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Legislation

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Contents

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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.

EN

([1]) Text with EEA relevance

II

(Non-legislative acts)

REGULATIONS

COMMISSION REGULATION (EU) No 1112/2010

of 1 December 2010

amending Regulation (EC) No 793/2006 laying down certain detailed rules for applying Council Regulation (EC) No 247/2006 laying down specific measures for agriculture in the outermost regions of the Union

THE EUROPEAN COMMISSION,

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

Having regard to Council Regulation (EC) No 247/2006 of 30 January 2006 laying down specific measures for agriculture in the outermost regions in the Union [1], and in particular Article 25 thereof,

Whereas:

(1) Following the adoption of Commission Regulation (EC) No 408/2009 [2] amending Regulation (EC) No 793/2006, and in particular Article 46a thereof, which specifies that reconstituted UHT milk intended for local consumption in Madeira is to incorporate at least 15 % of fresh cow's milk, it has emerged that all the fresh milk produced locally is used by the local cheese industry. To avoid any disruption to the economic balance already established and to ensure that fresh milk produced locally can be processed into high value added products, the obligation to incorporate a minimum rate should be abolished.

(2) Having regard to the amendment to Article 19(4) of Regulation (EC) No 247/2006 by Regulation (EU) No 641/2010 of the European Parliament and of the Council [3], which abolished, with effect from 1 January 2010, the Commission's obligation to determine an incorporation rate for fresh milk produced locally, Article 46a of Commission Regulation (EC) No 793/2006 [4] should also be repealed from that date.

(3) Regulation (EC) No 793/2006 should therefore be amended accordingly.

(4) The measures provided for in this Regulation are in accordance with the opinion of the Management Committee for Direct Payments,

HAS ADOPTED THIS REGULATION:

Article 1

Article 46a of Regulation (EC) No 793/2006 is deleted.

Article 2

This Regulation shall enter into force on 1 January 2010.

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

Done at Brussels, 1 December 2010.

For the Commission
The President
José Manuel BARROSO

[1] OJ L 42, 14.2.2006, p. 1.
[2] OJ L 123, 19.5.2009, p. 62.
[3] OJ L 194, 24.7.2010, p. 23.
[4] OJ L 145, 31.5.2006, p. 1.

COMMISSION REGULATION (EU) No 1113/2010

of 1 December 2010

fixing the coefficients applicable to cereals exported in the form of Scotch whisky for the period 2010/2011

THE EUROPEAN COMMISSION,

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

Having regard to Council Regulation (EC) No 1234/2007 of 22 October 2007 establishing a common organisation of agricultural markets and on specific provisions for certain agricultural products (Single CMO Regulation) [1],

Having regard to Commission Regulation (EC) No 1670/2006 of 10 November 2006 laying down certain detailed rules for the application of Council Regulation (EC) No 1784/2003 as regards the fixing and granting of adjusted refunds in respect of cereals exported in the form of certain spirit drinks [2], and in particular Article 5 thereof,

Whereas:

(1) Article 4(1) of Regulation (EC) No 1670/2006 lays down that the quantities of cereals eligible for the refund are to be the quantities placed under control and distilled, weighted by a coefficient to be fixed annually for each Member State concerned. The coefficient is to express the average ratio between the total quantities exported and the total quantities marketed of the spirit drink concerned, on the basis of the trend noted in those quantities during the number of years corresponding to the average ageing period of the spirit drink in question.

(2) According to the information provided by the United Kingdom in respect of the period 1 January to 31 December 2009, the average ageing period for Scotch whisky in 2009 was 7 years.

(3) The coefficients for the period 1 October 2010 to 30 September 2011 should therefore be fixed accordingly.

(4) Article 10 of Protocol 3 to the Agreement on the European Economic Area excludes the grant of refunds in respect of exports to Liechtenstein, Iceland and Norway. Moreover, the Union has concluded agreements abolishing export refunds with certain third countries. Under the terms of Article 7(2) of Regulation (EC) No 1670/2006, this should be taken into account in calculating the coefficients for 2010/2011.

(5) Commission Regulation (EC) No 1035/2009 of 30 October 2009 fixing the coefficients applicable to cereals exported in the form of Scotch whisky for the period 2009/2010 [3] has exhausted its effects, as it concerns the coefficients applicable for the year 2009/2010. For reasons of legal security and clarity, this Regulation should be repealed,

HAS ADOPTED THIS REGULATION:

Article 1

For the period 1 October 2010 to 30 September 2011, the coefficients provided for in Article 4 of Regulation (EC) No 1670/2006 applying to cereals used in the United Kingdom for manufacturing Scotch whisky shall be as set out in the Annex to this Regulation.

Article 2

Regulation (EC) No 1035/2009 is hereby repealed.

Article 3

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

It shall apply from 1 October 2010 to 30 September 2011.

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

Done at Brussels, 1 December 2010.

For the Commission
The President
José Manuel BARROSO

[1] OJ L 299, 16.11.2007, p. 1.
[2] OJ L 312, 11.11.2006, p. 33.
[3] OJ L 285, 31.10.2009, p. 3.

ANNEX

Coefficients applicable in the United Kingdom

Period of application	Coefficient applicable	
	to malted barley used in the production of malt whisky	to cereals used in the production of grain whisky
From 1 October 2010 to 30 September 2011	0,300	0,255

COMMISSION REGULATION (EU) No 1114/2010

of 1 December 2010

laying down detailed rules for the implementation of Council Regulation (EC) No 2494/95 as regards minimum standards for the quality of HICP weightings and repealing Commission Regulation (EC) No 2454/97

(Text with EEA relevance)

THE EUROPEAN COMMISSION,

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

Having regard to Council Regulation (EC) No 2494/95 ([1]) of 23 October 1995 concerning harmonised indices of consumer prices, and in particular Article 3 thereof,

Whereas:

(1) Harmonised Indices of Consumer Prices (HICP) are harmonised inflation measures required by the Commission and the European Central Bank for the performance of their functions pursuant to Article 140 of the Treaty on the Functioning of the European Union. HICPs are designed to facilitate international comparisons of consumer price inflation. They serve as important indicators for the management of monetary policy.

(2) Article 8(3) of Regulation (EC) No 2494/95 requires that the weightings of the HICP are updated with a frequency sufficient to meet the comparability and reliability requirements. HICPs based on weightings that are updated at different frequencies may fail to meet the comparability and the reliability requirements.

(3) Commission Regulation (EC) No 2454/97 ([2]) of 10 December 1997 laying down detailed rules for the implementation of Council Regulation (EC) No 2494/95 as regards minimum standards for the quality of HICP weightings laid down rules to ensure that HICPs were constructed using weightings which were sufficiently reliable and relevant for the purpose of international comparisons. Those rules should now be modified taking into account developments in the HICP domain. Therefore, the measures set out in this Regulation should replace those in Regulation (EC) No 2454/97, which should be repealed.

(4) Article 9 of Regulation (EC) No 2494/95 requires HICPs to be price indices of the Laspeyres-type. When relative prices of different goods and services change, consumers' expenditure patterns can change to an extent that makes it necessary for the weights of the corresponding expenditure groups, and in particular their underlying quantities, to be updated in order to ensure their relevance.

(5) By virtue of Article 4 of Regulation (EC) No 1749/96 ([3]) of 9 September 1996 on initial implementing measures for Council Regulation (EC) No 2494/95 concerning harmonised indices of consumer prices, the HICP should be compiled to include the price changes of newly significant goods or services and their relative expenditures.

(6) This Regulation should apply without prejudice to the minimum standards for the treatment of insurance weights in accordance with Commission Regulation (EC) No 1617/1999 ([4]) of 23 July 1999 laying down detailed rules for the implementation of Council Regulation (EC) No 2494/95 as regards minimum standards for the treatment of insurance in the HICP.

(7) Weights at the level of COICOP/HICP ([5]) divisions, groups and classes are required not to vary between months during the year unless under the provisions of the Commission Regulation (EC) No 330/2009 ([6]) of 22 April 2009 regarding minimum standards for the treatment of seasonal products in the HICP.

(8) This Regulation should not require Member States to carry out new statistical surveys or to carry out family budget surveys more frequently than once every 5 years, taking into consideration that Member States are required to compile national accounts in accordance with the European System of Accounts (ESA 1995) ([7]) and that the country weights, which are necessary for producing euro area, EU and other HICP aggregates, are based on national accounts data.

([1]) OJ L 257, 27.10.1995, p. 1.
([2]) OJ L 340, 11.12.1997, p. 24.
([3]) OJ L 229, 10.9.1996, p. 3.
([4]) OJ L 192, 24.7.1999, p. 9.
([5]) Classification of Individual Consumption by Purpose adapted to the needs of HICPs.
([6]) OJ L 103, 23.4.2009, p. 6.
([7]) Council Regulation (EC) No 2223/96 of 25 June 1996 on the European system of national and regional accounts in the Community, OJ L 310, 30.11.1996, p. 1.

(9) The principle of cost-effectiveness has been taken into account in accordance with Article 13 of Regulation (EC) No 2494/95.

(10) The European Central Bank has been consulted in accordance with Article 5(3) of Regulation (EC) No 2494/95.

(11) The measures provided for in this Regulation are in accordance with the opinion of the European Statistical System Committee,

HAS ADOPTED THIS REGULATION:

Article 1

Subject matter

The aim of this Regulation is to establish minimum standards for the quality of HICP weightings of the Harmonised Indices of Consumer Prices (HICPs).

Article 2

Definitions

For the purpose of this Regulation, the following definitions shall apply:

(1) the 'weighting reference period' of an HICP means the 12-month period of consumption or expenditure from which the weights are estimated for the compilation of the latest HICP index figures;

(2) 'sub-indices' mean the sub-indices laid down in Commission Regulation (EC) No 2214/96 ([1]) of 20 November 1996 concerning harmonised indices of consumer prices: transmission and dissemination of sub-indices of the HICP.

Article 3

Minimum standards for HICP weightings

(1) Each month, in current year t, Member States shall produce HICPs using sub-index weights which reflect the consumers' expenditure pattern in the weighting reference period and aim to be as representative as possible for consumers' expenditure patterns in the previous calendar year.

(2) Each year, Member States shall therefore review and update HICP sub-index weights taking into account preliminary national accounts data on consumption patterns of year t-2, except in exceptional and in duly motivated circumstances, as well as any available and relevant information from household budget surveys and other data sources which are sufficiently reliable for the purposes of the HICP.

(3) As regards weights below sub-index level, including those for Elementary Product Groups as defined in Regulation (EC) No 1749/96, Member States shall use weights which are in no case more than 7 years old.

