

Official Journal of the European Union

L 118



English edition

Legislation

Volume 64

7 April 2021

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⁽¹⁾ Text with EEA relevance.

EN

Acts whose titles are printed in light type are those relating to day-to-day management of agricultural matters, and are generally valid for a limited period.

The titles of all other acts are printed in bold type and preceded by an asterisk.

II

(Non-legislative acts)

REGULATIONS

COMMISSION IMPLEMENTING REGULATION (EU) 2021/566

of 30 March 2021

amending Implementing Regulation (EU) No 540/2011 as regards the extension of the approval periods of the active substances abamectin, *Bacillus subtilis* (Cohn 1872) strain QST 713, *Bacillus thuringiensis* subsp. Aizawai strains ABTS-1857 and GC-91, *Bacillus thuringiensis* subsp. Israeliensis (serotype H-14) strain AM65-52, *Bacillus thuringiensis* subsp. Kurstaki strains ABTS 351, PB 54, SA 11, SA12 and EG 2348, *Beauveria bassiana* strains ATCC 74040 and GHA, clodinafop, clopyralid, *Cydia pomonella* Granulovirus (CpGV), cyprodinil, dichlorprop-P, fenpyroximate, fosetyl, mepanipyrim, *Metarhizium anisopliae* (var. *anisopliae*) strain BIPESCO 5/F52, metconazole, metrafenone, pirimicarb, *Pseudomonas chlororaphis* strain MA342, pyrimethanil, *Pythium oligandrum* M1, rimsulfuron, spinosad, *Streptomyces* K61 (formerly '*S. griseoviridis*'), *Trichoderma asperellum* (formerly '*T. harzianum*') strains ICC012, T25 and TV1, *Trichoderma atroviride* (formerly '*T. harzianum*') strain T11, *Trichoderma gamsii* (formerly '*T. viride*') strain ICC080, *Trichoderma harzianum* strains T-22 and ITEM 908, triclopyr, trinexapac, triticonazole and ziram

(Text with EEA relevance)

THE EUROPEAN COMMISSION,

Having regard to the Treaty on the Functioning of the European Union,

Having regard to Regulation (EC) No 1107/2009 of the European Parliament and of the Council of 21 October 2009 concerning the placing of plant protection products on the market and repealing Council Directives 79/117/EEC and 91/414/EEC ⁽¹⁾, and in particular the first paragraph of Article 17 thereof,

Whereas:

- (1) Part A of the Annex to Commission Implementing Regulation (EU) No 540/2011 ⁽²⁾ sets out the active substances deemed to have been approved under Regulation (EC) No 1107/2009.
- (2) Commission Implementing Regulation (EU) 2020/421 ⁽³⁾ extended the approval period of the active substances abamectin, *Bacillus subtilis* (Cohn 1872) strain QST 713, *Bacillus thuringiensis* subsp. Aizawai strains ABTS-1857 and GC-91, *Bacillus thuringiensis* subsp. Israeliensis (serotype H-14) strain AM65-52, *Bacillus thuringiensis* subsp. Kurstaki

⁽¹⁾ OJ L 309, 24.11.2009, p. 1.

⁽²⁾ Commission Implementing Regulation (EU) No 540/2011 of 25 May 2011 implementing Regulation (EC) No 1107/2009 of the European Parliament and of the Council as regards the list of approved active substances (OJ L 153, 11.6.2011, p. 1).

⁽³⁾ Commission Implementing Regulation (EU) 2020/421 of 18 March 2020 amending Implementing Regulation (EU) No 540/2011 as regards the extension of the approval periods of the active substances abamectin, *Bacillus subtilis* (Cohn 1872) strain QST 713, *Bacillus thuringiensis* subsp. Aizawai strains ABTS-1857 and GC-91, *Bacillus thuringiensis* subsp. Israeliensis (serotype H-14) strain AM65-52, *Bacillus thuringiensis* subsp. Kurstaki strains ABTS 351, PB 54, SA 11, SA12 and EG 2348, *Beauveria bassiana* strains ATCC 74040 and GHA, clodinafop, clopyralid, *Cydia pomonella* Granulovirus (CpGV), cyprodinil, dichlorprop-P, fenpyroximate, fosetyl, *Lecanicillium muscarium* (formerly '*Verticillium lecanii*') strain Ve6, mepanipyrim, *Metarhizium anisopliae* (var. *anisopliae*) strain BIPESCO 5/F52, metconazole, metrafenone, *Phlebiopsis gigantea* strains FOC PG 410.3, VRA 1835 and VRA 1984, pirimicarb, *Pseudomonas chlororaphis* strain MA342, pyrimethanil, *Pythium oligandrum* M1, rimsulfuron, spinosad, *Streptomyces* K61 (formerly '*S. griseoviridis*'), *Trichoderma asperellum* (formerly '*T. harzianum*') strains ICC012, T25 and TV1, *Trichoderma atroviride* (formerly '*T. harzianum*') strains IMI 206040 and T11, *Trichoderma gamsii* (formerly '*T. viride*') strain ICC080, *Trichoderma harzianum* strains T-22 and ITEM 908, triclopyr, trinexapac, triticonazole and ziram (OJ L 84, 20.3.2020, p. 7).

