507-1 × MON-87411-9 × DAS-59122-7 for genetically modified maize 1507 × MON 87411 × 59122;
- (jj) MON-87460-4 × DAS-Ø1507-1 for genetically modified maize MON 87460 × 1507;
- (kk) MON-87460-4 × MON-87411-9 for genetically modified maize MON 87460 × 87411;
- (ll) MON-87460-4 × DAS-59122-7 for genetically modified maize MON 87460 × 59122;
- (mm) DAS-Ø1507-1 × MON-87411-9 for genetically modified maize 1507 × MON 87411;
- (nn) MON-87411-9 × DAS-59122-7 for genetically modified maize MON 87411 × 59122.

Authorisation

2. The following products are authorised for the purposes of Articles 4(2) and 16(2) of Regulation 1829/2003, in accordance with the conditions set out in this Schedule—

- (a) food and food ingredients containing, consisting of, or produced from genetically modified maize as referred to in paragraph 1;
- (b) feed containing, consisting of, or produced from genetically modified maize as referred to in paragraph 1; and
- (c) products containing or consisting of genetically modified maize as referred to in paragraph 1 for uses other than those in sub-paragraphs (a) and (b), with the exception of cultivation.

Labelling

3.—(1) For the purposes of the labelling requirements laid down in Articles 13(1) and 25(2) of Regulation 1829/2003, and in Article 4(6) of Regulation 1830/2003, the ‘name of the organism’ is ‘maize’.

(2) The words ‘not for cultivation’ must appear on the label of, and in documents accompanying, products containing or consisting of genetically modified maize as referred to in paragraph 1, with the exception of food and food ingredients.

Method for detection

4.—(1) For the purposes of Articles 7(3) and 19(3) of Regulation 1829/2003, the methods specified in sub-paragraph (2) are to be used for the detection of genetically modified maize referred to in paragraph 1.

(2) The methods are set out in—

- (a) for MON-87427-7, the document entitled “Event-specific Method for the Quantification of Maize MON 87427 Using Real-time PCR”, reference “EURL-VL-03/12VP” and dated 9 June 2015;
- (b) for MON-87460-4, the document entitled “Event-specific Method for the Quantification of Maize MON 87460 Using Real-time PCR”, reference “CRLVL04/09VP” and dated 18 January 2012;
- (c) for MON-89034-3, the document entitled “Event-specific Method for the Quantification of Maize Line MON 89034 Using Real-time PCR”, reference “CRLVL06/06VP” and dated 21 October 2008;

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- (d) for DAS-Ø15Ø7-1, the document entitled “Event-specific method for the quantitation of maize line TC1507 using real-time PCR”, reference “CRLVL02/04VP” and dated 9 March 2005;
 - (e) for MON-87411-9, the document entitled “Event-specific Method for the Quantification of maize MON 87411 by Real-time PCR”, reference “EURL-VL-01/15VP” and dated 4 July 2016;
 - (f) for DAS-59122-7, the document entitled “Event-specific method for the quantitation of maize 59122 using real-time PCR”, reference “CRLVL03/05VP – corrected version 1” and dated 8 June 2007.
- (3) The method of DNA extraction for use in the detection methods specified in sub-paragraph (2) is as set out in the document entitled “Report on the Validation of a DNA Extraction Method for Maize Seeds and Grains”, reference “CRLVL16/05XP corrected version 2” and dated 26 July 2017.
- (4) For the purposes of Articles 7(3) and 19(3) of Regulation 1829/2003—
- (a) the following reference materials are accessible via the American Oil Chemists’ Society—
 - (i) “AOCS 0512-A2” (for MON-87427-7);
 - (ii) “AOCS 0709-A2” (for MON-8746Ø-4);
 - (iii) “AOCS 0906-E2” (for MON-89Ø34-3);
 - (iv) “AOCS 0215-B” (for MON-87411-9);
 - (b) the following reference materials are accessible via the Joint Research Centre of the European Commission—
 - (i) “ERM®-BF418” (for DAS-Ø15Ø7-1);
 - (ii) “ERM®-BF424” (for DAS-59122-7).