(4) Member States shall review annually whether or not there have been any important and sustained market developments affecting quantities in the sub-divisions of COICOP/HICP, between the periods as described in paragraphs 2 and 3 and period t-1, in order to estimate weights that are as up-to-date as possible. Especially, consumption expenditure for sub-divisions of COICOP/HICP with known changes following administrative decisions and for products in fast-evolving markets shall be reviewed.

(5) Any adjustments made to weightings pursuant to this Article shall take effect with the index for January of year t. HICP weights for previous years shall be not revised, without prejudice to the possibility to correct mistakes in accordance with Article 4 of Commission Regulation (EC) No 1921/2001 ([2]) of 28 September 2001 laying down detailed rules for the implementation of Council Regulation (EC) No 2494/95 as regards minimum standards for revisions of the HICP and amending Regulation (EC) No 2602/2000. In any case, HICP weights shall take effect with the index for January each year and be price-updated to prices of the preceding December.

Article 4

Quality control

Member States shall provide the Commission (Eurostat) at its request with sufficient information on the weights used to construct the HICP, including the weighting reference period used, the outcome of the annual review and the adjustments made, for compliance with this Regulation to be evaluated.

Article 5

Application

The provisions of this Regulation shall take effect with the index for January 2012 at the latest.

Article 6

Repeal

Regulation (EC) No 2454/97 is repealed, as from January 2012. References to the repealed Regulation shall be construed as references to this Regulation.

([1]) OJ L 296, 21.11.1996, p. 8.

([2]) OJ L 261, 29.9.2001, p. 49.

Article 7

Entry into force

This Regulation shall enter into force on the 20th day following 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, 1 December 2010.

For the Commission
The President
José Manuel BARROSO

COMMISSION REGULATION (EU) No 1115/2010

of 1 December 2010

establishing the standard import values for determining the entry price of certain fruit and vegetables

THE EUROPEAN COMMISSION,

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

Having regard to Council Regulation (EC) No 1234/2007 of 22 October 2007 establishing a common organisation of agricultural markets and on specific provisions for certain agricultural products (Single CMO Regulation) [1],

Having regard to Commission Regulation (EC) No 1580/2007 of 21 December 2007 laying down implementing rules for Council Regulations (EC) No 2200/96, (EC) No 2201/96 and (EC) No 1182/2007 in the fruit and vegetable sector [2], and in particular Article 138(1) thereof,

Whereas:

Regulation (EC) No 1580/2007 lays down, pursuant to the outcome of the Uruguay Round multilateral trade negotiations, the criteria whereby the Commission fixes the standard values for imports from third countries, in respect of the products and periods stipulated in Annex XV, Part A thereto,

HAS ADOPTED THIS REGULATION:

Article 1

The standard import values referred to in Article 138 of Regulation (EC) No 1580/2007 are fixed in the Annex hereto.

Article 2

This Regulation shall enter into force on 2 December 2010.

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

Done at Brussels, 1 December 2010.

For the Commission,
On behalf of the President,
Jean-Luc DEMARTY
Director-General for Agriculture and Rural Development

[1] OJ L 299, 16.11.2007, p. 1.
[2] OJ L 350, 31.12.2007, p. 1.

ANNEX

Standard import values for determining the entry price of certain fruit and vegetables

(EUR/100 kg)

CN code	Third country code (1)	Standard import value
0702 00 00	AL	52,3
	MA	95,7
	MK	68,6
	TR	80,2
	ZZ	74,2
0707 00 05	EG	140,2
	JO	182,1
	TR	68,7
	ZZ	130,3
0709 90 70	MA	84,2
	TR	104,6
	ZZ	94,4
0805 20 10	MA	67,1
	ZZ	67,1
0805 20 30, 0805 20 50, 0805 20 70, 0805 20 90	HR	60,9
	IL	71,5
	TR	66,3
	UY	58,6
	ZZ	64,3
0805 50 10	AR	45,9
	TR	57,7
	UY	57,1
	ZZ	53,6
0808 10 80	AR	74,9
	AU	164,5
	BR	50,3
	CA	65,9
	CL	84,2
	CN	86,4
	CO	50,3
	MK	26,7
	NZ	79,7
	US	113,1
	ZA	114,1
	ZZ	82,7
0808 20 50	CN	76,9
	US	116,8
	ZZ	96,9

(1) Nomenclature of countries laid down by Commission Regulation (EC) No 1833/2006 (OJ L 354, 14.12.2006, p. 19). Code 'ZZ' stands for 'of other origin'.

DECISIONS

COUNCIL DECISION

of 29 November 2010

appointing a Finnish alternate member of the Committee of the Regions

(2010/733/EU)

THE COUNCIL OF THE EUROPEAN UNION,

Having regard to the Treaty on the Functioning of the European Union, and in particular Article 305 thereof,

Having regard to the proposal of the Finnish Government,

Whereas:

(1) On 22 December 2009 and on 18 January 2010, the Council adopted Decisions 2009/1014/EU and 2010/29/EU appointing the members and alternate members of the Committee of the Regions for the period from 26 January 2010 to 25 January 2015 [1].

(2) An alternate member's seat on the Committee of the Regions has become vacant following the end of the mandate of Mr Miikka SEPPÄLÄ,

HAS ADOPTED THIS DECISION:

Article 1

The following is hereby appointed to the Committee of the Regions as alternate member for the remainder of the current term of office, which runs until 25 January 2015:

Mr Antero SAKSALA

Pirkkalan kunnanvaltuuston jäsen

Article 2

This Decision shall take effect on the day of its adoption.

Done at Brussels, 29 November 2010.

For the Council
The President
K. PEETERS

[1] OJ L 348, 29.12.2009, p. 22, and OJ L 12, 19.1.2010, p. 11.

COMMISSION DECISION

of 30 November 2010

amending Decisions 2005/692/EC, 2005/734/EC, 2006/415/EC, 2007/25/EC and 2009/494/EC as regards avian influenza

(notified under document C(2010) 8282)

(Text with EEA relevance)

(2010/734/EU)

THE EUROPEAN COMMISSION,

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

Having regard to Council Directive 89/662/EEC of 11 December 1989 concerning veterinary checks in intra-Community trade with a view to the completion of the internal market ([1]), and in particular Article 9(4) thereof,

Having regard to Council Directive 90/425/EEC of 26 June 1990 concerning veterinary and zootechnical checks applicable in intra-Community trade in certain live animals and products with a view to the completion of the internal market ([2]), and in particular Article 10(4) thereof,

Having regard to Council Directive 91/496/EEC of 15 July 1991 laying down the principles governing the organization of veterinary checks on animals entering the Community from third countries and amending Directives 89/662/EEC, 90/425/EEC and 90/675/EEC ([3]), and in particular Article 18(7) thereof,

Having regard to Council Directive 97/78/EC of 18 December 1997 laying down the principles governing the organisation of veterinary checks on products entering the Community from third countries ([4]), and in particular Article 22(6) thereof,

Having regard to Regulation (EC) No 998/2003 of the European Parliament and of the Council of 26 May 2003 on the animal health requirements applicable to the non-commercial movement of pet animals and amending Council Directive 92/65/EEC ([5]), and in particular Article 18 thereof,

Having regard to Council Directive 2005/94/EC of 20 December 2005 on Community measures for the control of avian influenza and repealing Directive 92/40/EEC ([6]), and in particular Article 63(3) thereof,

Whereas:

(1) The Commission adopted several protection measures in relation to avian influenza, following the outbreaks of that disease in south-east Asia which started in December 2003 and was caused by the highly pathogenic avian influenza virus of the subtype H5N1.

(2) Those measures are laid down, in particular, in Commission Decision 2005/692/EC of 6 October 2005 concerning certain protection measures in relation to avian influenza in several third countries ([7]), Commission Decision 2005/734/EC of 19 October 2005 laying down biosecurity measures to reduce the risk of transmission of highly pathogenic avian influenza caused by Influenza virus A subtype H5N1 from birds living in the wild to poultry and other captive birds and providing for an early detection system in areas at particular risk ([8]) and in Commission Decision 2009/494/EC of 25 June 2009 concerning certain protection measures in relation to highly pathogenic avian influenza of subtype H5N1 in Croatia and Switzerland ([9]).

(3) The measures laid down in those Decisions are applicable until 31 December 2010. However, outbreaks of highly pathogenic avian influenza in wild birds and in poultry of the subtype H5N1 continue to occur in Member States and in third countries posing a risk to animal and human health.

(4) Given the epidemiological situation regarding avian influenza, it is appropriate to continue limiting the risks posed by imports of poultry, poultry products, pet birds and other consignments covered by those Decisions, as well as to maintain biosecurity measures, early detection systems and certain protection measures in relation to highly pathogenic avian influenza of the H5N1 subtype.

(5) The period of application of Decisions 2005/692/EC, 2005/734/EC, and 2009/494/EC should therefore be extended until 30 June 2012.

(6) In addition, Decision 2005/734/EC prohibits the use of decoy birds during bird-hunting in areas identified as being particularly at risk for the introduction of avian influenza. However, under certain conditions the competent authority may grant derogations for their use during bird-hunting and in the framework of

([1]) OJ L 395, 30.12.1989, p. 13.
([2]) OJ L 224, 18.8.1990, p. 29.
([3]) OJ L 268, 24.9.1991, p. 56.
([4]) OJ L 24, 30.1.1998, p. 9.
([5]) OJ L 146, 13.6.2003, p. 1.
([6]) OJ L 10, 14.1.2006, p. 16.
([7]) OJ L 263, 8.10.2005, p. 20.
([8]) OJ L 274, 20.10.2005, p. 105.
([9]) OJ L 166, 27.6.2009, p. 74.

Member States' programmes for avian influenza surveillance as provided for in Commission Decision 2005/732/EC of 17 October 2005 approving the programmes for the implementation of Member States' surveys for avian influenza in poultry and wild birds during 2005 and laying down reporting and eligibility rules for the Community financial contribution to the implementation costs of those programmes [1].

(7) Experience has shown that decoy birds are not only used during bird-hunting but also in the framework of research projects, ornithological studies and other activities which may pose similar risks in relation to the spread of avian influenza. The biosecurity measures of Decision 2005/734/EC should therefore apply to a wider use of decoy birds provided that the activity has been authorised by the competent authority in accordance to Article 2b(1)(d).

(8) Decision 2005/734/EC also refers to the use of decoy birds for sampling pursuant to Member States' programmes for avian influenza surveys as provided for in Decision 2005/732/EC. The surveys referred to in Decision 2005/732/EC have been completed within the given time period referred to in that Decision. Accordingly, Decision 2005/734/EC should be amended to refer to the surveillance programmes for avian influenza to be carried out by Member States under Directive 2005/94/EC.

