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Contents

I Acts adopted under the EC Treaty/Euratom Treaty whose publication is obligatory

REGULATIONS

- ★ **Council Regulation (EC) No 1134/2007 of 10 July 2007 amending Regulation (EC) No 2866/98 as regards the conversion rate to the euro for Malta** 1
- ★ **Council Regulation (EC) No 1135/2007 of 10 July 2007 amending Regulation (EC) No 2866/98 as regards the conversion rate to the euro for Cyprus** 2
- Commission Regulation (EC) No 1136/2007 of 1 October 2007 establishing the standard import values for determining the entry price of certain fruit and vegetables 3
- ★ **Commission Regulation (EC) No 1137/2007 of 1 October 2007 concerning the authorisation of *Bacillus subtilis* (O35) as a feed additive ⁽¹⁾** 5
- ★ **Commission Regulation (EC) No 1138/2007 of 1 October 2007 concerning the authorisation of a new use of benzoic acid (VevoVital) as a feed additive ⁽¹⁾** 8
- ★ **Commission Regulation (EC) No 1139/2007 of 1 October 2007 concerning the authorisation of L-arginine as a feed additive ⁽¹⁾** 11
- ★ **Commission Regulation (EC) No 1140/2007 of 1 October 2007 concerning the provisional authorisation of a new use of an additive already authorised in feedingstuffs ⁽¹⁾** 14
- ★ **Commission Regulation (EC) No 1141/2007 of 1 October 2007 concerning the authorisation of 3-phytase (ROVABIO PHY AP and ROVABIO PHY LC) as a feed additive ⁽¹⁾** 17
- ★ **Commission Regulation (EC) No 1142/2007 of 1 October 2007 concerning the authorisation of a new use of 3-phytase (Natuphos) as a feed additive ⁽¹⁾** 20

⁽¹⁾ Text with EEA relevance

(Continued overleaf)

★ Commission Regulation (EC) No 1143/2007 of 1 October 2007 amending Regulation (EC) No 256/2002 as regards the authorisation of the feed additive preparation of <i>Bacillus cereus</i> var. <i>toyoi</i> , belonging to the group of microorganisms ⁽¹⁾	23
★ Commission Regulation (EC) No 1144/2007 of 1 October 2007 amending Regulation (EC) No 1831/96 opening and providing for the administration of Community tariff quotas bound under GATT for certain fruit and vegetables and processed fruit and vegetable products from 1996	26

II Acts adopted under the EC Treaty/Euratom Treaty whose publication is not obligatory

DECISIONS

2007/633/EC:

★ Council Decision of 18 September 2007 on the conclusion of the Agreement between the European Community and New Zealand on certain aspects of air services	27
--	----

III Acts adopted under the EU Treaty

ACTS ADOPTED UNDER TITLE V OF THE EU TREATY

★ Council Joint Action 2007/634/CFSP of 1 October 2007 amending Joint Action 2007/113/CFSP amending and extending the mandate of the European Union Special Representative for Central Asia	28
★ Council common position 2007/635/CFSP of 1 October 2007 extending Common Position 2004/694/CFSP on further measures in support of the effective implementation of the mandate of the International Criminal Tribunal for the former Yugoslavia (ICTY)	30

Corrigenda

★ Corrigendum to Council Regulation (EC) No 1890/2005 of 14 November 2005 imposing a definitive anti-dumping duty and collecting definitively the provisional duty imposed on imports of certain stainless steel fasteners and parts thereof originating in the People's Republic of China, Indonesia, Taiwan, Thailand and Vietnam and terminating the proceeding on imports of certain stainless steel fasteners and parts thereof originating in Malaysia and the Philippines (OJ L 302, 19.11.2005)	31
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⁽¹⁾ Text with EEA relevance

I

(Acts adopted under the EC Treaty/Euratom Treaty whose publication is obligatory)

REGULATIONS

COUNCIL REGULATION (EC) No 1134/2007

of 10 July 2007

amending Regulation (EC) No 2866/98 as regards the conversion rate to the euro for Malta

THE COUNCIL OF THE EUROPEAN UNION,

Having regard to the Treaty establishing the European Community, and in particular Article 123(5) thereof,

Having regard to the proposal from the Commission,

Having regard to the opinion of the European Central Bank ⁽¹⁾,

Whereas:

- (1) Council Regulation (EC) No 2866/98 of 31 December 1998 on the conversion rates between the euro and the currencies of the Member States adopting the euro ⁽²⁾ determines the conversion rates as from 1 January 1999.
- (2) According to Article 4 of the 2003 Act of Accession, Malta is a Member State with a derogation as defined in Article 122 of the Treaty.
- (3) Pursuant to Council Decision 2007/504/EC of 10 July 2007 in accordance with Article 122(2) of the Treaty on the adoption by Malta of the single currency on 1 January 2008 ⁽³⁾, Malta fulfils the necessary conditions for the adoption of the single currency and the dero-

gation of Malta is abrogated with effect from 1 January 2008.