strains ABTS 351, PB 54, SA 11, SA12 and EG 2348, *Beauveria bassiana* strains ATCC 74040 and GHA, clodinafop, clopyralid, *Cydia pomonella* Granulovirus (CpGV), cyprodinil, dichlorprop-P, fenpyroximate, fosetyl, mepanipyrim, *Metarhizium anisopliae* (var. *anisopliae*) strain BIPESCO 5/F52, metconazole, metrafenone, pirimicarb, *Pseudomonas chlororaphis* strain MA342, pyrimethanil, *Pythium oligandrum* M1, rimsulfuron, spinosad, *Streptomyces* K61 (formerly 'S. griseoviridis'), *Trichoderma asperellum* (formerly 'T. harzianum') strains ICC012, T25 and TV1, *Trichoderma atroviride* (formerly 'T. harzianum') strain T11, *Trichoderma gamsii* (formerly 'T. viride') strain ICC080, *Trichoderma harzianum* strains T-22 and ITEM 908, triclopyr, trinexapac, triticonazole and ziram until 30 April 2021.

- (3) Applications for the renewal of the approval of those substances were submitted in accordance with Commission Implementing Regulation (EU) No 844/2012 ⁽⁴⁾.
- (4) Due to the fact that the assessment of those substances has been delayed for reasons beyond the control of the applicants, the approvals of those active substances are likely to expire before a decision has been taken on their renewal. It is therefore necessary to extend their approval periods.
- (5) In addition, an extension of the approval period is required for the active substances cyprodinil, dichlorprop-P, fosetyl, mepanipyrim, metconazole, metrafenone, pirimicarb, pyrimethanil, spinosad, triclopyr, trinexapac, triticonazole and ziram also to provide the time necessary to carry out an assessment relating to endocrine disrupting properties of those active substances in accordance with the procedure set out in Articles 13 and 14 of Implementing Regulation (EU) No 844/2012.
- (6) As regards cases where the Commission is to adopt a Regulation providing that the approval of an active substance referred to in the Annex to this Regulation is not renewed because the approval criteria are not satisfied, the Commission is to set the expiry date at the same date as before this Regulation or at the date of the entry into force of the Regulation providing that the approval of the active substance is not renewed, whichever date is later. As regards cases where the Commission is to adopt a Regulation providing for the renewal of an active substance referred to in the Annex to this Regulation, the Commission will endeavour to set, as appropriate under the circumstances, the earliest possible application date.
- (7) Implementing Regulation (EU) No 540/2011 should therefore be amended accordingly.
- (8) The measures provided for in this Regulation are in accordance with the opinion of the Standing Committee on Plants, Animals, Food and Feed,

HAS ADOPTED THIS REGULATION:

Article 1

The Annex to Implementing Regulation (EU) No 540/2011 is amended in accordance with the Annex to this Regulation.

Article 2

This Regulation shall enter into force on the twentieth day following that of its publication in the *Official Journal of the European Union*.

⁽⁴⁾ Commission Implementing Regulation (EU) No 844/2012 of 18 September 2012 setting out the provisions necessary for the implementation of the renewal procedure for active substances, as provided for in Regulation (EC) No 1107/2009 of the European Parliament and of the Council concerning the placing of plant protection products on the market (OJ L 252, 19.9.2012, p. 26).

This Regulation shall be binding in its entirety and directly applicable in all Member States.

Done at Brussels, 30 March 2021.