(9) Commission Decision 2006/415/EC of 14 June 2006 concerning certain protection measures in relation to highly pathogenic avian influenza of the subtype H5N1 in poultry in the Community and repealing Decision 2006/135/EC [2] lays down certain protection measures to be applied in the event of an outbreak of that disease. Pending a possible review of those measures, the period of application of that Decision should only be extended until 31 December 2011.

(10) Commission Decision 2007/25/EC of 22 December 2006 as regards certain protection measures in relation to highly pathogenic avian influenza and movements of pet birds accompanying their owners into the Community [3] lays down certain rules regarding the authorisation of the movement from third countries of live pet birds and refers to the list of third countries set out in Council Decision 79/542/EEC of 21 December 1976 drawing up a list of third countries or parts of third countries, and laying down animal and public health and veterinary certification conditions, for importation into the Community of certain live animals and their fresh meat [4].

(11) Commission Regulation (EU) No 206/2010 of 12 March 2010 laying down lists of third countries, territories or parts thereof authorised for the introduction into the European Union of certain animals and fresh meat and the veterinary certification requirements [5] replaces and repeals Decision 79/542/EEC. Accordingly, it is appropriate to update Decision 2007/25/EC by referring to Regulation (EU) No 206/2010.

(12) In addition, Article 1 of Decision 2007/25/EC and the model veterinary certificate set out in Annex II to that Decision, which refer to Chapter 2.1.14 of the Manual of Diagnostic Tests and Vaccines for Terrestrial Animals of the World Organisation for Animal Health (OIE), are out of date since the adoption of the revised Chapter on avian influenza in May 2009 and should be updated to refer to Chapter 2.3.4 of that manual. It is also necessary to make certain amendments to the owner's declaration set out in Annex III to that Decision in the light of experience. Decision 2007/25/EC should therefore be amended accordingly.

(13) Taking into account the animal health situation, it is also appropriate to extend the period of application of Decision 2007/25/EC until 30 June 2012.

(14) Decisions 2005/692/EC, 2005/734/EC, 2006/415/EC, 2007/25/EC and 2009/494/EC should therefore be amended accordingly.

(15) It is necessary to provide for a transitional period during which consignments of pet birds for which the necessary veterinary certificate and declaration of the owner have been issued in accordance with Decision 2007/25/EC, before the amendments made by this Decision, may continue to be introduced into the Union, in order to give the Member States and the industry time to adjust to the new rules.

(16) The measures provided for in this Decision are in accordance with the opinion of the Standing Committee on the Food Chain and Animal Health,

HAS ADOPTED THIS DECISION:

Article 1

In Article 7 of Decision 2005/692/EC, the date '31 December 2010' is replaced by '30 June 2012'.

Article 2

Decision 2005/734/EC is amended as follows:

1. in Article 1, paragraph 4 is replaced by the following:

[1] OJ L 274, 20.10.2005, p. 95.
[2] OJ L 164, 16.6.2006, p. 51.
[3] OJ L 8, 13.1.2007, p. 29.
[4] OJ L 146, 14.6.1979, p. 15.
[5] OJ L 73, 20.3.2010, p. 1.

'4. Member States shall regularly review the measures they have taken pursuant to paragraph 1, and in the light of the surveillance programmes that they have carried out in accordance with Article 4 of Council Directive 2005/94/EC (*), in order to adjust to the changing epidemiological and ornithological situation, the areas of their territory that they have identified as being particularly at risk from the introduction of avian influenza.

(*) OJ L 10, 14.1.2006, p. 16.';

2. in Article 2a(1), point (d) is replaced by the following:

'(d) the use of birds of the orders *Anseriformes* and *Charadriiformes* as decoy birds (decoy birds).';

3. in Article 2b(1), point (d) is amended as follows:

(a) the introductory phrase and (i) is replaced by the following:

'(d) the use of decoy birds:

(i) by decoy bird holders registered with the competent authority, under the strict supervision of the competent authority for the attraction of wild birds intended for sampling pursuant to the Member States' programmes for avian influenza, research projects, ornithological studies or any other activity approved by the competent authority; or';

(b) in (ii), the third indent is replaced by the following:

'— the recording and reporting of the health status of decoy birds and laboratory testing for avian influenza in the case of deaths of such birds and at the end of the period of use in the area identified at particular risk for the introduction of avian influenza,';

4. in Article 4, the date '31 December 2010' is replaced by '30 June 2012'.

Article 3

In Article 12 of Decision 2006/415/EC, the date '31 December 2010' is replaced by '31 December 2011'.

Article 4

Decision 2007/25/EC is amended as follows:

1. in Article 1(1), point (b) is amended as follows:

(a) point (i) is replaced by the following:

'(i) have undergone isolation for 30 days prior to export at the place of departure in a third country listed in Part 1 of Annex I or Part 1 of Annex II to Commission Regulation (EU) No 206/2010 (*), or

(*) OJ L 73, 20.3.2010, p. 1.';

(b) point (iv) is replaced by the following:

'(iv) have been in isolation for at least 10 days prior to export and have undergone a test to detect the avian influenza H5N1 antigen or genome as laid down in the Chapter on avian influenza of the Manual of Diagnostic tests and Vaccines for Terrestrial Animals, as regularly updated by the OIE, carried out on a sample taken not earlier than the third day of isolation.';

2. in Article 6, the date '31 December 2010' is replaced by '30 June 2012';

3. Annexes II and III are replaced by the text in the Annex to this Decision.

Article 5

In Article 3 of Decision 2009/494/EC, the date '31 December 2010' is replaced by '30 June 2012'.

Article 6

The Member States shall immediately take the necessary measures to comply with this Decision and publish those measures. They shall immediately inform the Commission thereof.

Article 7

For a transitional period until 31 March 2011, pet birds for which the veterinary certificate and declaration of the owner have been issued in accordance with Decision 2007/25/EC, before the amendments made by this Decision, may continue to be introduced into the Union.

Article 8

This Decision is addressed to the Member States.

Done at Brussels, 30 November 2010.

For the Commission

John DALLI

Member of the Commission

ANNEX

Annexes II and III to Decision 2007/25/EC are replaced by the following:

'ANNEX II

COUNTRY **Veterinary certificate to EU**

Part I: Details of dispatched consignment

I.1. Consignor Name Address Tel.	I.2. Certificate reference No	I.2.a.
	I.3. Central competent authority	
	I.4. Local competent authority	
I.5. Consignee Name Address Tel.	I.6.	
I.7. Country of origin ISO code I.8. Region of origin Code	I.9. Country of destination ISO code	I.10.
I.11. Place of origin Name Approval number Address Name Approval number Address Name Approval number Address	I.12. Place of destination Name Address Postal code	
I.13. Place of loading	I.14. Date of departure	
I.15. Means of transport Aeroplane ☐ Ship ☐ Railway wagon ☐ Road vehicle ☐ Other ☐ Identification Documentary references	I.16.	
	I.17. No(s) of CITES	
I.18. Description of commodity	I.19. Commodity code (HS code)	
	I.20. Quantity	
I.21.	I.22. Number of packages	
I.23. Seal/Container No	I.24.	
I.25. Commodities certified for: Pets ☐ Quarantine ☐		
I.26.	I.27. For import or admission into EU ☐	

I.28. Identification of the commodities

Species (Scientific name)	Identification system	Identification number	Quantity

COUNTRY **Pet birds**

Part II: Certification

II. Health information	II.a. Certificate reference number	II.b.

I, the undersigned official veterinarian of ..... (insert name of third country) certify that:

1. The country of dispatch is a member country of the World Organisation for Animal Health (OIE) and belongs to the OIE Regional Commission for ..... (insert name of Regional Commission).

2. The bird(s) described in point I.28 has/have been subjected today, within 48 hours or the last working day prior to the date of dispatch, to a clinical inspection and found free of obvious signs of disease;

([1]) *either* [3. The bird(s) complies/comply with at least one of the following conditions:

([1]) *either* [it/they comes/come from a third country listed in Part 1 of Annex I or Part 1 of Annex II to Regulation (EU) No 206/2010 and has/have been confined on the premises specified in point I.11 under official supervision for at least 30 days prior to the date of dispatch and effectively protected from contact with any other bird(s);]

([1]) *or* [it/they has/have been vaccinated on ..... [dd/mm/yyyy] and at least on one occasion re-vaccinated on ..... [dd/mm/yyyy] within the last 6 months and not later than 60 days prior to the date of dispatch, in accordance with the manufacturer's instructions against avian influenza using an H5 vaccine, which is not a live vaccine and approved for the species concerned in the third country of dispatch or at least in one Member State of the European Union;]

([1]) *or* [it/they has/have been isolated for at least 10 days prior to the date of dispatch and have been subjected to a test for the detection of avian influenza H5N1 antigen or genome, as prescribed in the Chapter 2.3.4 on avian influenza in the Manual of Diagnostic Tests and Vaccines for Terrestrial Animals of the OIE, as regularly updated, carried out on a sample taken on ..... [dd/mm/yyyy], not earlier than on the third day of isolation;]]

([1]) *or* [3. The owner/person responsible for the bird(s) has declared that he/she has made arrangements for the 30 days post-introduction quarantine in an approved quarantine facility or centre in accordance with Article 6 of Regulation (EC) No 318/2007].

4. The owner or the representative of the owner has declared that:

4.1. The bird(s) is/are "pet animal(s)" as defined in Article 3(a) of Regulation (EC) No 998/2003 intended for non-commercial movement.

4.2. During the period between the pre-movement veterinary inspection and the factual departure the bird(s) will remain isolated from any possible contact with any other bird(s).

([1]) *either* [4.3. The bird(s) has/have been confined on the premises for at least the 30 days immediately prior to the date of dispatch without coming into contact with any other bird(s).]

([1]) *or* [4.3. The bird(s) has/have undergone the 10 days pre-movement isolation.]

([1]) *or* [4.3. Arrangements have been made for the 30 days post-introduction quarantine of the bird(s) at the quarantine premises of .....]

Notes

Part I

— Box reference I.8: Provide the code of territory as appearing in Part 1 of Annex I or in Part 1 of Annex II to Regulation (EU) No 206/2010,

— Box reference I.15: Registration number (railway wagons or container and lorries), flight number (aircraft) or name (ship) is to be provided,

— Box reference I.19: Use the appropriate HS codes: 01.06.31, 01.06.32, 01.06.39,

— Box reference I.23: For containers or boxes, the container number and the seal number (if applicable) should be included,

— Box reference I.28: *Identification system:* The birds must bear:

An individual number which permits tracing of their premises of origin. Specify the identification system (such as clip, leg band, microchip, transponder, tag).