- (4) The introduction of the euro in Malta requires the adoption of the conversion rate between the euro and the Maltese lira. This conversion rate should be set at 0,4293 lira per 1 euro, which corresponds to the current central rate of the lira in the exchange rate mechanism (ERM II).

- (5) Regulation (EC) No 2866/98 should therefore be amended accordingly,

HAS ADOPTED THIS REGULATION:

Article 1

In Article 1 of Regulation (EC) No 2866/98, the following line shall be inserted between the conversion rates applicable to the Luxembourg franc and the Dutch guilder:

‘= 0,429300 Maltese liras’.

Article 2

This Regulation shall enter into force on 1 January 2008.

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

Done at Brussels, 10 July 2007.

For the Council

The President

F. TEIXEIRA DOS SANTOS

⁽¹⁾ OJ C 160, 13.7.2007, p. 1.

⁽²⁾ OJ L 359, 31.12.1998, p. 1. Regulation as last amended by Regulation (EC) No 1086/2006 (OJ L 195, 15.7.2006, p. 1).

⁽³⁾ OJ L 186, 18.7.2007, p. 32.

COUNCIL REGULATION (EC) No 1135/2007
of 10 July 2007
amending Regulation (EC) No 2866/98 as regards the conversion rate to the euro for Cyprus

THE COUNCIL OF THE EUROPEAN UNION,

Having regard to the Treaty establishing the European Community, and in particular Article 123(5) thereof,

Having regard to the proposal from the Commission,

Having regard to the opinion of the European Central Bank ⁽¹⁾,

Whereas:

- (1) Council Regulation (EC) No 2866/98 of 31 December 1998 on the conversion rates between the euro and the currencies of the Member States adopting the euro ⁽²⁾ determines the conversion rates as from 1 January 1999.
- (2) According to Article 4 of the 2003 Act of Accession, Cyprus is a Member State with a derogation as defined in Article 122 of the Treaty.
- (3) Pursuant to Council Decision 2007/503/EC of 10 July 2007 in accordance with Article 122(2) of the Treaty on the adoption by Cyprus of the single currency on 1 January 2008 ⁽³⁾, Cyprus fulfils the necessary conditions for the adoption of the single currency and the deroga-

tion of Cyprus is abrogated with effect from 1 January 2008.

- (4) The introduction of the euro in Cyprus requires the adoption of the conversion rate between the euro and the Cyprus pound. This conversion rate should be set at 0,585274 pound per 1 euro, which corresponds to the current central rate of the pound in the exchange rate mechanism (ERM II).
- (5) Regulation (EC) No 2866/98 should therefore be amended accordingly,

HAS ADOPTED THIS REGULATION:

Article 1

In Article 1 of Regulation (EC) No 2866/98, the following line shall be inserted between the conversion rates applicable to the Italian lira and the Luxembourg franc:

‘= 0,585274 Cyprus pounds’.

Article 2

This Regulation shall enter into force on 1 January 2008.

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

Done at Brussels, 10 July 2007.

For the Council

The President

F. TEIXEIRA DOS SANTOS

⁽¹⁾ OJ C 160, 13.7.2007, p. 1.

⁽²⁾ OJ L 359, 31.12.1998, p. 1. Regulation as last amended by Regulation (EC) No 1086/2006 (OJ L 195, 15.7.2006, p. 1).

⁽³⁾ OJ L 186, 18.7.2007, p. 29.

COMMISSION REGULATION (EC) No 1136/2007**of 1 October 2007****establishing the standard import values for determining the entry price of certain fruit and vegetables**

THE COMMISSION OF THE EUROPEAN COMMUNITIES,

Having regard to the Treaty establishing the European Community,

Having regard to Commission Regulation (EC) No 3223/94 of 21 December 1994 on detailed rules for the application of the import arrangements for fruit and vegetables ⁽¹⁾, and in particular Article 4(1) thereof,

Whereas:

- (1) Regulation (EC) No 3223/94 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 the Annex thereto.

- (2) In compliance with the above criteria, the standard import values must be fixed at the levels set out in the Annex to this Regulation,

HAS ADOPTED THIS REGULATION:

Article 1

The standard import values referred to in Article 4 of Regulation (EC) No 3223/94 shall be fixed as indicated in the Annex hereto.

Article 2

This Regulation shall enter into force on 2 October 2007.

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

Done at Brussels, 1 October 2007.

For the Commission

Jean-Luc DEMARTY

*Director-General for Agriculture and
Rural Development*

⁽¹⁾ OJ L 337, 24.12.1994, p. 66. Regulation as last amended by Regulation (EC) No 756/2007 (OJ L 172, 30.6.2007, p. 41).