For the Commission
The President
Ursula VON DER LEYEN

ANNEX

Part A of the Annex to Implementing Regulation (EU) No 540/2011 is amended as follows:

- (1) in the sixth column, expiration of approval, of row 74, Ziram, the date is replaced by '30 April 2022';
- (2) in the sixth column, expiration of approval, of row 89, *Pseudomonas chlororaphis* strain MA 342, the date is replaced by '30 April 2022';
- (3) in the sixth column, expiration of approval, of row 90, Mepanipyrim, the date is replaced by '30 April 2022';
- (4) in the sixth column, expiration of approval, of row 123, Clodinafop, the date is replaced by '30 April 2022';
- (5) in the sixth column, expiration of approval, of row 124, Pirimicarb, the date is replaced by '30 April 2022';
- (6) in the sixth column, expiration of approval, of row 125, Rimsulfuron, the date is replaced by '30 April 2022';
- (7) in the sixth column, expiration of approval, of row 127, Triticonazole, the date is replaced by '30 April 2022';
- (8) in the sixth column, expiration of approval, of row 129, Clopyralid, the date is replaced by '30 April 2022';
- (9) in the sixth column, expiration of approval, of row 130, Cyprodinil, the date is replaced by '30 April 2022';
- (10) in the sixth column, expiration of approval, of row 131, Fosetyl, the date is replaced by '30 April 2022';
- (11) in the sixth column, expiration of approval, of row 132, Trinexapac, the date is replaced by '30 April 2022';
- (12) in the sixth column, expiration of approval, of row 133, Dichlorprop-P, the date is replaced by '30 April 2022';
- (13) in the sixth column, expiration of approval, of row 134, Metconazole, the date is replaced by '30 April 2022';
- (14) in the sixth column, expiration of approval, of row 135, Pyrimethanil, the date is replaced by '30 April 2022';
- (15) in the sixth column, expiration of approval, of row 136, Triclopyr, the date is replaced by '30 April 2022';
- (16) in the sixth column, expiration of approval, of row 137, Metrafenone, the date is replaced by '30 April 2022';
- (17) in the sixth column, expiration of approval, of row 138, *Bacillus subtilis* (Cohn 1872) strain QST 713, the date is replaced by '30 April 2022';
- (18) in the sixth column, expiration of approval, of row 139, Spinosad, the date is replaced by '30 April 2022';
- (19) in the sixth column, expiration of approval, of row 193, *Bacillus thuringiensis* subsp. Aizawai strain ABTS-1857 and strain GC-91, the date is replaced by '30 April 2022';
- (20) in the sixth column, expiration of approval, of row 194, *Bacillus thuringiensis* subsp. israeliensis (serotype H-14) strain AM65-52, the date is replaced by '30 April 2022';
- (21) in the sixth column, expiration of approval, of row 195, *Bacillus thuringiensis* subsp. kurstaki strain ABTS 351, strain PB 54, strain SA 11, strain SA 12, strain EG 2348, the date is replaced by '30 April 2022';
- (22) in the sixth column, expiration of approval, of row 197, *Beauveria bassiana* strain ATCC 74040, strain GHA, the date is replaced by '30 April 2022';
- (23) in the sixth column, expiration of approval, of row 198, *Cydia pomonella* Granulovirus (CpGV), the date is replaced by '30 April 2022';

- (24) in the sixth column, expiration of approval, of row 200, *Metarhizium anisopliae* var. *anisopliae* (formerly *Metarhizium anisopliae*) strain BIPESCO 5/ F52, the date is replaced by '30 April 2022';
 - (25) in the sixth column, expiration of approval, of row 202, *Pythium oligandrum* strain M1, the date is replaced by '30 April 2022';
 - (26) in the sixth column, expiration of approval, of row 203, *Streptomyces* K61 (formerly *S. griseoviridis*) strain K61, the date is replaced by '30 April 2022';
 - (27) in the sixth column, expiration of approval, of row 204, *Trichoderma atroviride* (formerly *T. harzianum*) strain T11, the date is replaced by '30 April 2022';
 - (28) in the sixth column, expiration of approval, of row 206, *Trichoderma harzianum* Rifai strain T-22, stain ITEM 908, the date is replaced by '30 April 2022';
 - (29) in the sixth column, expiration of approval, of row 207, *Trichoderma asperellum* (formerly *T. harzianum*) strain ICC012, strain T25, strain TV1, the date is replaced by '30 April 2022';
 - (30) in the sixth column, expiration of approval, of row 208, *Trichoderma gamsii* (formerly *T. viride*) strain ICC080, the date is replaced by '30 April 2022';
 - (31) in the sixth column, expiration of approval, of row 210, Abamectin, the date is replaced by '30 April 2022';
 - (32) in the sixth column, expiration of approval, of row 213, Fenpyroximate, the date is replaced by '30 April 2022'.
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COMMISSION IMPLEMENTING REGULATION (EU) 2021/567**of 6 April 2021****approving the low-risk active substance aqueous extract from the germinated seeds of sweet *Lupinus albus* in accordance with Regulation (EC) No 1107/2009 of the European Parliament and of the Council concerning the placing of plant protection products on the market, and amending Commission Implementing Regulation (EU) No 540/2011****(Text with EEA relevance)**