COUNTRY **Pet birds**

II. Health information	II.a. Certificate reference number	II.b.
Part II: (1) Keep as appropriate. This certificate is valid for 10 days. In the case of transport by boat the validity is prolonged by the time of the sea voyage.		
Official veterinarian or official inspector Name (in capital letters): Qualification and title: Local veterinary unit: LVU No: Date: Signature: Stamp:		

ANNEX III

DECLARATION

I, the undersigned owner [a]/person responsible for the bird(s) on behalf of the owner [a] declare:

1. The bird(s) is/are accompanying the undersigned person and are not intended to be sold or transferred to another owner.

2. The bird(s) will remain under the responsibility of the undersigned person during its/their non-commercial movement.

3. During the period between the pre-movement veterinary inspection and the actual departure the bird(s) will remain isolated from any possible contact with any other bird(s); and

4. *[a] either* [The bird(s) has/have been confined on the premises for a period of at least 30 days immediately prior to the date of dispatch without coming into contact with any other bird(s).]

   [a] or [The bird(s) has/have undergone the 10 days pre-movement isolation.]

   [a] or [I have made arrangements for the 30 days post-introduction quarantine of the bird(s) at the quarantine premises of ....., as indicated in the corresponding Certificate.]

....

(Date and Place) (Signature)

[a] Keep as appropriate.'

COMMISSION DECISION

of 1 December 2010

on financial aid from the Union for the year 2011 for certain European Union reference laboratories in the field of animal health and live animals

(notified under document C(2010) 8344)

(Only the Danish, English, French, German, Spanish and Swedish texts are authentic)

(2010/735/EU)

THE EUROPEAN COMMISSION,

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

Having regard to Council Decision 2009/470/EC of 25 May 2009 on expenditure in the veterinary field [1], and in particular Article 31(1) thereof,

Having regard to Regulation (EC) No 882/2004 of the European Parliament and of the Council of 29 April 2004 on official controls performed to ensure the verification of compliance with feed and food law, animal health and animal welfare rules [2], and in particular Article 32(7) thereof,

Whereas:

(1) Pursuant to Article 31(1) of Decision 2009/470/EC European Union reference laboratories in the field of animal health and live animals may be granted Union aid.

(2) Commission Regulation (EC) No 1754/2006 of 28 November 2006 laying down detailed rules for the granting of Community financial assistance to Community reference laboratories for feed and food and the animal health sector [3] provides that the financial assistance from the Union is to be granted if the approved work programmes are efficiently carried out and the beneficiaries supply all the necessary information within certain time limits.

(3) In accordance with Article 2 of Regulation (EC) No 1754/2006 the relationship between the Commission and European Union reference laboratories is laid down in a partnership agreement which is supported by a multiannual work programme.

(4) The Commission has assessed the work programmes and corresponding budget estimates submitted by the European Union reference laboratories for the year 2011.

(5) Accordingly, Union financial assistance should be granted to the European Union reference laboratories designated to carry out the functions and duties provided for in the following acts:

— Council Directive 92/35/EEC of 29 April 1992 laying down control rules and measures to combat African horse sickness [4],

— Council Directive 92/66/EEC of 14 July 1992 introducing Community measures for the control of Newcastle disease [5],

— Council Directive 92/119/EEC of 17 December 1992 introducing general Community measures for the control of certain animals diseases and specific measures relating to swine vesicular disease [6],

— Council Directive 93/53/EEC of 24 June 1993 introducing minimum Community measures for the control of certain fish diseases [7],

— Council Directive 95/70/EC of 22 December 1995 introducing minimum Community measures for the control of certain diseases affecting bivalve molluscs [8],

— Council Decision 2000/258/EC of 20 March 2000 designating a specific institute responsible for establishing the criteria necessary for standardising the serological tests to monitor the effectiveness of rabies vaccines [9],

[1] OJ L 155, 18.6.2009, p. 30.
[2] OJ L 165, 30.4.2004, p. 1.
[3] OJ L 331, 29.11.2006, p. 8.
[4] OJ L 157, 10.6.1992, p. 19.
[5] OJ L 260, 5.9.1992, p. 1.
[6] OJ L 62, 15.3.1993, p. 69.
[7] OJ L 175, 19.7.1993, p. 23.
[8] OJ L 332, 30.12.1995, p. 33.
[9] OJ L 79, 30.3.2000, p. 40.

— Council Directive 2000/75/EC of 20 November 2000 laying down specific provisions for the control and eradication of bluetongue ([1]),

— Council Directive 2001/89/EC of 23 October 2001 on Community measures for the control of classical swine fever ([2]),

— Council Directive 2002/60/EC of 27 June 2002 laying down specific provisions for the control of African swine fever and amending Directive 92/119/EEC as regards Teschen disease and African swine fever ([3]),

— Council Directive 2003/85/EC of 29 September 2003 on Community measures for the control of foot-and-mouth disease repealing Directive 85/511/EEC and Decisions 89/531/EEC and 91/665/EEC and amending Directive 92/46/EEC ([4]),

— Council Decision 96/463/EC of 23 July 1996 designating the reference body responsible for collaborating in rendering uniform the testing methods and the assessment of the results for pure-bred breeding animals of the bovine species ([5]),

— Regulation (EC) No 882/2004 for brucellosis,

— Council Directive 2005/94/EC of 20 December 2005 on Community measures for the control of avian influenza and repealing Directive 92/40/EEC ([6]),

— Council Directive 2006/88/EC of 24 October 2006 on animal health requirements for aquaculture animals and products thereof, and on the prevention and control of certain diseases in aquatic animals ([7]),

— Commission Regulation (EC) No 180/2008 of 28 February 2008 concerning the Community reference laboratory for equine diseases other than African horse sickness and amending Annex VII to Regulation (EC) No 882/2004 of the European Parliament and of the Council ([8]),

— Commission Regulation (EC) No 737/2008 of 28 July 2008 designating the Community reference laboratories for crustacean diseases, rabies and bovine tuberculosis, laying down additional responsibilities and tasks for the Community reference laboratories for rabies and bovine tuberculosis and amending Annex VII to Regulation (EC) No 882/2004 of the European Parliament and of the Council ([9]).

(6) Financial assistance for the operation and organisation of workshops of European Union reference laboratories should also be in conformity with the eligibility rules laid down in Regulation (EC) No 1754/2006.

(7) Regulation (EC) No 1754/2006 lays down eligibility rules for the workshops organised by the European Union reference laboratories. It also limits the financial assistance to a maximum of 32 participants in workshops. Derogations to that limitation should be provided in accordance with Article 13(3) of Regulation (EC) No 1754/2006 to some European Union reference laboratories that needs support for attendance by more than 32 participants in order to achieve the best outcome of its workshops. Derogations can be obtained in case a European Union reference laboratory takes the leadership and responsibility of organising a workshop with another European Union reference laboratory.

(8) In accordance with Articles 3(2)(a) and 13 of Council Regulation (EC) No 1290/2005 of 21 June 2005 on the financing of the common agricultural policy ([10]), animal disease eradication and control programmes (veterinary measures) shall be financed from the European Agricultural Guarantee Fund (EAGF). Furthermore, Article 13, second paragraph of that regulation foresees that in duly justified exceptional cases, for measures and programmes covered by Decision 2009/470/EC, expenditure relating to administrative and personnel costs incurred by Member States and beneficiaries of aid from the EAGF shall be borne by the Fund. For financial control purposes, Articles 9, 36 and 37 of Regulation (EC) No 1290/2005 are to apply.

(9) The measures provided for in this Decision are in accordance with the opinion of the Standing Committee on the Food Chain and Animal Health,

HAS ADOPTED THIS DECISION:

Article 1

For African horse sickness, the Union grants financial assistance to the Laboratorio Central de Sanidad Animal de Algete, Algete (Madrid), Spain, to carry out the functions and duties set out in Annex III to Directive 92/35/EEC.

([1]) OJ L 327, 22.12.2000, p. 74.
([2]) OJ L 316, 1.12.2001, p. 5.
([3]) OJ L 192, 20.7.2002, p. 27.
([4]) OJ L 306, 22.11.2003, p. 1.
([5]) OJ L 192, 2.8.1996, p. 19.
([6]) OJ L 10, 14.1.2006, p. 16.
([7]) OJ L 328, 24.11.2006, p. 14.
([8]) OJ L 56, 29.2.2008, p. 4.
([9]) OJ L 201, 30.7.2008, p. 29.
([10]) OJ L 209, 11.8.2005, p. 1.

The Union's financial assistance shall be at the rate of 100 % of the eligible costs as defined in Regulation (EC) No 1754/2006 to be incurred by that laboratory for the work programme and shall amount to a maximum of EUR 105 000 for the period from 1 January to 31 December 2011.

Article 2

For Newcastle disease, the Union grants financial assistance to the Veterinary Laboratories Agency (VLA, ex-CVL), New Haw, Weybridge, United Kingdom, to carry out the functions and duties set out in Annex V to Directive 92/66/EEC.

The Union's financial assistance shall be at the rate of 100 % of the eligible costs as defined in Regulation (EC) No 1754/2006 to be incurred by that laboratory for the work programme and shall amount to a maximum of EUR 90 850 for the period from 1 January to 31 December 2011, of which a maximum of EUR 850 shall be dedicated to the organisation of a restricted technical workshop on Newcastle disease.

Article 3

For swine vesicular disease, the Union grants financial assistance to the AFRC Institute for Animal Health, Pirbright Laboratory, Pirbright, United Kingdom, to carry out the functions and duties set out in Annex III to Directive 92/119/EEC.

The Union's financial assistance shall be at the rate of 100 % of the eligible costs as defined in Regulation (EC) No 1754/2006 to be incurred by that laboratory for the work programme and shall amount to a maximum of EUR 130 000 for the period from 1 January to 31 December 2011.

Article 4

For fish diseases, the Union grants financial assistance to the Technical University of Denmark, National Veterinary Institute, Department of Poultry, Fish and Fur Animals, Aarhus, Denmark, to carry out the functions and duties set out in Annex VI to Directive 2006/88/EC.

The Union's financial assistance shall be at the rate of 100 % of the eligible costs as defined in Regulation (EC) No 1754/2006 to be incurred by that laboratory for the work programme and shall amount to a maximum of EUR 318 000 for the period from 1 January to 31 December 2011, of which a maximum of EUR 40 000 shall be dedicated to the organisation of a technical workshop on fish diseases.

Article 5

For diseases of bivalve molluscs, the Union grants financial assistance to Ifremer, La Tremblade, France, to carry out the functions and duties set out in Annex VI to Directive 2006/88/EC.