ANNEX

to Commission Regulation of 1 October 2007 establishing the 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	MK	49,9
	TR	109,3
	XS	28,3
	ZZ	62,5
0707 00 05	JO	151,2
	TR	61,3
	ZZ	106,3
0709 90 70	TR	116,7
	ZZ	116,7
0805 50 10	AR	73,1
	TR	99,4
	UY	82,6
	ZA	65,1
	ZZ	80,1
0806 10 10	BR	253,6
	IL	284,6
	MK	11,8
	TR	119,6
	US	210,4
	ZZ	176,0
0808 10 80	AR	87,7
	AU	165,9
	BR	45,1
	CL	73,7
	CN	79,8
	NZ	93,9
	US	99,9
	ZA	77,4
	ZZ	90,4
0808 20 50	CN	63,8
	TR	121,2
	ZA	77,8
	ZZ	87,6

⁽¹⁾ Country nomenclature as fixed by Commission Regulation (EC) No 1833/2006 (OJ L 354, 14.12.2006, p. 19). Code 'ZZ' stands for 'of other origin'.

COMMISSION REGULATION (EC) No 1137/2007
of 1 October 2007
concerning the authorisation of *Bacillus subtilis* (O35) as a feed additive
(Text with EEA relevance)

THE COMMISSION OF THE EUROPEAN COMMUNITIES,

Having regard to the Treaty establishing the European Community,

Having regard to Regulation (EC) No 1831/2003 of the European Parliament and of the Council of 22 September 2003 on additives for use in animal nutrition⁽¹⁾, and in particular Article 9(2) thereof,

Whereas:

- (1) Regulation (EC) No 1831/2003 provides for the authorisation of additives for use in animal nutrition and for the grounds and procedures for granting such authorisation.
- (2) In accordance with Article 7 of Regulation (EC) No 1831/2003, an application was submitted for the authorisation of the preparation set out in the Annex to this Regulation. That application was accompanied by the particulars and documents required under Article 7(3) of Regulation (EC) No 1831/2003.
- (3) The application concerns the authorisation of the preparation of *Bacillus subtilis* DSM 17299 (O35) as a feed additive for chickens for fattening to be classified in the additive category 'zootechnical additives'.
- (4) The European Food Safety Authority (the Authority) concluded in its opinion of 17 October 2006 that the preparation of *Bacillus subtilis* DSM 17299 (O35) does not have an adverse effect on animal health, human health or the environment⁽²⁾. It further concluded that that preparation does not present any other risk which

would, in accordance with Article 5(2) of Regulation (EC) No 1831/2003, exclude authorisation. The opinion of the Authority recommends appropriate measures for user safety. It does not consider that there is a need for specific requirements of post market monitoring. This opinion also verifies the report on the method of analysis of the feed additive in feed submitted by the Community Reference Laboratory set up by Regulation (EC) No 1831/2003.

- (5) The assessment of that preparation shows that the conditions for authorisation, provided for in Article 5 of Regulation (EC) No 1831/2003, are satisfied. Accordingly, the use of that preparation should be authorised, as specified in the Annex to this Regulation.
- (6) The measures provided for in this Regulation are in accordance with the opinion of the Standing Committee on the Food Chain and Animal Health,

HAS ADOPTED THIS REGULATION:

Article 1

The preparation specified in the Annex, belonging to the additive category 'zootechnical additives' and to the functional group 'gut flora stabilisers', is authorised as an additive in animal nutrition subject to the conditions laid down in that Annex.

Article 2

This Regulation shall enter into force on the 20th day following its publication in the *Official Journal of the European Union*.

⁽¹⁾ OJ L 268, 18.10.2003, p. 29. Regulation as amended by Commission Regulation (EC) No 378/2005 (OJ L 59, 5.3.2005, p. 8).

⁽²⁾ Opinion of the Scientific Panel on Additives and Products or Substances used in Animal Feed and the Scientific Panel on Genetically Modified Organisms on the safety and efficacy of the microbiological product 'O35', a preparation of *Bacillus subtilis*, as a feed additive for chickens for fattening in accordance with Regulation (EC) No 1831/2003. Adopted on 17 October 2006. *The EFSA Journal* (2006) 406, p. 1-11.

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

Done at Brussels, 1 October 2007.

For the Commission
Markos KYPRIANOU
Member of the Commission

ANNEX

Identification number of the additive	Name of the holder of authorisation	Additive (Trade name)	Composition, chemical formula, description, analytical method	Species or category of animal	Maximum age	Minimum content		Maximum content	Other provisions	End of period of authorisation
						CFU/kg of complete feedingstuff with a moisture content of 12 %				
Category of zootechnical additives. Functional group: gut flora stabilisers										
4b1821	Chr. Hansen A/S	<i>Bacillus subtilis</i> DSM 17299 (O35)	Additive composition: Preparation of <i>Bacillus subtilis</i> DSM 17299 Containing a minimum of $1,6 \times 10^9$ CFU/g of additive Characterisation of the active substance: <i>Bacillus subtilis</i> DSM 17299 spore concentrate Analytical method ⁽¹⁾ : Enumeration spread plate method using tryptone soya agar with pre-heat treatment of feed samples	Chickens for fattening	—	8×10^8		$1,6 \times 10^9$	1. In the directions for use of the additive and premixture, indicate the storage temperature, storage life and stability to pelleting. 2. The use is permitted in feed containing the permitted coccidiostats: diclazuril, halofuginone and robenidine.	22 October 2017