THE EUROPEAN COMMISSION,

Having regard to the Treaty on the Functioning of the European Union,

Having regard to Regulation (EC) No 1107/2009 of the European Parliament and of the Council of 21 October 2009 concerning the placing of plant protection products on the market and repealing Council Directives 79/117/EEC and 91/414/EEC ⁽¹⁾, and in particular Article 13(2) in conjunction with Article 22(1) thereof,

Whereas:

- (1) In accordance with Article 7(1) of Regulation (EC) No 1107/2009, CEV SA submitted to the Netherlands on 7 June 2016 an application for the approval of the active substance aqueous extract from the germinated seeds of sweet *Lupinus albus*.
- (2) In accordance with Article 9(3) of Regulation (EC) No 1107/2009, the Netherlands, as rapporteur Member State, notified the applicant, the other Member States, the Commission and the European Food Safety Authority ('the Authority') of the admissibility of the application on 18 January 2017.
- (3) On 1 April 2019, the rapporteur Member State submitted a draft assessment report to the Commission with a copy to the Authority, assessing whether that active substance can be expected to meet the approval criteria provided for in Article 4 of Regulation (EC) No 1107/2009.
- (4) The Authority complied with Article 12(1) of Regulation (EC) No 1107/2009. In accordance with Article 12(3) of that Regulation, it requested that the applicant supplies additional information to the Member States, the Commission and the Authority. The assessment of the additional information by the rapporteur Member State was submitted to the Authority in the format of an updated draft assessment report on 3 March 2020.
- (5) On 19 June 2020, the Authority communicated to the applicant, the Member States and the Commission its conclusion ⁽²⁾ on whether the active substance aqueous extract from the germinated seeds of sweet *Lupinus albus* can be expected to meet the approval criteria provided for in Article 4 of Regulation (EC) No 1107/2009. The Authority made its conclusion available to the public.
- (6) On 23 October 2020, the Commission presented to the Standing Committee on Plants, Animals, Food and Feed a review report and a draft regulation regarding aqueous extract from the germinated seeds of sweet *Lupinus albus*.
- (7) The applicant was given the possibility to submit comments on the review report.
- (8) It has been established with respect to one or more representative uses of at least one plant protection product containing the active substance, and in particular the uses which were examined and detailed in the review report, that the approval criteria provided for in Article 4 of Regulation (EC) No 1107/2009 are satisfied.

⁽¹⁾ OJ L 309, 24.11.2009, p. 1.

⁽²⁾ EFSA (European Food Safety Authority), 2020. Conclusion on the peer review of the pesticide risk assessment of the active substance Aqueous extract from the germinated seeds of sweet *Lupinus albus*; *EFSA Journal* 2020;18(7):6190, 45 pp.; <https://doi.org/10.2903/j.efsa.2020.6190>. Available online: www.efsa.europa.eu.

- (9) The Commission further considers that aqueous extract from the germinated seeds of sweet *Lupinus albus* is a low-risk active substance pursuant to Article 22 of Regulation (EC) No 1107/2009. Aqueous extract from the germinated seeds of sweet *Lupinus albus* is not a substance of concern and it fulfils the conditions set out in point 5.1 of Annex II to Regulation (EC) No 1107/2009.
- (10) It is therefore appropriate to approve aqueous extract from the germinated seeds of sweet *Lupinus albus*.
- (11) In accordance with Article 13(2) of Regulation (EC) No 1107/2009 in conjunction with Article 6 thereof and in the light of current scientific and technical knowledge, it is necessary to include certain conditions and restrictions. It is in particular appropriate to require further confirmatory information.
- (12) In accordance with Article 13(4) of Regulation (EC) No 1107/2009, the Annex to Commission Implementing Regulation (EU) No 540/2011 ⁽³⁾ should be amended accordingly.
- (13) The measures provided for in this Regulation are in accordance with the opinion of the Standing Committee on Plants, Animals, Food and Feed,

HAS ADOPTED THIS REGULATION:

Article 1

Approval of the active substance

The active substance aqueous extract from the germinated seeds of sweet *Lupinus albus*, as specified in Annex I, is approved subject to the conditions laid down in that Annex.