Monitoring plan for environmental effects

5.—(1) The authorisation holder must ensure that the monitoring plan for environmental effects, which accompanied the application for authorisation of the genetically modified maize referred to in paragraph 1, reference number “RP1180” submitted to the Food Safety Authority on 2 July 2021, is implemented.

(2) The authorisation holder must submit to the Food Safety Authority annual reports on the implementation of, and the results of the activities set out in, the monitoring plan in accordance with the format set out in Annex 2 of Decision 2009/770.

Authorisation holder

6.—(1) The name and address of the authorisation holder is Bayer CropScience LP, 800 N. Lindbergh Boulevard, St. Louis, Missouri 63167, United States of America.

(2) The authorisation holder is represented in Great Britain by Bayer CropScience Limited, 230 Cambridge Science Park, Milton Road, Cambridge, England, CB4 0WB.

SCHEDULE 5

Regulation 3

Authorisation of the placing on the market of products containing, consisting of, or produced from genetically modified maize 1507 × MIR162 × MON 810 × NK603 and its sub-combinations

Genetically modified organism and unique identifier

1. For the purposes of Articles 7(3) and 19(3) of Regulation 1829/2003, the following unique identifiers are specified for genetically modified maize 1507 × MIR 162 × MON 810 × NK 603 and its sub-combinations as follows—

- (a) DAS-Ø15Ø7-1 × SYN-IR162-4 × MON-ØØ81Ø-6 × MON-ØØ6Ø3-6 for genetically modified maize 1507 × MIR162 × MON 810 × NK603;
- (b) DAS-Ø15Ø7-1 × SYN-IR162-4 × MON-ØØ81Ø-6 for genetically modified maize 1507 × MIR162 × MON 810;
- (c) DAS-Ø15Ø7-1 × SYN-IR162-4 × MON-ØØ6Ø3-6 for genetically modified maize 1507 × MIR162 × NK603;
- (d) SYN-IR162-4 × MON-ØØ81Ø-6 × MON-ØØ6Ø3-6 for genetically modified maize MIR162 × MON 810 × NK603;
- (e) SYN-IR162-4 × MON-ØØ81Ø-6 for genetically modified maize MIR162 × MON 810.

Authorisation

2. The following products are authorised for the purposes of Articles 4(2) and 16(2) of Regulation 1829/2003, in accordance with the conditions set out in this Schedule—

- (a) food and food ingredients containing, consisting of, or produced from genetically modified maize as referred to in paragraph 1;
- (b) feed containing, consisting of, or produced from genetically modified maize as referred to in paragraph 1; and
- (c) products containing or consisting of genetically modified maize as referred to in paragraph 1 for uses other than those in sub-paragraphs (a) and (b), with the exception of cultivation.

Labelling

3.—(1) For the purposes of the labelling requirements laid down in Articles 13(1) and 25(2) of Regulation 1829/2003, and in Article 4(6) of Regulation 1830/2003, the ‘name of the organism’ is ‘maize’.

(2) The words ‘not for cultivation’ must appear on the label of, and in documents accompanying, products containing or consisting of genetically modified maize as referred to in paragraph 1, with the exception of food and food ingredients.

Method for detection

4.—(1) For the purposes of Articles 7(3) and 19(3) of Regulation 1829/2003, the methods specified in sub-paragraph (2) are to be used for the detection of genetically modified maize referred to in paragraph 1.

(2) The methods are set out in—

- (a) for DAS-Ø15Ø7-1, the document entitled “Event-specific method for the quantitation of maize line TC1507 using real-time PCR”, “Version B”, reference “JRC 113269” and dated 24 September 2018;

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- (b) for SYN-IR162-4, the document entitled “Event-specific Method for the Quantification of Maize MIR162 Using Real-time PCR”, reference “CRLVL08/08VP” and dated 31 January 2011;
- (c) for MON-ØØ81Ø-6, the document entitled “CRL assessment on the validation of an event specific method for the relative quantitation of maize line MON 810 DNA using real-time PCR as carried out by Federal Institute for Risk Assessment (BfR)”, reference “CRL-VL-25/04VR” and dated 10 March 2006;
- (d) for MON-ØØ6Ø3-6, the document entitled “Event-specific method for the quantitation of maize line NK603 using real-time PCR”, reference “CRLVL27/04VP” and dated 10 January 2005.