⁽¹⁾ Details of the analytical methods are available at the following address of the Community Reference Laboratory: www.irmm.jrc.be/crl-feed-additives

COMMISSION REGULATION (EC) No 1138/2007**of 1 October 2007****concerning the authorisation of a new use of benzoic acid (VevoVital) as a feed additive****(Text with EEA relevance)**

THE COMMISSION OF THE EUROPEAN COMMUNITIES,

Having regard to the Treaty establishing the European Community,

Having regard to Regulation (EC) No 1831/2003 of the European Parliament and of the Council of 22 September 2003 on additives for use in animal nutrition⁽¹⁾, and in particular Article 9(2) thereof,

Whereas:

(1) Regulation (EC) No 1831/2003 provides for the authorisation of additives for use in animal nutrition and for the grounds and procedures for granting such authorisation.

(2) In accordance with Article 7 of Regulation (EC) No 1831/2003, an application was submitted for the authorisation of the preparation set out in the Annex to this Regulation. That application was accompanied by the particulars and documents required under Article 7(3) of Regulation (EC) No 1831/2003.

(3) The application concerns the authorisation of a new use the preparation benzoic acid (VevoVital) as a feed additive for pigs for fattening, to be classified in the additive category 'zootechnical additives'.

(4) The use of the preparation of benzoic acid (VevoVital) was authorised for weaned piglets by Commission Regulation (EC) No 1730/2006⁽²⁾.

(5) New data were submitted in support of an application for authorisation for pigs for fattening. The European

Food Safety Authority (the Authority) concluded in its opinion of 7 March 2007 that benzoic acid (VevoVital) does not have an adverse effect on animal health, human health or the environment⁽³⁾. It further concluded that benzoic acid (VevoVital) does not present any other risk which would, in accordance with Article 5(2) of Regulation (EC) No 1831/2003, exclude authorisation. According to that opinion, the use of that preparation does not have an adverse effect on this additional animal. The opinion of the Authority recommends appropriate measures for user safety. It does not consider that there is a need for specific requirements of post market monitoring. This opinion also verifies the report on the method of analysis of the feed additive in feed submitted by the Community Reference Laboratory set up by Regulation (EC) No 1831/2003.

(6) The assessment of that preparation shows that the conditions for authorisation, provided for in Article 5 of Regulation (EC) No 1831/2003, are satisfied. Accordingly, the use of that preparation should be authorised, as specified in the Annex to this Regulation.

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

HAS ADOPTED THIS REGULATION:

Article 1

The preparation specified in the Annex, belonging to the additive category 'zootechnical additives' and to the functional group 'other zootechnical additives', is authorised as an additive in animal nutrition subject to the conditions laid down in that Annex.

Article 2

This Regulation shall enter into force on the 20th day following that of its publication in the *Official Journal of the European Union*.

⁽¹⁾ OJ L 268, 18.10.2003, p. 29. Regulation as amended by Commission Regulation (EC) No 378/2005 (OJ L 59, 5.3.2005, p. 8).

⁽²⁾ OJ L 325, 24.11.2006, p. 9.

⁽³⁾ Opinion of the Scientific Panel on Additives and Products or Substances used in Animal Feed on the safety and efficacy of the product VevoVital (benzoic acid) as a feed additive for pigs for fattening in accordance with Regulation (EC) No 1831/2003. Adopted on 7 March 2007. The EFSA Journal (2007) 457, p. 1 to 14.

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

Done at Brussels, 1 October 2007.

For the Commission
Markos KYPRIANOU
Member of the Commission

COMMISSION REGULATION (EC) No 1139/2007
of 1 October 2007
concerning the authorisation of L-arginine as a feed additive
(Text with EEA relevance)

THE COMMISSION OF THE EUROPEAN COMMUNITIES,

Having regard to the Treaty establishing the European Community,

Having regard to Regulation (EC) No 1831/2003 of the European Parliament and of the Council of 22 September 2003 on additives for use in animal nutrition ⁽¹⁾, and in particular Article 9(2) thereof,

Whereas:

(1) Regulation (EC) No 1831/2003 provides for the authorisation of additives for use in animal nutrition and for the grounds and procedures for granting such authorisation.

(2) An application for authorisation has been submitted for L-arginine as an aminoacid.

(3) As the application for authorisation was submitted before the date of application of Regulation (EC) No 1831/2003, it was submitted under Council Directive 82/471/EEC of 30 June 1982 concerning certain products used in animal nutrition ⁽²⁾. As from 18 October 2004, aminoacids, their salts and analogues fall under the scope of Regulation (EC) No 1831/2003. The application is therefore to be considered as an application under Article 7 of Regulation (EC) No 1831/2003.

