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STATUTORY INSTRUMENTS

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**2023 No. 235**

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

**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-21023-5) pursuant to Regulation [\(EC\) No 1829/2003](#) of the European Parliament and of the Council<sup>(1)</sup> 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 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<sup>(2)</sup> 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—

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(1) EUDN 2011/891, amended by [S.I. 2019/705](#).

(2) EUDN 2012/84, 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 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<sup>(3)</sup> 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:

(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 maize combining three different single GM events (MON89034 × 1507 × MON88017 (MON-89034-3 × DAS-01507-1 × MON-88017-3), MON89034 ×

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

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(4) 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**

- (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-305423-1) pursuant to Regulation (EC) No 1829/2003 of the European Parliament and of the Council(5) 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—

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(4) EUDN 2013/650, amended by S.I. 2019/705.

(5) EUDN 2015/698, 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) 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(6) 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.
- (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(7) 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.

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(6) EUDN 2016/1215, amended by S.I. 2019/705.

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

(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<sup>(8)</sup> 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.”.

#### **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<sup>(9)</sup> 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.”.

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<sup>(8)</sup> EUDN 2017/1212, amended by [S.I. 2019/705](#).

<sup>(9)</sup> EUDN 2017/2448, amended by [S.I. 2019/705](#).

### **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<sup>(10)</sup> 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 DAS-44406-6, pursuant to Regulation (EC) No 1829/2003 of the European Parliament and of the Council on genetically modified food and feed<sup>(11)</sup> 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—

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<sup>(10)</sup> EUDN 2017/2449, amended by S.I. 2019/705.

<sup>(11)</sup> EUDN 2017/2450, amended by S.I. 2019/705.

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(**12**) 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:**

(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(**13**) 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.”.

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

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

(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](#)(**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 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:**

(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(**15**) 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:**

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(14) EUDN 2018/1110, amended by [S.I. 2019/705](#).

(15) EUDN 2019/1304, 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) 2019/1306**

**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(16) 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—

“(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(17) 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.”.

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

(17) EUDN 2019/2085, amended by S.I. 2019/705.

(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<sup>(18)</sup> 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.”.

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<sup>(18)</sup> EUDN 2019/2086, amended by S.I. 2019/705.