The Union's financial assistance shall be at the rate of 100 % of the eligible costs as defined in Regulation (EC) No 1754/2006 to be incurred by that institute for the work programme and shall amount to a maximum of EUR 130 000 for the period from 1 January to 31 December 2011.

Article 6

For bluetongue, the Union grants financial assistance to the AFRC Institute for Animal Health, Pirbright Laboratory, Pirbright, United Kingdom, to carry out the functions and duties set out in Annex II(B) to Directive 2000/75/EC.

The Union's financial assistance shall be at the rate of 100 % of the eligible costs as defined in Regulation (EC) No 1754/2006 to be incurred by that laboratory for the work programme and shall amount to a maximum of EUR 300 000 for the period from 1 January to 31 December 2011.

Article 7

For classical swine fever, the Union grants financial assistance to the Institut für Virologie der Tierärztlichen Hochschule Hannover, Hannover, Germany, to carry out the functions and duties set out in Annex IV to Directive 2001/89/EC.

The Union's financial assistance shall be at the rate of 100 % of the eligible costs as defined in Regulation (EC) No 1754/2006 to be incurred by that institute for the work programme and shall amount to a maximum of EUR 340 000 for the period from 1 January to 31 December 2011 of which a maximum of EUR 49 000 shall be dedicated to the organisation of a technical workshop on classical swine fever.

Article 8

For African swine fever, the Union grants financial assistance to the Centro de Investigación en Sanidad Animal, Valdeolmos, Madrid, Spain, to carry out the functions and duties set out in Annex V to Directive 2002/60/EC.

The Union's financial assistance shall be at the rate of 100 % of the eligible costs as defined in Regulation (EC) No 1754/2006 to be incurred by that research centre for the work programme and shall amount to a maximum of EUR 200 000 for the period from 1 January to 31 December 2011, of which a maximum of EUR 40 000 shall be dedicated to the organisation of a technical workshop on African swine fever.

Article 9

For foot-and-mouth disease, the Union grants financial assistance to the Institute for Animal Health, Pirbright Laboratory, of the Biotechnology and Biological Sciences Research Council (BBSRC), Pirbright, United Kingdom, to carry out the functions and duties set out in Annex XVI to Directive 2003/85/EC.

The Union's financial assistance shall be at the rate of 100 % of the eligible costs as defined in Regulation (EC) No 1754/2006 to be incurred by that laboratory for the work programme and shall amount to a maximum of EUR 360 000 for the period from 1 January to 31 December 2011.

Article 10

For collaborating in rendering uniform the testing methods and the assessment of the results for pure-bred breeding animals of the bovine species, the Union grants financial assistance to the Interbull Centre, Department of Animal Breeding and Genetics, Swedish University of Agricultural Sciences, Uppsala, Sweden, to carry out the functions and duties set out in Annex II to Decision 96/463/EC.

The Union's financial assistance shall be at the rate of 100 % of the eligible costs as defined in Regulation (EC) No 1754/2006 to be incurred by that centre for the work programme and shall amount to a maximum of EUR 150 000 for the period from 1 January to 31 December 2011.

Article 11

For brucellosis, the Union grants financial assistance to ANSES (ex-AFSSA), Laboratoire d'études et de recherches en pathologie animale et zoonoses, Maisons-Alfort, France, to carry out the functions and duties set out in Article 32(2) of Regulation (EC) No 882/2004.

The Union's financial assistance shall be at the rate of 100 % of the eligible costs as defined in Regulation (EC) No 1754/2006 to be incurred by that laboratory for the work programme and shall amount to a maximum of EUR 275 000 for the period from 1 January to 31 December 2011, of which a maximum of EUR 25 000 shall be dedicated to the organisation of a technical workshop on brucellosis.

Article 12

For avian influenza, the Union grants financial assistance to the Veterinary Laboratories Agency (VLA, ex-CVL), New Haw, Weybridge, United Kingdom, to carry out the functions and duties set out in Annex VII to Directive 2005/94/EC.

The Union's financial assistance shall be at the rate of 100 % of the eligible costs as defined in Regulation (EC) No 1754/2006 to be incurred by that laboratory for the work programme and shall amount to a maximum of EUR 385 850 for the period from 1 January to 31 December 2011, of which a maximum of EUR 850 shall be dedicated to the organisation of a restricted technical workshop on avian influenza.

Article 13

For crustacean diseases, the Union grants financial assistance to the Centre for Environment, Fisheries & Aquaculture Science (Cefas), Weymouth Laboratory, United Kingdom, to carry out the functions and duties set out in Part I Annex VI to Directive 2006/88/EC.

The Union's financial assistance shall be at the rate of 100 % of the eligible costs as defined in Regulation (EC) No 1754/2006 to be incurred by that laboratory for the work programme and shall amount to a maximum of EUR 150 000 for the period from 1 January to 31 December 2011 of which a maximum of EUR 40 000 shall be dedicated to the organisation of a technical workshop on crustacean diseases.

Article 14

For equine diseases other than African Horse Sickness, the Union grants financial assistance to ANSES (ex-AFSSA), Laboratoire d'études et de recherches en pathologie animale et zoonoses/Laboratoire d'études et de recherche en pathologie equine, France, to carry out the functions and duties set out in the Annex to Regulation (EC) No 180/2008.

The Union's financial assistance shall be at the rate of 100 % of the eligible costs as defined in Regulation (EC) No 1754/2006 to be incurred by that laboratory for the work programme and shall amount to a maximum of EUR 540 000 for the period from 1 January to 31 December 2011 of which a maximum of EUR 40 000 shall be dedicated to the organisation of a technical workshop on equine diseases.

Article 15

For rabies, the Union grants financial assistance to ANSES (ex-AFSSA), Laboratoire d'études sur la rage et la pathologie des animaux sauvages, Nancy, France, to carry out the functions and duties set out in Annex I to Regulation (EC) No 737/2008.

The Union's financial assistance shall be at the rate of 100 % of the eligible costs as defined in Regulation (EC) No 1754/2006 to be incurred by that laboratory for the work programme and shall amount to a maximum of EUR 275 000 for the period from 1 January to 31 December 2011, of which a maximum of EUR 25 000 shall be dedicated to the organisation of a technical workshop on rabies.

Article 16

For tuberculosis, the Union grants financial assistance to the Laboratorio de Vigilancia Veterinaria (Visavet) of the Facultad de Veterinaria, Universidad Complutense de Madrid, Madrid, Spain, to carry out the functions and duties set out in Annex II to Regulation (EC) No 737/2008.

The Union's financial assistance shall be at the rate of 100 % of the eligible costs as defined in Regulation (EC) No 1754/2006 to be incurred by that laboratory for the work programme and shall amount to a maximum of EUR 245 000 for the period from 1 January to 31 December 2011, of which a maximum of EUR 30 000 shall be dedicated to the organisation of a technical workshop on tuberculosis.

Article 17

This Decision is addressed to:

— for African horse sickness: Laboratorio Central de Sanidad Animal, Ministerio de Agricultura, PESCA y Alimentación, Ctra. De Algete km. 8, Valdeolmos, 28110, Algete (Madrid), Spain;

— for Newcastle disease: Veterinary Laboratories Agency, Weybridge, New Haw, Addelstone, Surrey KT15 3NB, United Kingdom;

— for swine vesicular disease: AFRC Institute for Animal Health, Pirbright Laboratory, Pirbright, Woking, Surrey GU24 ONF, United Kingdom;

— for fish diseases: the Technical University of Denmark, National Veterinary Institute, Department of Poultry, Fish and Fur Animals, Hangøvej 2, 8200-Århus Denmark;

— for diseases of bivalve molluscs: Ifremer, B.P. 133 17390 La Tremblade, France;

— for bluetongue: AFRC Institute for Animal Health, Pirbright Laboratory, Pirbright, Woking, Surrey GU24 ONF, United Kingdom;

— for classical swine fever: Institut für Virologie der Tierärztlichen Hochschule, Bischofsholer Damm 15, 3000 Hannover, Germany;

— for African swine fever: Centro de Investigación en Sanidad Animal, Ctra. De Algete a El Casar, Valdeolmos 28130, Madrid, Spain;

— for foot-and-mouth disease: AFRC Institute for Animal Health, Pirbright Laboratory, Pirbright, Woking, Surrey GU24 ONF, United Kingdom;

— Interbull Centre, Department of Animal Breeding and Genetics SLU, Swedish University of Agricultural Sciences, Box: 7023, SE-750 07 Uppsala Sweden;

— for brucellosis: ANSES, Laboratoire d'études et de recherches en pathologie animale et zoonoses, 23 avenue du Général de Gaulle 94 706 Maisons-Alfort, Cedex France;

— for Avian influenza: Veterinary Laboratories Agency, Weybridge, New Haw, Addelstone, Surrey KT15 3NB, United Kingdom;

— for crustacean diseases: Centre for Environment, Fisheries & Aquaculture Science (Cefas), Weymouth Laboratory, The Nothe, Barrack Road, Weymouth, Dorset DT4 8UB, United Kingdom;

— for equine diseases: ANSES, Laboratoire d'études et de recherches en pathologie animale et zoonoses, 23 avenue du Général de Gaulle 94 706 Maisons-Alfort, Cedex France;

— for rabies: ANSES, Laboratoire d'études sur la rage et la pathologie des animaux sauvages, site de Nancy, Domaine de Pixérécourt, BP 9, 54220 Malzéville, France;

— for tuberculosis: Visavet — Laboratorio de vigilancia veterinaria, Facultad de Veterinaria, Universidad Complutense de Madrid, Avda. Puerta de Hierro, s/n. Ciudad Universitaria, 28040 Madrid, Spain.

Done at Brussels, 1 December 2010.

For the Commission

John DALLI

Member of the Commission

COMMISSION DECISION

of 1 December 2010

as regards a Union financial contribution for the year 2011, to certain European Union reference laboratories in the feed and food control area

(notified under document C(2010) 8350)

(Only the Danish, Dutch, English, French, German, Italian, Spanish and Swedish texts are authentic)

(2010/736/EU)

THE EUROPEAN COMMISSION,

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

Having regard to Regulation (EC) No 882/2004 of the European Parliament and of the Council of 29 April 2004 on official controls performed to ensure the verification of compliance with feed and food law, animal health and animal welfare rules [1], and in particular Article 32(7) thereof,

Whereas:

(1) European Union reference laboratories in the food and feed control area may be granted a Union financial contribution in accordance with Article 31 of Council Decision 2009/470/EC of 25 May 2009 on expenditure in the veterinary field [2].

(2) Commission Regulation (EC) No 1754/2006 of 28 November 2006 laying down detailed rules for the granting of Community financial assistance to Community reference laboratories for feed and food and the animal health sector [3] provides that the financial contribution from the Union is to be granted if the approved work programmes are efficiently carried out and that the beneficiaries supply all the necessary information within certain time limits.