(4) In order to comply with the requirements laid down in Article 7 of Regulation (EC) No 1831/2003, additional information was submitted in support of the application.

(5) The application concerns authorisation of L-arginine as a feed additive for all animal species, to be classified in the additive category 'nutritional additives' and the functional group 'aminoacids, their salts and analogues'.

(6) The European Food Safety Authority (the Authority) concluded in its opinions of 17 April 2007 ⁽³⁾ that L-arginine does not have an adverse effect on animal health, human health or the environment. It further concluded that L-arginine does not present any other risk which would, in accordance with Article 5(2) of Regulation (EC) No 1831/2003, exclude authorisation. It does not consider that there is a need for specific requirements of post market monitoring. The report on the method of analysis of the feed additive in feed was submitted by the Community Reference Laboratory set up by Regulation (EC) No 1831/2003 to the Authority. The assessment of that preparation shows that the conditions for authorisation, provided for in Article 5 of Regulation (EC) No 1831/2003, are satisfied. Accordingly, the use of that preparation should be authorised, as specified in the Annex to this Regulation.

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

HAS ADOPTED THIS REGULATION:

Article 1

The preparation specified in the Annex, belonging to the additive category 'nutritional additives' and to the functional group 'aminoacids, their salts and analogues', is authorised as an additive in animal nutrition subject to the conditions laid down in that Annex.

Article 2

This Regulation shall enter into force on the 20th day following its publication in the *Official Journal of the European Union*.

⁽¹⁾ OJ L 268, 18.10.2003, p. 29. Regulation as amended by Commission Regulation (EC) No 378/2005 (OJ L 59, 5.3.2005, p. 8).

⁽²⁾ OJ L 213, 21.7.1982, p. 8. Directive as last amended by Commission Directive 2004/116/EC (OJ L 379, 24.12.2004, p. 81).

⁽³⁾ Opinion of the Scientific Panel on Additives and Products or Substances used in Animal Feed on the safety and efficacy of the product L-arginine produced by fermentation from *Corynebacterium glutamicum* (ATCC-13870) for all animal species. Adopted on 17 April 2007. The EFSA Journal (2007) 473, p. 1-19.

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

Done at Brussels, 1 October 2007.

For the Commission
Markos KYPRIANOU
Member of the Commission

ANNEX

Identification number of the additive	Name of the holder of the authorisation	Additive	Composition, chemical formula, description, analytical method	Species or category of animal	Maximum age	Minimum content	Maximum content	Other provisions	End of period of authorisation
						mg/kg of complete feedstuff with a moisture content of 12 %	Maximum content of 12 %		
Category of nutritional additives. Functional group: amino acids, their salts and analogues									
3c3.6.1	—	L-arginine	Characterisation of the additive: L-arginine 98 % produced by <i>Corynebacterium glutamicum</i> (ATCC 13870) C₆H₁₄N₄O₂ Analytical method: Community method for the determination of amino acids (Commission Directive 98/64/EC amending Directive 71/393/EEC ⁽¹⁾)	All species	—	—	—	—	22.10.2017

⁽¹⁾ OJ L 257, 19.9.1998, p. 14.

(1) OJ L 257, 19.9.1998, p. 14.

COMMISSION REGULATION (EC) No 1140/2007**of 1 October 2007****concerning the provisional authorisation of a new use of an additive already authorised in feedingstuffs****(Text with EEA relevance)**

THE COMMISSION OF THE EUROPEAN COMMUNITIES,

Having regard to the Treaty establishing the European Community,

Having regard to Council Directive 70/524/EEC of 23 November 1970 concerning additives in feedingstuffs ⁽¹⁾, and in particular Articles 3, and 9e(1) thereof,

Having regard to Regulation (EC) No 1831/2003 of the European Parliament and of the Council of 22 September 2003 on additives for use in animal nutrition ⁽²⁾, and in particular Article 25 thereof,

Whereas:

(1) Regulation (EC) No 1831/2003 provides for the authorisation of additives for use in animal nutrition.

(2) Article 25 of Regulation (EC) No 1831/2003 lays down transitional measures for applications for the authorisation of feed additives submitted in accordance with Directive 70/524/EEC before the date of application of Regulation (EC) No 1831/2003.

(3) The application for the authorisation of a new use of the additive listed in the Annex to this Regulation was submitted before the date of application of Regulation (EC) No 1831/2003.

⁽¹⁾ OJ L 270, 14.12.1970, p. 1. Directive as last amended by Commission Regulation (EC) No 1800/2004 (OJ L 317, 16.10.2004, p. 37).

⁽²⁾ OJ L 268, 18.10.2003, p. 29. Regulation as amended by Commission Regulation (EC) No 378/2005 (OJ L 59, 5.3.2005, p. 8).