(3) The method of DNA extraction for use in the detection methods specified in sub-paragraph (2) is as set out in the document entitled “Report on the In-house Validation of a DNA Extraction Method from Ground Maize Seeds and Validated DNA Extraction Method”, reference “EURL-VL-02/14XP” and dated 10 April 2018.

(4) For the purposes of Articles 7(3) and 19(3) of Regulation 1829/2003—

- (a) the following reference materials are accessible via the Joint Research Centre of the European Commission—
 - (i) ERM®-BF418 (for DAS-Ø15Ø7);
 - (ii) ERM®-BF413 (for MON-ØØ81Ø-6);
 - (iii) ERM®-BF415 (for MON-ØØ6Ø3-6);
- (b) the reference material AOCs 1208-A3 (for SYN-IR162-4) is accessible via the American Oil Chemists’ Society.

Monitoring plan for environmental effects

5.—(1) The authorisation holder must ensure that the monitoring plan for environmental effects, which accompanied the application for authorisation of genetically modified maize referred to in paragraph 1, reference number “RP1184” submitted to the Food Safety Authority on 5 July 2021, is implemented.

(2) The authorisation holder must submit to the Food Safety Authority annual reports on the implementation of, and the results of the activities set out in, the monitoring plan in accordance with the format set out in Annex 2 of Decision 2009/770.

Authorisation holder

6.—(1) The name and address of the authorisation holder is Corteva Agriscience LLC, 9330 Zionsville Road, Indianapolis, Indiana, 46268, 1054, USA.

(2) The authorisation holder is represented in Great Britain by Corteva Agriscience UK Limited, CPC2 Capital Park, Fulbourn, Cambridge, England, CB21 5XE.

SCHEDULE 6

Regulation 3

Authorisation of the placing on the market of products containing, consisting of, or produced from genetically modified cotton GHB614 × T304-40 × GHB119

Genetically modified organism and unique identifier

1. For the purposes of Articles 7(3) and 19(3) of Regulation 1829/2003, the following unique identifier BCS-GH002-5 × BCS-GH004-7 × BCS-GH005-8 is specified for genetically modified cotton GHB614 × T304-40 × GHB119.

Authorisation

2. The following products are authorised for the purposes of Articles 4(2) and 16(2) of Regulation 1829/2003, in accordance with the conditions set out in this Schedule—

- (a) food and food ingredients containing, consisting of, or produced from genetically modified cotton BCS-GH002-5 × BCS-GH004-7 × BCS-GH005-8;
- (b) feed containing, consisting of, or produced from genetically modified cotton BCS-GH002-5 × BCS-GH004-7 × BCS-GH005-8; and
- (c) products containing or consisting of genetically modified cotton BCS-GH002-5 × BCS-GH004-7 × BCS-GH005-8 for uses other than those provided for in sub-paragraphs (a) and (b), with the exception of cultivation.

Labelling

3.—(1) For the purposes of the labelling requirements laid down in Articles 13(1) and 25(2) of Regulation 1829/2003, and in Article 4(6) of Regulation 1830/2003, the ‘name of the organism’ is ‘cotton’.

(2) The words ‘not for cultivation’ must appear on the label of, and in documents accompanying, products containing or consisting of genetically modified cotton BCS-GH002-5 × BCS-GH004-7 × BCS-GH005-8, with the exception of food and food ingredients.

Method for detection

4.—(1) For the purposes of Articles 7(3) and 19(3) of Regulation 1829/2003, the methods specified in sub-paragraph (2) are to be used for the detection of genetically modified cotton BCS-GH002-5 × BCS-GH004-7 × BCS-GH005-8.

(2) The methods are set out in—

- (a) for BCS-GH002-5, the document entitled “Event-specific Method for the Quantification of Cotton Line GHB614 Using Real-time PCR”, reference “CRLVL14/07VP” and dated 5 September 2008;
- (b) for BCS-GH004-7, the document entitled “Event-specific Method for the Quantification of Cotton T304-40 using Real-time PCR”, reference “EURL-VL-05/11VP” and dated 19 December 2012.
- (c) for BCS-GH005-8, the document entitled “Event-specific Method for the Quantification of Cotton GHB119 Using Real-time PCR”, reference EURL VL04/11 VP and dated 11 October 2012.