(3) In accordance with Article 2 of Regulation (EC) No 1754/2006 the relationship between the Commission and each European Union reference laboratory is laid down in a partnership agreement which is supported by a multiannual work programme.

(4) The Commission has assessed the work programmes and corresponding budget estimates submitted by the European Union reference laboratories for the year 2011.

(5) Accordingly, a Union financial contribution should be granted to the European Union reference laboratories designated in order to co-finance their activities to carry out the functions and duties provided for in Regulation (EC) No 882/2004. The Union's financial contribution should be at the rate of 100 % of eligible costs as defined in Regulation (EC) No 1754/2006.

(6) Regulation (EC) No 1754/2006 lays down eligibility rules for the workshops organised by the European Union reference laboratories. It also limits the financial assistance to a maximum of 32 participants in workshops. Derogations to that limitation should be provided in accordance with Article 13(3) of Regulation (EC) No 1754/2006 to some European Union reference laboratory that needs support for attendance by more than 32 participants in order to achieve the best outcome of its workshops. Derogations can be obtained in case a European Union reference laboratory takes the leadership and responsibility when organising a workshop with another European Union reference laboratory.

(7) In accordance with Article 3(2)(a) of Council Regulation (EC) No 1290/2005 of 21 June 2005 on the financing of the common agricultural policy [4], animal disease eradication and control programmes (veterinary measures) shall be financed from the European Agricultural Guarantee Fund (EAGF). Furthermore, Article 13, second paragraph of that Regulation foresees that in duly justified exceptional cases, for measures and programmes covered by Council Decision 90/424/EEC of 26 June 1990 on expenditure in the veterinary field [5], expenditure relating to administrative and personnel costs incurred by Member States and beneficiaries of aid from the EAGF shall be borne by the Fund. For financial control purposes, Articles 9, 36 and 37 of Regulation (EC) No 1290/2005 are to apply.

(8) The measures provided for in this Decision are in accordance with the opinion of the Standing Committee of the Food Chain and Animal Health,

[1] OJ L 165, 30.4.2004, p. 1.
[2] OJ L 155, 18.6.2009, p. 30.
[3] OJ L 331, 29.11.2006, p. 8.
[4] OJ L 209, 11.8.2005, p. 1.
[5] OJ L 224, 18.8.1990, p. 19.

HAS ADOPTED THIS DECISION:

Article 1

1. The European Union grants financial aid to the Laboratoire d'études et de recherches sur la qualité des aliments et sur les procédés agroalimentaires (Lerqap), of the Agence nationale de sécurité sanitaire de l'alimentation, de l'environnement et du travail (Anses, ex-AFSSA), Maisons-Alfort, France, to carry out the functions and duties provided for in Article 32 of Regulation (EC) No 882/2004, for the analysis and testing of milk and milk products.

For the period from 1 January 2011 to 31 December 2011, that financial aid shall not exceed EUR 355 820.

2. In addition to the maximum amount provided for in paragraph 1, the Union grants financial aid to the laboratory referred to in paragraph 1 for the organisation of workshops. That aid shall not exceed EUR 23 000.

Article 2

1. The European Union grants financial aid to the Rijksinstituut voor Volksgezondheid en Milieu (RIVM), Bilthoven, the Netherlands, to carry out the functions and duties provided for in Article 32 of Regulation (EC) No 882/2004, for the analysis and testing of zoonoses (salmonella).

For the period from 1 January 2011 to 31 December 2011, that financial aid shall not exceed EUR 373 450.

2. In addition to the maximum amount provided for in paragraph 1, the Union grants financial aid to the laboratory referred to in paragraph 1 for the organisation of workshops. That aid shall not exceed EUR 30 300.

Article 3

1. The European Union grants financial aid to the Laboratorio de Biotoxinas Marinas, Agencia Española de Seguridad Alimentaria y Nutrición (Ministerio de Sanidad y Política Social), Vigo, Spain, to carry out the functions and duties provided for in Article 32 of Regulation (EC) No 882/2004, for the monitoring of marine biotoxins.

For the period from 1 January 2011 to 31 December 2011, that financial aid shall not exceed EUR 283 302.

2. In addition to the maximum amount provided for in paragraph 1, the Union grants financial aid to the laboratory referred to in paragraph 1 for the organisation of workshops. That aid shall not exceed EUR 44 500.

Article 4

1. The European Union grants financial aid to the laboratory of the Centre for Environment, Fisheries and Aquaculture Science, Weymouth, United Kingdom, to carry out the functions and duties provided for in Article 32 of Regulation (EC) No 882/2004, for the monitoring of viral and bacteriological contamination of bivalve molluscs.

For the period from 1 January 2011 to 31 December 2011, that financial aid shall not exceed EUR 289 832.

2. In addition to the maximum amount provided for in paragraph 1, the Union grants financial aid to the laboratory referred to in paragraph 1 for the organisation of workshops. That aid shall not exceed EUR 46 800.

Article 5

1. The European Union grants a financial contribution to the Laboratoire d'études et de recherches sur la qualité des aliments et sur les procédés agroalimentaires (Lerqap), of the Agence nationale de sécurité sanitaire de l'alimentation, de l'environnement et du travail (Anses, ex-AFSSA), Maisons-Alfort, France, to carry out the functions and duties provided for in Article 32 of Regulation (EC) No 882/2004, for the analysis and testing of *Listeria monocytogenes*.

For the period from 1 January 2011 to 31 December 2011, that financial contribution shall not exceed EUR 426 065.

2. In addition to the maximum amount provided for in paragraph 1, the Union grants a financial contribution to the laboratory referred to in paragraph 1 for the organisation of workshops. That contribution shall not exceed EUR 23 000.

Article 6

1. The European Union grants financial aid to the Laboratoire d'études et de recherches sur la qualité des aliments et sur les procédés agroalimentaires (Lerqap), of the Agence nationale de sécurité sanitaire de l'alimentation, de l'environnement et du travail (Anses, ex-AFSSA), Maisons-Alfort, France, to carry out the functions and duties provided for in Article 32 of Regulation (EC) No 882/2004, for the analysis and testing of coagulase positive *Staphylococci*, including *Staphylococcus aureus*.

For the period from 1 January 2011 to 31 December 2011, that financial contribution shall not exceed EUR 361 615.

2. In addition to the maximum amount provided for in paragraph 1, the Union grants a financial contribution to the laboratory referred to in paragraph 1 for the organisation of workshops. That contribution shall not exceed EUR 23 000.

Article 7

1. The European Union grants a financial contribution to the Istituto Superiore di Sanità (ISS), Rome, Italy, to carry out the functions and duties provided for in Article 32 of Regulation (EC) No 882/2004, for the analysis and testing of *Escherichia coli*, including Verotoxigenic *E. Coli* (VTEC).

For the period from 1 January 2011 to 31 December 2011, that financial contribution shall not exceed EUR 269 296.

2. In addition to the maximum amount provided for in paragraph 1, the Union grants a financial contribution to the laboratory referred to in paragraph 1 for the organisation of workshops. That contribution shall not exceed EUR 22 000.

Article 8

1. The European Union grants a financial contribution to the Statens Veterinärmedicinska Anstalt (SVA), Uppsala, Sweden, to carry out the functions and duties provided for in Article 32 of Regulation (EC) No 882/2004, for the monitoring of *Campylobacter*.

For the period from 1 January 2011 to 31 December 2011, that financial contribution shall not exceed EUR 305 386.

2. In addition to the maximum amount provided for in paragraph 1, the Union grants a financial contribution to the laboratory referred to in paragraph 1 for the organisation of workshops. That contribution shall not exceed EUR 30 000.

Article 9

1. The European Union grants a financial contribution to the Istituto Superiore di Sanità (ISS), Rome, Italy, to carry out the functions and duties provided for in Article 32 of Regulation (EC) No 882/2004, in respect of analysis and testing of parasites (in particular *Trichinella, Echinococcus* and *Anisakis*).

For the period from 1 January 2011 to 31 December 2011, that financial contribution shall not exceed EUR 325 010.

2. In addition to the maximum amount provided for in paragraph 1, the Union grants a financial contribution to the laboratory referred to in paragraph 1 for the organisation of workshops. That contribution shall not exceed EUR 30 000.

Article 10

1. The European Union grants a financial contribution to the Fødevareinstituttet, Danmarks Tekniske Universitet (DTU), Copenhagen, Denmark, to carry out the functions and duties provided for in Article 32 of Regulation (EC) No 882/2004, for the monitoring of antimicrobial resistance.

For the period from 1 January 2011 to 31 December 2011, that financial contribution shall not exceed EUR 387 534.

2. In addition to the maximum amount provided for in paragraph 1, the Union grants a financial contribution to the laboratory referred to in paragraph 1 for the organisation of workshops. That contribution shall not exceed EUR 67 000.

Article 11

1. The European Union grants a financial contribution to the Veterinary Laboratories Agency, Addlestone, United Kingdom, to carry out the functions and duties provided in Chapter B of Annex X to Regulation (EC) No 999/2001 of the European Parliament and of the Council [1], in particular for the monitoring of transmissible spongiform encephalopathies.

For the period from 1 January 2011 to 31 December 2011, that financial contribution shall not exceed EUR 737 901.

2. In addition to the maximum amount provided for in paragraph 1, the Union grants a financial contribution to the laboratory referred to in paragraph 1 for the organisation of workshops. That contribution shall not exceed EUR 70 200.

3. By way of derogation from Article 13(1) of Regulation (EC) No 1754/2006, the laboratory referred to in paragraph 1 shall be entitled to claim financial assistance for attendance by a maximum of 50 participants at one of its workshops referred to in paragraph 2 of this Article.

Article 12

1. The European Union grants a financial contribution to the Centre wallon de Recherches agronomiques (CRA-W), Gembloux, Belgium, to carry out the functions and duties provided for in Article 32 of Regulation (EC) No 882/2004, for the analysis and testing of animal proteins in feedingstuffs.

For the period from 1 January 2011 to 31 December 2011, that financial contribution shall not exceed EUR 581 716.

2. In addition to the maximum amount provided for in paragraph 1, the Union grants a financial contribution to the laboratory referred to in paragraph 1 for the organisation of workshops. That contribution shall not exceed EUR 30 000.

[1] OJ L 147, 31.5.2001, p. 1.

Article 13

1. The European Union grants financial aid to the Rijksinstituut voor Volksgezondheid en Milieu (RIVM), Bilthoven, the Netherlands, to carry out the functions and duties provided for in Article 32 of Regulation (EC) No 882/2004, for residues of certain substances listed in Annex I to Council Directive 96/23/EC [1] and referred to by Annex VII, Section I, point 12(a) of Regulation (EC) No 882/2004.