(4) Initial comments on this application, as provided for in Article 4(4) of Directive 70/524/EEC, were forwarded to the Commission before the date of application of Regulation (EC) No 1831/2003. This application is therefore to continue to be treated in accordance with Article 4 of Directive 70/524/EEC.

(5) The use of the enzyme preparation endo-1,3(4)-beta-glucanase produced by *Aspergillus aculeatus* (CBS 589.94), endo-1,4-beta-glucanase produced by *Trichoderma longibrachiatum* (CBS 592.94), alpha-amylase produced by *Bacillus amyloliquefaciens* (DSM 9553), bacil-lolysin produced by *Bacillus amyloliquefaciens* (DSM 9554) and endo-1,4-beta-xylanase produced by *Trichoderma viride* (NIBH FERM BP 4842) was authorised without a time limit for chickens for fattening by Commission Regulation (EC) No 358/2005 ⁽³⁾.

(6) New data were submitted in support of an application for authorisation for four years of that preparation for laying hens. The European Food Safety Authority (EFSA) delivered its opinion on that use on 18 April 2007. The assessment shows that the conditions laid down in Article 9e(1) of Directive 70/524/EEC for such authorisation are satisfied. Accordingly, the use of that preparation, as specified in Annex to this Regulation, should be provisionally authorised for four years.

(7) The assessment of this application shows that certain procedures should be required to protect workers from exposure to the additives set out in the Annexes. Such protection should be assured by the application of Council Directive 89/391/EEC of 12 June 1989 on the introduction of measures to encourage improvements in the safety and health of workers at work ⁽⁴⁾.

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

⁽³⁾ OJ L 57, 3.3.2005, p. 3.

⁽⁴⁾ OJ L 183, 29.6.1989, p. 1. Directive as last amended by Directive 2007/30/EC of the European Parliament and of the Council (OJ L 165, 27.6.2007, p. 21).

HAS ADOPTED THIS REGULATION:

Article 1

The preparation belonging to the group 'Enzymes', as specified in the Annex, is authorised provisionally for four years as an additive in animal nutrition under the conditions laid down in that Annex.

Article 2

This Regulation shall enter into force on the 20th 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, 1 October 2007.

For the Commission

Markos KYPRIANOU

Member of the Commission

ANNEX

EC No	Additive	Chemical formula, description	Species or category of animal	Maximum age	Minimum content		Maximum content	Other provisions	End of period of authorisation
					Unit of activity/kg of complete feedingstuff				
Enzymes									
53	Endo-1,3(4)-beta-glucanase EC 3.2.1.6 Endo-1,4-beta-glucanase EC 3.2.1.4 Alpha-amylase EC 3.2.1.1 Bacillolysin EC 3.4.24.28 Endo-1,4-beta-xylanase EC 3.2.1.8	Preparation of endo-1,3(4)-beta-glucanase produced by <i>Aspergillus aculeatus</i> (CBS 589/94), endo-1,4-beta-glucanase produced by <i>Trichoderma longibrachiatum</i> (CBS 592/94), alpha-amylase produced by <i>Bacillus amyloliquefaciens</i> (DSM 9553), bacillolysin produced by <i>Bacillus amyloliquefaciens</i> (DSM 9554) and endo-1,4-beta-xylanase produced by <i>Trichoderma viride</i> (NIBH FERM BP 4842) having a minimum activity of: Endo-1,3(4)-beta-glucanase: 2 350 U/g ⁽¹⁾ Endo-1,4-beta-glucanase: 4 000 U/g ⁽²⁾ Alpha-amylase: 400 U/g ⁽³⁾ Bacillolysin: 450 U/g ⁽⁴⁾ Endo-1,4-beta-xylanase: 20 000 U/g ⁽⁵⁾	Laying hens	—	Endo-1,3(4)-beta-glucanase: 587 Endo-1,4-beta-glucanase: 1 000 U Alpha-amylase: 100 U Bacillolysin: 112 U Endo-1,4-beta-xylanase: 5 000 U	—	1. In the directions for use of the additive and premixture, indicate the storage temperature, storage life, and stability to pelleting. 2. Recommended dose per kilogram of complete feedingsstuff: — endo-1,3(4)-beta-glucanase: 587-2 350 U — endo-1,4-beta-glucanase: 1 000-4 000 U — alpha-amylase: 100-400 U — bacillolysin: 112-450 U — endo-1,4-beta-xylanase: 5 000-20 000 U 3. For use in compound feed rich in non-starch polysaccharides (mainly beta-glucans and especially arabinoxylans), e.g. containing more than 30 % of wheat.	22 October 2011	
				</					

⁽¹⁾ 1 U is the amount of enzyme which liberates 0,0056 micromoles of reducing sugars (glucose equivalents) from barley beta-glucan per minute at pH 7,5 and 30 °C.

⁽²⁾ 1 U is the amount of enzyme which liberates 0,0056 micromoles of reducing sugars (glucose equivalents) from carboxymethylcellulose per minute at pH 4,8 and 50 °C.