(3) The method of DNA extraction for use in the detection methods specified in sub-paragraph (2) is set out in the document entitled “Cotton Seeds Sampling and DNA Extraction Report on the

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Validation of DNA Extraction Method from Cotton Seeds”, reference “CRLVL13/04XP” dated 14 March 2007.

- (4) For the purposes of Articles 7(3) and 19(3) of Regulation 1829/2003—
- (a) the reference material “AOCS 1108-A6” (for BCS-GHØØ2-5) is accessible via the American Oil Chemists’ Society;
 - (b) the following reference materials are accessible via the Joint Research Centre of the European Commission—
 - (i) ERM®-BF429 (for BCS-GHØØ4-7);
 - (ii) ERM®-BF428 (for BCS-GHØØ5-8).

Monitoring plan for environmental effects

5.—(1) The authorisation holder must ensure that the monitoring plan for environmental effects, which accompanied the application for authorisation of the genetically modified cotton referred to in paragraph 1, reference number “RP1205” submitted to the Food Safety Authority on 28 July 2021, is implemented.

(2) The authorisation holder must submit to the Food Safety Authority annual reports on the implementation of, and the results of the activities set out in, the monitoring plan in accordance with the format set out in Annex 2 of Decision 2009/770.

Authorisation holder

6.—(1) The name and address of the authorisation holder is BASF Agricultural Solutions Seed US LLC, 100 Park Avenue, Florham Park, New Jersey 07932, United States of America.

(2) The authorisation holder is represented in Great Britain by BASF Plc, 2 Stockport Exchange, Railway Road, Stockport, Cheshire, England, SK1 3GG.

SCHEDULE 7

Regulation 3

Renewal of the authorisation of the placing on the market of products containing, consisting of, or produced from genetically modified maize MON 88017 × MON 810

Genetically modified organism and unique identifier

1. For the purposes of Articles 7(3) and 19(3) of Regulation 1829/2003, the unique identifier MON-88Ø17-3 × MON-ØØ81Ø-6 is specified for genetically modified maize MON 88017 × MON 810.

Authorisation

2. The following products are authorised⁽²⁷⁾ for the purposes of Articles 4(2) and 16(2) of Regulation 1829/2003, in accordance with the conditions set out in this Schedule—

- (a) food and food ingredients containing, consisting of, or produced from genetically modified maize MON-88Ø17-3 × MON-ØØ81Ø-6;
- (b) feed containing, consisting of, or produced from genetically modified maize MON-88Ø17-3 × MON-ØØ81Ø-6; and

⁽²⁷⁾ This authorisation is a renewal of the authorisation previously granted under Commission [Decision 2010/429/EU](#). That instrument is revoked by regulation 22 of these Regulations.

- (c) products containing or consisting of genetically modified maize MON-88Ø17-3 × MON-ØØ81Ø-6 for uses other than those provided for in sub-paragraphs (a) and (b), with the exception of cultivation.

Labelling

3.—(1) For the purposes of the labelling requirements laid down in Articles 13(1) and 25(2) of Regulation 1829/2003, and in Article 4(6) of Regulation 1830/2003, the ‘name of the organism’ is ‘maize’.

(2) The words ‘not for cultivation’ must appear on the label of, and in documents accompanying, the products containing or consisting of the genetically modified maize MON-88Ø17-3 × MON-ØØ81Ø-6, with the exception of food and food ingredients.

Method for detection

4.—(1) For the purposes of Articles 7(3) and 19(3) of Regulation 1829/2003, the methods specified in sub-paragraph (2) are to be used for the detection of genetically modified maize MON-88Ø17-3 × MON-ØØ81Ø-6.

(2) The methods are set out in—

- (a) for MON-88Ø17-3, the document entitled “Event-specific Method for the Quantification of Maize Line MON 88017 Using Real-time PCR”, reference “CRLVL16/05VP corrected version 1” and dated 30 March 2010;
- (b) for MON-ØØ81Ø-6, the document entitled “CRL assessment on the validation of an event specific method for the relative quantitation of maize line MON 810 DNA using real-time PCR as carried out by Federal Institute for Risk Assessment (BfR)”, reference CRL-VL-25/04VR” and dated 10 March 2006.