Article 2

Amendments to Implementing Regulation (EU) No 540/2011

The Annex to Implementing Regulation (EU) No 540/2011 is amended in accordance with Annex II to this Regulation.

Article 3

Entry into force

This Regulation shall enter into force on the twentieth day following that of its publication in the *Official Journal of the European Union*.

This Regulation shall be binding in its entirety and directly applicable in all Member States.

Done at Brussels, 6 April 2021.

For the Commission
The President
Ursula VON DER LEYEN

⁽³⁾ Commission Implementing Regulation (EU) No 540/2011 of 25 May 2011 implementing Regulation (EC) No 1107/2009 of the European Parliament and of the Council as regards the list of approved active substances (OJ L 153, 11.6.2011, p. 1).

Common Name, Identification Numbers	IUPAC Name	Purity ⁽¹⁾	Date of approval	Expiration of approval	Specific provisions
Aqueous extract from the germinated seeds of sweet <i>Lupinus albus</i> CAS No: Not available for the extract BLAD protein: 1219521-95-5 CIPAC No: Not allocated	Not applicable	The minimum purity is not relevant for the extract. BLAD protein content: 195 – 210 g/kg. The following relevant impurities (of toxicological, ecotoxicological and/or environmental concern) in the active substance as manufactured were identified: Total quinolizidine alkaloids (QA): (lupanine, 13α-OH-lupanine, 13α-angeloyloxylupanine, lupinine, albine, angustofoline, 13α-tigloyloxylupanine, α-isolupanine, tetrahydrohombifoline, multiflorine, sparteine) Maximum content: provisionally set at 0,05 g/kg	27 April 2021	27 April 2036	For the implementation of the uniform principles as referred to in Article 29(6) of Regulation (EC) No 1107/2009, the conclusions of the review report on aqueous extract from the germinated seeds of sweet <i>Lupinus albus</i>, and in particular Appendices I and II thereto, shall be taken into account. In this overall assessment, Member States shall pay particular attention to the necessary labelling instructions concerning the measures to address foaming and stability of dilutions of the formulation. The applicant shall submit to the Commission, the Member States and the Authority confirmatory information as regards: 1. the technical specification of the active substance as manufactured (based on commercial scale production) and the compliance of the toxicity batches with the confirmed technical specification; and 2. in particular, the maximum content of the quinolizidine alkaloids (lupanine, 13α-OH-lupanine, 13α-angeloyloxylupanine, lupinine, albine, angustofoline, 13α-tigloyloxylupanine, α-isolupanine, tetrahydrohombifoline, multiflorine, sparteine). The applicant shall submit the information referred in points 1 and 2 by 27 October 2021.

⁽¹⁾ Further details on identity and specification of active substance are provided in the review report.

ANNEX II

In Part D of the Annex to Implementing Regulation (EU) No 540/2011, the following entry is added:

No	Common Name, Identification Numbers	IUPAC Name	Purity ⁽¹⁾	Date of approval	Expiration of approval	Specific provisions
28	Aqueous extract from the germinated seeds of sweet <i>Lupinus albus</i> CAS No: Not available for the extract BLAD protein: 1219521-95-5 CIPAC No: Not allocated	Not applicable	The minimum purity is not relevant for the extract. BLAD protein content: 195 – 210 g/kg. The following relevant impurities (of toxicological, ecotoxicological and/or environmental concern) in the active substance as manufactured were identified: Total quinolizidine alkaloids (QA): (lupanine, 13α-OH-lupanine, 13α-angeloyloxylupanine, lupinine, albine, angustofoline, 13α-tigloyloxylupanine, α-isolupanine, tetrahydrohombifoline, multiflorine, sparteine) Maximum content: provisionally set at 0,05 g/kg	27 April 2021	27 April 2036	For the implementation of the uniform principles as referred to in Article 29(6) of Regulation (EC) No 1107/2009, the conclusions of the review report on aqueous extract from the germinated seeds of sweet <i>Lupinus albus</i>, and in particular Appendices I and II thereto, shall be taken into account. In this overall assessment, Member States shall pay particular attention to the necessary labelling instructions concerning the measures to address foaming and stability of dilutions of the formulation. The applicant shall submit to the Commission, the Member States and the Authority confirmatory information as regards: 1. the technical specification of the active substance as manufactured (based on commercial scale production) and the compliance of the toxicity batches with the confirmed technical specification; and 2. in particular on the maximum content of the quinolizidine alkaloids (lupanine, 13α-OH-lupanine, 13α-angeloyloxylupanine, lupinine, albine, angustofoline, 13α-tigloyloxylupanine, α-isolupanine, tetrahydrohombifoline, multiflorine, sparteine). The applicant shall submit the information referred in points 1 and 2 by 27 October 2021.'