For the period from 1 January 2011 to 31 December 2011, that financial aid shall not exceed EUR 464 000.

2. In addition to the maximum amount provided for in paragraph 1, the Union grants financial aid to the laboratory referred to in paragraph 1 for the organisation of workshops. That aid shall not exceed EUR 25 000.

Article 14

1. The European Union grants financial aid to the Laboratoire d'études et de recherches sur les médicaments vétérinaires et les désinfectants de L'Agence nationale de sécurité sanitaire de l'alimentation, de l'environnement et du travail (Anses, ex-AFSSA), Fougères, France, to carry out the functions and duties provided for in Article 32 of Regulation (EC) No 882/2004, for residues of certain substances listed in Annex I to Directive 96/23/EC and referred to by Annex VII, Section I, point 12 (a) of Regulation (EC) No 882/2004.

For the period from 1 January 2011 to 31 December 2011, that financial aid shall not exceed EUR 464 000.

2. In addition to the maximum amount provided for in paragraph 1, the Union grants financial aid to by the laboratory referred to in paragraph 1 for the organisation of workshops. That aid shall not exceed EUR 25 000.

Article 15

1. The European Union grants financial aid to the Bundesamt für Verbraucherschutz und Lebensmittelsicherheit (BVL), Berlin, Germany, to carry out the functions and duties provided for in Article 32 of Regulation (EC) No 882/2004, for residues of certain substances listed in Annex I to Directive 96/23/EC and referred to by Annex VII, Section I, point 12(a) of Regulation (EC) No 882/2004.

For the period from 1 January 2011 to 31 December 2011, that financial aid shall not exceed EUR 464 000.

2. In addition to the maximum amount provided for in paragraph 1, the Union grants financial aid to the laboratory referred to in paragraph 1 for the organisation of workshops. That aid shall not exceed EUR 25 000.

Article 16

1. The European Union grants financial aid to the Istituto Superiore di Sanità (ISS), Rome, Italy, to carry out the functions and duties provided for in Article 32 of Regulation (EC) No 882/2004, for residues of certain substances listed in Annex I to Directive 96/23/EC and referred to by Annex VII, Section I, point 12(a) of Regulation (EC) No 882/2004.

For the period from 1 January 2011 to 31 December 2011, that financial aid shall not exceed EUR 283 000.

2. In addition to the maximum amount provided for in paragraph 1, the Union grants financial aid to the laboratory referred to in paragraph 1 for the organisation of workshops. That aid shall not exceed EUR 25 000.

Article 17

1. The European Union grants a financial contribution to the Chemisches und Veterinäruntersuchungsamt (CVUA) Freiburg, Germany, to carry out the functions and duties provided for in Article 32 of Regulation (EC) No 882/2004, for the analysis and testing of residues of pesticides in food of animal origin and commodities with high fat content.

For the period from 1 January 2011 to 31 December 2011, that financial contribution shall not exceed EUR 198 900.

2. In addition to the maximum amount provided for in paragraph 1, the Union grants financial aid to the laboratory referred to in paragraph 1 for the organisation of workshops. That aid shall not exceed EUR 110 000.

3. By way of derogation from Article 13(1) of Regulation (EC) No 1754/2006, the laboratory referred to in paragraph 1 shall be entitled to claim financial assistance for attendance by a maximum of 110 participants at one of its workshops referred to in paragraph 2 of this Article.

Article 18

The European Union grants a financial contribution to the Fødevareinstituttet, Danmarks Tekniske Universitet (DTU), Copenhagen, Denmark, to carry out the functions and duties provided for in Article 32 of Regulation (EC) No 882/2004, for the analysis and testing of residues of pesticides in cereals and feedingstuffs.

[1] OJ L 125, 23.5.1996, p. 10.

For the period from 1 January 2011 to 31 December 2011, that financial contribution shall not exceed EUR 198 900.

Article 19

1. The European Union grants a financial contribution to the Laboratorio Agrario de la Generalitat Valenciana (LAGV)/Grupo de Residuos de Plaguicidas de la Universidad de Almería (PRRG), Spain to carry out the functions and duties provided for in Article 32 of Regulation (EC) No 882/2004, for the analysis and testing of residues of pesticides in fruits and vegetables, including commodities with high water and high acid content.

For the period from 1 January 2011 to 31 December 2011, that financial contribution shall not exceed EUR 447 000.

2. In addition to the maximum amount provided for in paragraph 1, the Union grants a financial contribution to the laboratory referred to in paragraph 1 for the organisation of workshops. That contribution shall not exceed EUR 10 000.

Article 20

The European Union grants a financial contribution to the Chemisches und Veterinäruntersuchungsamt (CVUA), Stuttgart, Germany, to carry out the functions and duties provided for in Article 32 of Regulation (EC) No 882/2004, for the analysis and testing of residues of pesticides by single residue methods.

For the period from 1 January 2011 to 31 December 2011, that financial contribution shall not exceed EUR 365 000.

Article 21

1. The European Union grants a financial contribution to the Chemisches und Veterinäruntersuchungsamt (CVUA), Freiburg, Germany, to carry out by the functions and duties provided for in Article 32 of Regulation (EC) No 882/2004, for the analysis and testing of dioxins and PCBs in feed and food.

For the period from 1 January 2011 to 31 December 2011, that financial contribution shall not exceed EUR 470 000.

2. In addition to the maximum amount provided for in paragraph 1, the Union grants a financial contribution to the laboratory referred to in paragraph 1 for the organisation of workshops. That contribution shall not exceed EUR 55 000.

Article 22

The Union's financial contribution referred to in Articles 1 to 21 shall be at the rate of 100 % of eligible costs as defined in Regulation (EC) No 1754/2006.

Article 23

This Decision is addressed to the:

— for milk and milk products: Laboratoire d'études et de recherches sur la qualité des aliments et sur les procédés agroalimentaires (Lerqap), of the Agence nationale de sécurité sanitaire de l'alimentation, de l'environnement et du travail (Anses), 23 avenue du Général de Gaulle, 94700 Maisons-Alfort, France,

— for the analysis and testing of zoonoses (salmonella): Rijksinstituut voor Volksgezondheid en Milieu (RIVM), Postbus 1, Anthony van Leeuwenhoeklaan 9, 3720 BA Bilthoven, The Netherlands,

— for the monitoring of marine biotoxins: Laboratorio de Biotoxinas Marinas, Agencia Española de Seguridad Alimentaria y Nutrición (Ministerio de Sanidad y Política Social), Estación Maritima, s/n, 36200 Vigo, Spain,

— for monitoring the viral and bacteriological contamination of bivalve molluscs: Laboratory of the Centre for Environment, Fisheries and Aquaculture Science (CEFAS), Weymouth laboratory, Barrack Road, The Nothe, Weymouth, Dorset, DT4 8UB, United Kingdom,

— for *Listeria monocytogenes*: Laboratoire d'Etudes et de Recherches sur la Qualité des Aliments et sur les Procédés Agro-alimentaires (Lerqap), of the Agence nationale de sécurité sanitaire de l'alimentation, de l'environnement et du travail (Anses), 23 avenue du Général de Gaulle, 94700 Maisons-Alfort, France,

— for coagulase positive *Staphylococci*, including *Staphylococcus aureus*: Laboratoire d'études et de recherches sur la qualité des aliments et sur les procédés agro-alimentaires (Lerqap), of the Agence nationale de sécurité sanitaire de l'alimentation, de l'environnement et du travail (Anses), 23 avenue du Général de Gaulle, 94700 Maisons-Alfort, France,

— for *Escherichia coli*, including Verotoxigenic *E. Coli* (VTEC): Istituto Superiore di Sanità (ISS), Viale Regina Elena 299, 00161 Roma, Italy,

— for Campylobacter: Statens Veterinärmedicinska Anstalt (SVA), Ulls väg 2 B, 75189 Uppsala, Sweden,

— for parasites (in particular *Trichinella*, *Echinococcus* and *Anisakis*): Istituto Superiore di Sanità (ISS), Viale Regina Elena 299, 00161 Roma, Italy,

— for antimicrobial resistance: Fødevareinstituttet, Danmarks Tekniske Universitet (DTU), Bülowsvej 27, 1790 Copenhagen V, Denmark,

— for transmissible spongiform encephalopathies (TSEs): Veterinary Laboratories Agency, Woodham Lane, New Haw, Addlestone, Surrey KT15 3NB, United Kingdom (Ms Marion Simmons, tel. +441932357564),

— for animal proteins in feedingstuffs: Centre wallon de Recherches agronomiques (CRA-W), Chaussée de Namur 24, 5030 Gembloux, Belgium,

— for residues: Rijksinstituut voor Volksgezondheid en Milieu (RIVM), Postbus 1, Anthony van Leeuwenhoeklaan 9, 3720 BA Bilthoven, The Netherlands,

— for residues: Laboratoire d'études et de recherches sur les médicaments vétérinaires et les désinfectants de L'Agence nationale de sécurité sanitaire de l'alimentation, de l'environnement et du travail (Anses), Site de Fougères, BP 90203, 35302 Fougères, France,

— for residues: Bundesamt für Verbraucherschutz und Lebensmittelsicherheit, Postfach 100214, Mauerstrasse 39-42, 10562 Berlin, Germany,

— for residues: Istituto Superiore di Sanità (ISS), Viale Regina Elena 299, 00161 Roma, Italy,

— for the analysis and testing of residues of pesticides in food of animal origin: Chemisches und Veterinäruntersuchungsamt (CVUA), Postfach 100462, Bissierstrasse 5, 79114 Freiburg, Germany,

— for the analysis and testing of residues of pesticides in cereals: Fødevareinstituttet, Danmarks Tekniske Universitet (DTU), Department of Food Chemistry, Moerkhoej Bygade 19, 2860 Soeborg, Denmark,

— for the analysis and testing of residues of pesticides in fruits and vegetables: Laboratorio Agrario de la Generalitat Valenciana (LAGV)/Grupo de Residuos de Plaguicidas de la Universidad de Almería (PRRG), Ctra. Sacramento s/n, La Canada de San Urbano, 04120 Almeria, Spain,

— for the analysis and testing of residues of pesticides by single residue methods: Chemisches und Veterinäruntersuchungsamt (CVUA), Postfach 1206, Schaflandstrasse 3/2, 70736 Stuttgart, Germany,

— for the analysis and testing of dioxins and PCBs in feed and food: Chemisches und Veterinäruntersuchungsamt (CVUA), Postfach 100462, Bissierstrasse 5, 79114 Freiburg, Germany.