(3) The method of DNA extraction for use in the detection methods specified in sub-paragraph (2) is as set out in the document entitled “Report on the Validation of a DNA Extraction Method for Maize Seeds and Grains”, reference “CRLVL16/05XP” and dated 13 October 2008.

(4) For the purposes of Articles 7(3) and 19(3) of Regulation 1829/2003—

- (a) the reference material “AOCS 0406-D2” (for MON-88Ø17-3) is accessible via the American Oil Chemists’ Society;
- (b) the reference material “ERM®-BF413” (for MON-ØØ81Ø-6) is accessible via the Joint Research Centre of the European Commission.

Monitoring plan for environmental effects

5.—(1) The authorisation holder must ensure that the monitoring plan for environmental effects, which accompanied the application for the renewal of the authorisation of genetically modified maize MON-88Ø17-3 × MON-ØØ81Ø-6, reference number “RP1179” submitted to the Food Safety Authority on 2 July 2021, is implemented.

(2) The authorisation holder must submit to the Food Safety Authority annual reports on the implementation of, and the results of the activities set out in, the monitoring plan in accordance with the format set out in Annex 2 of Decision 2009/770.

Authorisation holder

6.—(1) The name and address of the authorisation holder is Bayer CropScience LP, 800 N. Lindbergh Boulevard, St. Louis, Missouri 63167, United States of America.

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(2) The authorisation holder is represented in Great Britain by Bayer CropScience Limited, 230 Cambridge Science Park, Milton Road, Cambridge, England, CB4 0WB.

SCHEDULE 8

Regulation 3

Renewal of the authorisation of the placing on the market of products containing or consisting of genetically modified oilseed rape GT73 (other than food)

Genetically modified organism and unique identifier

1. For the purposes of Articles 7(3) and 19(3) of Regulation 1829/2003, the unique identifier MON- 00073-7 is specified for genetically modified oilseed rape GT73.

Authorisation

2. The following products are authorised⁽²⁸⁾ for the purposes of Articles 4(2) and 16(2) of Regulation 1829/2003, in accordance with the conditions set out in this Schedule—

- (a) feed containing or consisting of genetically modified oilseed rape MON- 00073-7; and
- (b) products containing or consisting of genetically modified oilseed rape MON- 00073-7 for uses other than those provided for in sub-paragraph (a) and other than food, with the exception of cultivation.

Labelling

3.—(1) For the purposes of the labelling requirements laid down in Articles 13(1) and 25(2) of Regulation 1829/2003, and in Article 4(6) of Regulation 1830/2003, the ‘name of the organism’ is ‘oilseed rape’.

(2) The words ‘not for cultivation’ must appear on the label of, and in documents accompanying, the products containing or consisting of the genetically modified oilseed rape MON- 00073-7, referred to in paragraph 2.

Method for detection

4.—(1) For the purposes of Articles 7(3) and 19(3) of Regulation 1829/2003, the method specified in sub-paragraph (2) is to be used for the detection of genetically modified oilseed rape MON- 00073-7.

(2) The method is set out in the document entitled “Event-specific Method for the Quantification of Oilseed Rape Line RT73 Using Real-time PCR”, reference “CRLVL26/04VP” and dated 7 February 2007.

(3) The method of DNA extraction for use in the detection method specified in sub-paragraph (2) is set out in the document entitled “Report on the Validation of an Oilseed Rape DNA Extraction Method from Seeds”, “Corrected version 1”, reference “CRLVL26/04XP Version 1” and dated 25 July 2017.

(4) For the purposes of Articles 7(3) and 19(3) of Regulation 1829/2003, the reference material “AOCS 0304-B3” is accessible via the American Oil Chemists’ Society.

⁽²⁸⁾ This authorisation is a renewal of the authorisation previously granted pursuant to Commission Decision 2005/635/EC concerning the placing on the market, in accordance with Directive 2001/18/EC of the European Parliament and of the Council, of an oilseed rape product (*Brassica napus* L., GT73 line) genetically modified for tolerance to the herbicide glyphosate (OJ No L 228, 3.9.2005 p11).

Monitoring plan for environmental effects

5.—(1) The authorisation holder must ensure that the monitoring plan for environmental effects, which accompanied the application for the renewal of the authorisation of genetically modified oilseed rape MON- 00073-7, reference number “RP1263” submitted to the Food Safety Authority on 22 September 2021, is implemented.