⁽¹⁾ Further details on identity and specification of active substance are provided in the review report.

COMMISSION IMPLEMENTING REGULATION (EU) 2021/568**of 6 April 2021****amending Annex I to Regulation (EC) No 798/2008 as regards the entry for the United Kingdom in the list of third countries, territories, zones or compartments from which certain poultry commodities may be imported into and transit through the Union in relation to highly pathogenic avian influenza****(Text with EEA relevance)**

THE EUROPEAN COMMISSION,

Having regard to the Treaty on the Functioning of the European Union,

Having regard to Council Directive 2002/99/EC of 16 December 2002 laying down the animal health rules governing the production, processing, distribution and introduction of products of animal origin for human consumption ⁽¹⁾, and in particular the introductory phrase of Article 8, the first subparagraph of paragraph 1 of Article 8, paragraph 4 of Article 8 and Article 9(4) thereof,

Having regard to Council Directive 2009/158/EC of 30 November 2009 on animal health conditions governing intra-Community trade in, and imports from third countries of, poultry and hatching eggs ⁽²⁾, and in particular Articles 23(1), 24(2) and 25(2) thereof,

Whereas:

- (1) Commission Regulation (EC) No 798/2008 ⁽³⁾ lays down veterinary certification requirements for imports into and transit, including storage during transit, through the Union of poultry and poultry products ('the commodities'). It provides that the commodities are only to be imported into and transit through the Union from the third countries, territories, zones or compartments listed in columns 1 and 3 of the table in Part 1 of Annex I thereto.
- (2) Regulation (EC) No 798/2008 also lays down the conditions for a third country, territory, zone or compartment to be considered as free from highly pathogenic avian influenza (HPAI).
- (3) In accordance with the Agreement on the withdrawal of the United Kingdom of Great Britain and Northern Ireland from the European Union and the European Atomic Energy Community (Withdrawal Agreement), and in particular Article 5(4) of the Protocol on Ireland/Northern Ireland in conjunction with Annex 2 to that Protocol, Directives 2002/99/EC and 2009/158/EC, as well as the Commission acts based on them, apply to and in the United Kingdom in respect of Northern Ireland after the end of the transition period provided for in the Withdrawal Agreement.
- (4) Therefore, the United Kingdom, excluding Northern Ireland, is listed in the table in Part 1 of Annex I to Regulation (EC) No 798/2008 as a third country from which imports into and transit through the Union of certain poultry commodities are authorised from certain parts of its territory depending on the presence of HPAI. That regionalisation of the United Kingdom is set out in Part 1 of Annex I to Regulation (EC) No 798/2008, as amended by Commission Implementing Regulation (EU) 2021/256 ⁽⁴⁾.
- (5) On 29 March 2021, the United Kingdom confirmed the presence of HPAI of subtype H5N8 in a poultry holding in East Staffordshire.

⁽¹⁾ OJ L 18, 23.1.2003, p. 11.

⁽²⁾ OJ L 343, 22.12.2009, p. 74.

⁽³⁾ Commission Regulation (EC) No 798/2008 of 8 August 2008 laying down a list of third countries, territories, zones or compartments from which poultry and poultry products may be imported into and transit through the Community and the veterinary certification requirements (OJ L 226, 23.8.2008, p. 1).

⁽⁴⁾ Commission Implementing Regulation (EU) 2021/256 of 18 February 2021 amending Annex I to Regulation (EC) No 798/2008 as regards the entry for the United Kingdom in the list of third countries, territories, zones or compartments from which certain poultry commodities may be imported into and transit through the Union in relation to highly pathogenic avian influenza (OJ L 58, 19.2.2021, p. 36).