⁽³⁾ 1 U is the amount of enzyme which liberates 1 micromole of glucose from a cross-linked starch polymer per minute at pH 7,5 and 37 °C.

⁽⁴⁾ 1 U is the amount of enzyme which solubilises one microgram of azo-casein substrate per minute at pH 7,5 and 37 °C.

⁽⁵⁾ 1 U is the amount of enzyme which liberates 0,0067 micromoles of reducing sugars (xylose equivalents) from birchwood xylan per minute at pH 5,3 and 50° C.

COMMISSION REGULATION (EC) No 1141/2007**of 1 October 2007****concerning the authorisation of 3-phytase (ROVABIO PHY AP and ROVABIO PHY LC) as a feed additive****(Text with EEA relevance)**

THE COMMISSION OF THE EUROPEAN COMMUNITIES,

Having regard to the Treaty establishing the European Community,

Having regard to Regulation (EC) No 1831/2003 of the European Parliament and of the Council of 22 September 2003 on additives for use in animal nutrition⁽¹⁾, and in particular Article 9(2) thereof,

Whereas:

- (1) Regulation (EC) No 1831/2003 provides for the authorisation of additives for use in animal nutrition and for the grounds and procedures for granting such authorisation.
- (2) In accordance with Article 7 of Regulation (EC) No 1831/2003, an application was submitted for the authorisation of the preparation set out in the Annex to this Regulation. That application was accompanied by the particulars and documents required under Article 7(3) of Regulation (EC) No 1831/2003.
- (3) The application concerns the authorisation of the preparation of 3-phytase produced by *Penicillium funiculosum* (CBS 111 433) (ROVABIO PHY AP and ROVABIO PHY LC), as a feed additive for chickens for fattening, laying hens, piglets (weaned) and pigs for fattening to be classified in the additive category 'zootechnical additives'.
- (4) The European Food Safety Authority (the Authority) concluded in its opinion of 17 April 2007 and 22 March 2007 that the preparation 3-phytase produced by *Penicillium funiculosum* (CBS 111 433) (ROVABIO

PHY AP and ROVABIO PHY LC) does not have an adverse effect on animal health, human health or the environment⁽²⁾. It further concluded that that preparation does not present any other risk which would, in accordance with Article 5(2) of Regulation (EC) No 1831/2003, exclude authorisation. The opinion of the Authority recommends appropriate measures for user safety. It does not consider that there is a need for specific requirements of post market monitoring. This opinion also verifies the report on the method of analysis of the feed additive submitted by the Community Reference Laboratory set up by Regulation (EC) No 1831/2003.

- (5) The assessment of that preparation shows that the conditions for authorisation, provided for in Article 5 of Regulation (EC) No 1831/2003, are satisfied. Accordingly, the use of that preparation should be authorised, as specified in the Annex to this Regulation.
- (6) The measures provided for in this Regulation are in accordance with the opinion of the Standing Committee on the Food Chain and Animal Health,

HAS ADOPTED THIS REGULATION:

Article 1

The preparation specified in the Annex, belonging to the additive category 'zootechnical additives' and to the functional group 'digestibility enhancers', is authorised as an additive in animal nutrition subject to the conditions laid down in that Annex.

Article 2

This Regulation shall enter into force on the 20th day following its publication in the *Official Journal of the European Union*.

⁽¹⁾ OJ L 268, 18.10.2003, p. 29. Regulation as amended by Commission Regulation (EC) No 378/2005 (OJ L 59, 5.3.2005, p. 8).

⁽²⁾ Opinion of the Scientific Panel on Additives and Products or Substances used in Animal Feed and the Scientific Panel on Genetically Modified Organisms on the safety and efficacy of the enzyme preparation of 3-phytase produced by *Penicillium funiculosum* (CBS 111 433) (ROVABIO PHY AP and ROVABIO PHY LC), as a feed additive for chickens for fattening, laying hens, piglets (weaned) and pigs for fattening in accordance with Regulation (EC) No 1831/2003. Adopted on 17 April 2007 and 22 March 2007. *The EFSA Journal* (2007) 471, 1 to 29.

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

Done at Brussels, 1 October 2007.