(2) The authorisation holder must submit to the Food Safety Authority annual reports on the implementation of, and the results of the activities set out in, the monitoring plan in accordance with the format set out in Annex 2 of Decision 2009/770.

Authorisation holder

6.—(1) The name and address of the authorisation holder is Bayer CropScience LP, 800 N. Lindbergh Boulevard, St. Louis, Missouri 63167, United States of America.

(2) The authorisation holder is represented in Great Britain by Bayer CropScience Limited, 230 Cambridge Science Park, Milton Road, Cambridge, England, CB4 0WB.

EXPLANATORY NOTE

(This note is not part of the Regulations)

These Regulations authorise the placing on the market of specified genetically modified food and feed products in England. They also modify the terms of existing authorisations.

Part 2 and Schedules 1 to 8 contain the authorisations, in relation to England, of products containing, and consisting of, or produced from the specified genetically modified organisms, in accordance with Articles 7(3) and 19(3) of Regulation (EC) No. 1829/2003 of the European Parliament and of the Council on genetically modified food and feed (“Regulation 1829/2003”) as follows—

- (a) Schedule 1 is a new authorisation for genetically modified soybean DAS-81419-2 × DAS-44406-6;
- (b) Schedule 2 is a new authorisation for genetically modified soybean DAS-81419-2;
- (c) Schedule 3 is a new authorisation for genetically modified soybean SYHT0H2;
- (d) Schedule 4 is a new authorisation for genetically modified maize MON 87427 × MON 87460 × MON 89034 × 1507 × MON 87411 × 59122 and its sub-combinations;
- (e) Schedule 5 is a new authorisation for genetically modified maize 1507 × MIR162 × MON 810 × NK603 and its sub-combinations;
- (f) Schedule 6 is a new authorisation for genetically modified cotton GHB614 × T304-40 × GHB119;
- (g) Schedule 7 renews the authorisation for genetically modified maize MON 88017 × MON 810;
- (h) Schedule 8 renews the authorisation for genetically modified oilseed rape GT73. This renewal is limited to products containing or consisting of genetically modified oilseed rape GT73 (other than food). It does not cover products “produced from” that genetically modified organism.

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Authorisations granted by these Regulations are valid for a period of ten years in accordance with Articles 7(4) and 19(4) of Regulation 1829/2003 and will expire at the end of 25th April 2033. This is subject to Articles 11(4) and 23(4) of that Regulation, which provide for an extension of the authorisation period in certain circumstances where an application for renewal has been submitted.

Part 3 amends, in relation to England, 18 retained EU Decisions containing existing authorisations for genetically modified food and feed. The amendments all relate to changes to the names and addresses of the respective authorisation holders, and their representatives in Great Britain.

Part 4 revokes, in relation to England, the retained EU Decision containing the previous authorisation for the products now authorised by Schedule 7.

In each Schedule, paragraph 4 specifies the methods for detection, including sampling, which have been validated for use in relation to the authorised products. The documents referred to have been published at <http://gmo-crl.jrc.ec.europa.eu/statusofdossiers.aspx>. Hard copies of these documents are available for inspection at the Food Standards Agency, Floors 6 and 7, Clive House, 70 Petty France, London SW1H 9EX, at reasonable times, following a written request to that address or to GM.Inquiries@food.gov.uk.

Information on the authorisation of the products will be entered in the register of genetically modified food and feed referred to in Article 28(1) of Regulation 1829/2003.

The authorisations made by this instrument will be notified through the Biosafety Clearing-House to the Parties to the Cartagena Protocol on Biosafety to the Convention on Biological Diversity, pursuant to Article 9(1) and Article 15(1)(e) of Regulation (EC) No. 1946/2003 of the European Parliament and of the Council on transboundary movements of genetically modified organisms.

Further information, including in relation to the register, or the information notified pursuant to the Cartagena Protocol, can be obtained from the Food Standards Agency, Floors 6 and 7, Clive House, 70 Petty France, London SW1H 9EX or by writing to GM.Inquiries@food.gov.uk.

An impact assessment has not been produced for this instrument as no, or no significant, impact on the public, private or voluntary sector is foreseen.