- (6) The veterinary authorities of the United Kingdom placed a 10 km control zone around the affected holding and implemented a stamping-out policy in order to control the presence of HPAI and limit the spread of that disease. Furthermore, the veterinary authorities of the United Kingdom confirmed that they immediately suspended issuing veterinary certificates for consignments of commodities intended for export to the Union from the whole of the territory of the United Kingdom, excluding Northern Ireland.
- (7) The United Kingdom has submitted information to the Commission on the epidemiological situation on its territory and the measures it has taken to prevent the further spread of HPAI which has now been evaluated by the Commission. On the basis of that evaluation, it is appropriate to place restrictions on the introduction into the Union of commodities from the area the veterinary authorities of the United Kingdom have placed under restrictions due to the current HPAI outbreak in East Staffordshire.
- (8) The entry for the United Kingdom in the table in Part 1 of Annex I to Regulation (EC) No 798/2008 should, therefore, be amended to take account of the current epidemiological situation in that third country.
- (9) Annex I to Regulation (EC) No 798/2008 should therefore be amended accordingly.
- (10) The measures provided for in this Regulation are in accordance with the opinion of the Standing Committee on Plants, Animals, Food and Feed,

HAS ADOPTED THIS REGULATION:

Article 1

Part 1 of Annex I to Regulation (EC) No 798/2008 is amended in accordance with the Annex to this Regulation.

Article 2

This Regulation shall enter into force on the third day following that of its publication in the *Official Journal of the European Union*.

This Regulation shall be binding in its entirety and directly applicable in all Member States.

Done at Brussels, 6 April 2021.

For the Commission
The President
Ursula VON DER LEYEN

ANNEX

In Part 1 of Annex I to Regulation (EC) No 798/2008, the entry for the United Kingdom is replaced by the following:

GB - United Kingdom (*)	GB-0	Whole country	SPF							
			EP, E							
GB-1		The whole country of the United Kingdom, excluding area GB-2	BPP, BPR, DOC, DOR, HEP, HER, SRP, SRA, LT20		N			A		
			WGM							
			POU, RAT		N					
GB-2		The territory of the United Kingdom corresponding to:								
GB-2.1		North Yorkshire County: The area contained within a circle of a radius of 10 km, centered on WGS84 dec. coordinates N54.30 and W1.47	BPP, BPR, DOC, DOR, HEP, HER, SRP, SRA, LT20		N P2	1.1.2021	6.1.2021	A		
			WGM		P2	1.1.2021	6.1.2021			
			POU, RAT		N P2	1.1.2021	6.1.2021			
GB-2.2		North Yorkshire County: The area contained within a circle of a radius of 10 km, centered on WGS84 dec. coordinates N54.29 and W1.45	BPP, BPR, DOC, DOR, HEP, HER, SRP, SRA, LT20		N P2	1.1.2021	8.1.2021	A		
			WGM		P2	1.1.2021	8.1.2021			
			POU, RAT		N P2	1.1.2021	8.1.2021			
GB-2.3		Norfolk County: The area contained within a circle of a radius of 10 km, centered on WGS84 dec. coordinates N52.49 and E0.95	BPP, BPR, DOC, DOR, HEP, HER, SRP, SRA, LT20		N P2	1.1.2021	10.1.2021	A		
			WGM		P2	1.1.2021	10.1.2021			
			POU, RAT		N P2	1.1.2021	10.1.2021			
GB-2.4		Norfolk County: The area contained within a circle of a radius of 10 km, centered on WGS84 dec. coordinates N52.72 and E0.15	BPP, BPR, DOC, DOR, HEP, HER, SRP, SRA, LT20		N P2	1.1.2021	11.1.2021	A		
			WGM		P2	1.1.2021	11.1.2021			
			POU, RAT		N P2	1.1.2021	11.1.2021			