Done at Brussels, 1 December 2010.

For the Commission

John DALLI

Member of the Commission

CORRIGENDA

Corrigendum to Commission Decision 2010/432/EU of 28 July 2010 authorising the placing on the market of products containing, consisting of, or produced from genetically modified maize 1507x59122 (DAS-Ø15Ø7-1xDAS-59122-7) pursuant to Regulation (EC) No 1829/2003 of the European Parliament and of the Council

(Official Journal of the European Union L 202 of 4 August 2010)

On page 14, Annex, point (b):

for: 'The genetically modified DAS-Ø15Ø7-1xDAS-59122-7 maize, as described in the application, is produced by crosses between maize containing DAS-Ø15Ø7 and DAS-59122-7 events and expresses the Cry1F protein which confers protection against certain lepidopteran pests, the Cry34Ab1 and Cry35Ab1 proteins which confer protection against certain coleopteran pests and the PAT protein, used as a selectable marker, which confers tolerance to the glufosinate-ammonium herbicide.',

read: 'The genetically modified DAS-Ø15Ø7-1xDAS-59122-7 maize, as described in the application, is produced by crosses between maize containing DAS-Ø15Ø7-1 and DAS-59122-7 events and expresses the Cry1F protein which confers protection against certain lepidopteran pests, the Cry34Ab1 and Cry35Ab1 proteins which confer protection against certain coleopteran pests and the PAT protein, used as a selectable marker, which confers tolerance to the glufosinate-ammonium herbicide.';

on page 14, Annex, point (d), first indent:

for: 'event specific real-time quantitative PCR based methods for genetically modified maize DAS-Ø15Ø7 and DAS-59122-7 maize validated on DAS-Ø15Ø7-1xDAS-59122-7 maize,',

read: 'event specific real-time quantitative PCR based methods for genetically modified maize DAS-Ø15Ø7-1 and DAS-59122-7 maize validated on DAS-Ø15Ø7-1xDAS-59122-7 maize,';

on page 14, Annex, point (d), third indent:

for: 'reference material: ERM®-BF418 (for DAS-Ø15Ø7) and ERM®-BF424 (for DAS-59122-7) accessible via the Joint Research Centre (JRC) of the European Commission, Institute for Reference Materials and Measurements (IRMM) at https://irmm.jrc.ec.europa.eu/rmcatalogue',

read: 'reference material: ERM®-BF418 (for DAS-Ø15Ø7-1) and ERM®-BF424 (for DAS-59122-7) accessible via the Joint Research Centre (JRC) of the European Commission, Institute for Reference Materials and Measurements (IRMM) at https://irmm.jrc.ec.europa.eu/rmcatalogue'.

Corrigendum to Commission Decision 2010/428/EU of 28 July 2010 authorising the placing on the market of products containing, consisting of, or produced from genetically modified maize 59122x1507xNK603 (DAS-59122-7xDAS-Ø15Ø7xMON-ØØ6Ø3-6) pursuant to Regulation (EC) No 1829/2003 of the European Parliament and of the Council

(Official Journal of the European Union L 201 of 3 August 2010)

On page 41, title:

for: 'Commission Decision 2010/428/EU of 28 July 2010 authorising the placing on the market of products containing, consisting of, or produced from genetically modified maize 59122x1507xNK603 (DAS-59122-7xDAS-Ø15Ø7xMON-ØØ6Ø3-6) pursuant to Regulation (EC) No 1829/2003 of the European Parliament and of the Council',

read: 'Commission Decision 2010/428/EU of 28 July 2010 authorising the placing on the market of products containing, consisting of, or produced from genetically modified maize 59122x1507xNK603 (DAS-59122-7xDAS-Ø15Ø7-1xMON-ØØ6Ø3-6) pursuant to Regulation (EC) No 1829/2003 of the European Parliament and of the Council';

on page 42, Article 1:

for: 'Genetically modified maize (*Zea mays* L.) 59122x1507xNK603, as specified in point (b) of the Annex to this Decision, is assigned the unique identifier DAS-59122-7xDAS-Ø15Ø7xMON-ØØ6Ø3-6, as provided for in Regulation (EC) No 65/2004.',

read: 'Genetically modified maize (*Zea mays* L.) 59122x1507xNK603, as specified in point (b) of the Annex to this Decision, is assigned the unique identifier DAS-59122-7xDAS-Ø15Ø7-1xMON-ØØ6Ø3-6, as provided for in Regulation (EC) No 65/2004.';

on page 42, Article 2(a):

for: 'foods and food ingredients containing, consisting of, or produced from DAS-59122-7xDAS-Ø15Ø7xMON-ØØ6Ø3-6 maize;',

read: 'foods and food ingredients containing, consisting of, or produced from DAS-59122-7xDAS-Ø15Ø7-1xMON-ØØ6Ø3-6 maize;';

on page 42, Article 2(b):

for: 'feed containing, consisting of, or produced from DAS-59122-7xDAS-Ø15Ø7xMON-ØØ6Ø3-6 maize;',

read: 'feed containing, consisting of, or produced from DAS-59122-7xDAS-Ø15Ø7-1xMON-ØØ6Ø3-6 maize;';

on page 42, Article 2(c):

for: 'products other than food and feed containing or consisting of DAS-59122-7xDAS-Ø15Ø7xMON-ØØ6Ø3-6 maize for the same uses as any other maize with the exception of cultivation.',

read: 'products other than food and feed containing or consisting of DAS-59122-7xDAS-Ø15Ø7-1xMON-ØØ6Ø3-6 maize for the same uses as any other maize with the exception of cultivation.';

on page 42, Article 3(2):

for: 'The words "not for cultivation" shall appear on the label of and in documents accompanying products containing or consisting of DAS-59122-7xDAS-Ø15Ø7xMON-ØØ6Ø3-6 maize referred to in Article 2(b) and (c).',

read: 'The words "not for cultivation" shall appear on the label of and in documents accompanying products containing or consisting of DAS-59122-7xDAS-Ø15Ø7-1xMON-ØØ6Ø3-6 maize referred to in Article 2(b) and (c).';

on page 44, Annex, point (b)(1):

for: 'foods and food ingredients containing, consisting of, or produced from DAS-59122-7xDAS-Ø15Ø7xMON-ØØ6Ø3-6 maize',

read: 'foods and food ingredients containing, consisting of, or produced from DAS-59122-7xDAS-Ø15Ø7-1xMON-ØØ6Ø3-6 maize';

on page 44, Annex, point (b)(2):

for: 'feed containing, consisting of, or produced from DAS-59122-7xDAS-Ø15Ø7xMON-ØØ6Ø3-6 maize',

read: 'feed containing, consisting of, or produced from DAS-59122-7xDAS-Ø15Ø7-1xMON-ØØ6Ø3-6 maize';

on page 44, Annex, point (b)(3):

for: 'products other than food and feed containing or consisting of DAS-59122-7xDAS-Ø15Ø7xMON-ØØ6Ø3-6 maize for the same uses as any other maize with the exception of cultivation.',

read: 'products other than food and feed containing or consisting of DAS-59122-7xDAS-Ø15Ø7-1xMON-ØØ6Ø3-6 maize for the same uses as any other maize with the exception of cultivation.';

on page 44, Annex, point (b):

for: 'The genetically modified DAS-59122-7xDAS-Ø15Ø7xMON-ØØ6Ø3-6 maize, as described in the application, is produced by crosses between maize containing DAS-59122-7, DAS-Ø15Ø7 and MON-ØØ6Ø3-6 events and expresses the Cry34Ab1 and Cry35Ab1 proteins which confer protection against certain coleopteran pests, the Cry1F protein which confers protection against certain lepidopteran pests, the PAT protein, used as a selectable marker, which confers tolerance to the glufosinate-ammonium herbicide and the CP4 EPSPS protein which confers tolerance to glyphosate herbicide.',

read: 'The genetically modified DAS-59122-7xDAS-Ø15Ø7-1xMON-ØØ6Ø3-6 maize, as described in the application, is produced by crosses between maize containing DAS-59122-7, DAS-Ø15Ø7-1 and MON-ØØ6Ø3-6 events and expresses the Cry34Ab1 and Cry35Ab1 proteins which confer protection against certain coleopteran pests, the Cry1F protein which confers protection against certain lepidopteran pests, the PAT protein, used as a selectable marker, which confers tolerance to the glufosinate-ammonium herbicide and the CP4 EPSPS protein which confers tolerance to glyphosate herbicide.';

on page 44, Annex, point (c)(2):

for: 'the words "not for cultivation" shall appear on the label of and in documents accompanying products containing or consisting of DAS-59122-7xDAS-Ø15Ø7xMON-ØØ6Ø3-6 maize referred to in Article 2(b) and (c) of this Decision.',

read: 'the words "not for cultivation" shall appear on the label of and in documents accompanying products containing or consisting of DAS-59122-7xDAS-Ø15Ø7-1xMON-ØØ6Ø3-6 maize referred to in Article 2(b) and (c) of this Decision.';

on page 44, Annex, point (d), first indent:

for: 'event specific real-time quantitative PCR-based methods for genetically modified maize DAS-59122-7, DAS-Ø15Ø7 and MON-ØØ6Ø3-6 maize validated on DAS-59122-7xDAS-Ø15Ø7xMON-ØØ6Ø3-6 maize;',

read: 'event specific real-time quantitative PCR-based methods for genetically modified maize DAS-59122-7, DAS-Ø15Ø7-1 and MON-ØØ6Ø3-6 maize validated on DAS-59122-7xDAS-Ø15Ø7-1xMON-ØØ6Ø3-6 maize;';

on page 44, Annex, point (d), third indent:

for: 'reference material: ERM®-BF424 (for DAS-59122-7), ERM®-BF418 (for DAS-Ø15Ø7) and ERM®-BF415 (for MON-ØØ6Ø3-6) accessible via the Joint Research Centre (JRC) of the European Commission, Institute for Reference Materials and Measurements (IRMM) at https://irmm.jrc.ec.europa.eu/rmcatalogue',

read: 'reference material: ERM®-BF424 (for DAS-59122-7), ERM®-BF418 (for DAS-Ø15Ø7-1) and ERM®-BF415 (for MON-ØØ6Ø3-6) accessible via the Joint Research Centre (JRC) of the European Commission, Institute for Reference Materials and Measurements (IRMM) at https://irmm.jrc.ec.europa.eu/rmcatalogue';

on page 44, Annex, point (e):

for: 'DAS-59122-7xDAS-Ø15Ø7xMON-ØØ6Ø3-6.',

read: 'DAS-59122-7xDAS-Ø15Ø7-1xMON-ØØ6Ø3-6.'.

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