GB-2.5	Derbyshire County: The area contained within a circle of a radius of 10 km, centered on WGS84 dec. coordinates N52.93 and W1.57	BPP, BPR, DOC, DOR, HEP, HER, SRP, SRA, LT20		N P2	1.1.2021	17.1.2021	A		
		WGM		P2	1.1.2021	17.1.2021			
		POU, RAT		N P2	1.1.2021	17.1.2021			
GB-2.6	North Yorkshire County: The area contained within a circle of a radius of 10 km, centered on WGS84 dec. coordinates N54.37 and W2.16	BPP, BPR, DOC, DOR, HEP, HER, SRP, SRA, LT20		N P2	1.1.2021	19.1.2021	A		
		WGM		P2	1.1.2021	19.1.2021			
		POU, RAT		N P2	1.1.2021	19.1.2021			
GB-2.7	Orkney Islands: The area contained within a circle of a radius of 10 km, centered on WGS84 dec. coordinates N59.28 and W2.44	BPP, BPR, DOC, DOR, HEP, HER, SRP, SRA, LT20		N P2	1.1.2021	20.1.2021	A		
		WGM		P2	1.1.2021	20.1.2021			
		POU, RAT		N P2	1.1.2021	20.1.2021			
GB-2.8	Dorset County: The area contained within a circle of a radius of 10 km, centered on WGS84 dec. coordinates N51.06 and W2.27	BPP, BPR, DOC, DOR, HEP, HER, SRP, SRA, LT20		N P2	1.1.2021	20.1.2021	A		
		WGM		P2	1.1.2021	20.1.2021			
		POU, RAT		N P2	1.1.2021	20.1.2021			
GB-2.9	Norfolk County: The area contained within a circle of a radius of 10 km, centered on WGS84 dec. coordinates N52.52 and E0.96	BPP, BPR, DOC, DOR, HEP, HER, SRP, SRA, LT20		N P2	1.1.2021	23.1.2021	A		
		WGM		P2	1.1.2021	23.1.2021			
		POU, RAT		N P2	1.1.2021	23.1.2021			
GB-2.10	Norfolk County: The area contained within a circle of a radius of 10 km, centered on WGS84 dec. coordinates N52.52 and E0.95	BPP, BPR, DOC, DOR, HEP, HER, SRP, SRA, LT20		N P2	1.1.2021	28.1.2021	A		
		WGM		P2	1.1.2021	28.1.2021			
		POU, RAT		N P2	1.1.2021	28.1.2021			
GB-2.11	Norfolk County: The area contained within a circle of a radius of 10.4 km, centered on WGS84 dec. coordinates N52.53 and E0.66	BPP, BPR, DOC, DOR, HEP, HER, SRP, SRA, LT20		N P2	1.1.2021	7.2.2021	A		
		WGM		P2	1.1.2021	7.2.2021			
		POU, RAT		N P2	1.1.2021	7.2.2021			

GB-2.12	Devon County: The area contained within a circle of a radius of 10 km, centred on WGS84 dec. coordinates N50.70 and W3.36	BPP, BPR, DOC, DOR, HEP, HER, SRP, SRA, LT20		N P2	1.1.2021	31.1.2021	A		
		WGM		P2	1.1.2021	31.1.2021			
		POU, RAT		N P2	1.1.2021	31.1.2021			
GB-2.13	Near Amlwch, Isle of Anglesey, Wales: The area contained within a circle of a radius of 10 km, centred on WGS84 dec. coordinates N53.38 and W4.30	BPP, BPR, DOC, DOR, HEP, HER, SRP, SRA, LT20		N P2	27.1.2021		A		
		WGM		P2	27.1.2021				
		POU, RAT		N P2	27.1.2021				
GB-2.14	Near Redcar, Redcar and Cleveland, England: The area contained within a circle of a radius of 10km, centred on WGS84 dec, coordinates N54.57 and W1.07	BPP, BPR, DOC, DOR, HEP, HER, SRP, SRA, LT20		N P2	8.2.2021		A		
		WGM		P2	8.2.2021				
		POU, RAT		N P2	8.2.2021				
GB-2.15	Glenrothes, Fife, Scotland: The area contained within a circle of a radius of 10km, centred on WGS84 dec, coordinates N56.23 and W3.02	BPP, BPR, DOC, DOR, HEP, HER, SRP, SRA, LT20		N P2	12.2.2021		A		
		WGM		P2	12.2.2021				
		POU, RAT		N P2	12.2.2021				
GB-2.16	Staffordshire county, Derbyshire county: The area contained with a circle of a radius of 10km, centred on WGS84 dec, coordinates N52.99 and W1.85	BPP, BPR, DOC, DOR, HEP, HER, SRP, SRA, LT20		N P2	29.3.2021		A		
		WGM		P2	29.3.2021				
		POU, RAT		N P2	29.3.2021				

(*) In accordance with the Agreement on the withdrawal of the United Kingdom of Great Britain and Northern Ireland from the European Union and the European Atomic Energy Community, and in particular Article 5(4) of the Protocol on Ireland/Northern Ireland in conjunction with Annex 2 to that Protocol, for the purposes of this Annex references to the United Kingdom do not include Northern Ireland.*

ISSN 1977-0677 (electronic edition)
ISSN 1725-2555 (paper edition)



Publications Office
of the European Union
L-2985 Luxembourg
LUXEMBOURG

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