For the Commission
Markos KYPRIANOU
Member of the Commission

ANNEX

Identification number of the additive	Name of the holder of authorisation	Additive (Trade name)	Composition, chemical formula, description, analytical method	Species or category of animal	Maximum age	Minimum content		Maximum content	Other provisions	End of period of authorisation
						Units of activity/kg of complete feedingstuff with a moisture content of 12 %				
Category of zootechnical additives. Functional group: Digestibility enhancers										
4a1	Adisseo	3-phytase EC 3.1.3.8 (ROVABIO PHY AP and ROVABIO PHY LC)	Additive composition: Preparation of 3-phytase produced by <i>Penicillium funiculosum</i> (CBS 111 433) having a minimum activity of: Solid form: 2 500 RPU ⁽¹⁾ /g Liquid form: 1 000 RPU/ml Characterisation of the active substance: 3-phytase produced by <i>Penicillium funiculosum</i> (CBS 111 433) Analytical methods ⁽²⁾ Colorimetric method measuring inorganic phosphate released by the enzyme from phytate substrate.	Chickens for fattening	—	350 RPU			1. In the directions for use of the additive and premixture, indicate the storage temperature, storage life, and stability to pelleting. 2. For use in feed containing more than 0,23 % phytin bound phosphorus. 3. For piglets (weaned) up to 35 kg of body weight. 4. For user safety: breathing protection, safety glasses and gloves should be worn during handling. 5. Recommended doses for kilogram of complete feedingsuffs: — chickens for fattening: 350-500 RPU, 350-500 RPU, — laying hens: 300-500 RPU, 300-500 RPU, — piglets (weaned): 250-500 RPU, 250-500 RPU, — pigs for fattening: 350-500 RPU.	22.10.2017
				Laying hens	—	300 RPU				
				Piglets (weaned)	—	250 RPU				
				Pigs for fattening	—	350 RPU				

(¹) 1 RPU is the amount of enzyme that releases 1 micromole inorganic phosphate per minute from sodium phytate as substrate under defined conditions (pH 5,5 and 37 °C).

(²) Details of the analytical methods are available at the following address of the Community Reference Laboratory: www.irmm.jrc.be/crl-feed-additives

⁽¹⁾ 1 RPU is the amount of enzyme that releases 1 micromole inorganic phosphate per minute from sodium phytate as substrate under defined conditions (pH 5,5 and 37 °C).

⁽²⁾ Details of the analytical methods are available at the following address of the Community Reference Laboratory: www.irmm.jrc.be/crl-feed-additives

COMMISSION REGULATION (EC) No 1142/2007**of 1 October 2007****concerning the authorisation of a new use of 3-phytase (Natuphos) as a feed additive****(Text with EEA relevance)**

THE COMMISSION OF THE EUROPEAN COMMUNITIES,

Having regard to the Treaty establishing the European Community,

Having regard to Regulation (EC) No 1831/2003 of the European Parliament and of the Council of 22 September 2003 on additives for use in animal nutrition⁽¹⁾, and in particular Article 9(2) thereof,

Whereas:

(1) Regulation (EC) No 1831/2003 provides for the authorisation of additives for use in animal nutrition and for the grounds and procedures for granting such authorisation.

(2) In accordance with Article 7 of Regulation (EC) No 1831/2003, an application was submitted for the authorisation of the preparation set out in the Annex to this Regulation. That application was accompanied by the particulars and documents required under Article 7(3) of Regulation (EC) No 1831/2003.

(3) The application concerns authorisation of a new use of the enzyme preparation 3-phytase (Natuphos 5 000, Natuphos 5 000 G, Natuphos 5 000 L, Natuphos 10 000 G and Natuphos 10 000 L) produced by *Aspergillus niger* (CBS 101 672) for laying hens and turkeys for fattening to be classified in the additive category 'zootechnical additives'.

(4) The use of that enzyme preparation was authorised for weaned piglets, pigs for fattening and chickens for fattening by Commission Regulation (EC) No 243/2007⁽²⁾.

(5) The European Food Safety Authority (the Authority) concluded in its opinion of 17 April 2007 that the enzyme preparation 3-phytase (Natuphos 5 000, Natuphos 5 000 G, Natuphos 5 000 L, Natuphos 10 000 G and Natuphos 10 000 L) produced by *Aspergillus niger* (CBS 101 672) does not have an adverse effect on animal health, human health or the environment⁽³⁾. It further concluded that preparation does not present any other risk which would, in accordance with Article 5(2) of Regulation (EC) No 1831/2003, exclude authorisation. According to that opinion, the use of that preparation does not have an adverse effect on these additional animal categories. The opinion of the Authority recommends appropriate measures for user safety. It does not consider that there is a need for specific requirements of post market monitoring. This opinion also verifies the report on the method of analysis of the feed additive in feed submitted by the Community Reference Laboratory set up by Regulation (EC) No 1831/2003.

(6) The assessment of that preparation shows that the conditions for authorisation, provided for in Article 5 of Regulation (EC) No 1831/2003, are satisfied. Accordingly, the use of that preparation should be authorised, as specified in the Annex to this Regulation.

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

HAS ADOPTED THIS REGULATION:

Article 1

The preparation specified in the Annex, belonging to the additive category 'zootechnical additives' and to the functional group 'digestibility enhancers', is authorised as an additive in animal nutrition subject to the conditions laid down in that Annex.

Article 2

This Regulation shall enter into force on the 20th day following its publication in the *Official Journal of the European Union*.

⁽¹⁾ OJ L 268, 18.10.2003, p. 29. Regulation as amended by Commission Regulation (EC) No 378/2005 (OJ L 59, 5.3.2005, p. 8).

⁽²⁾ OJ L 73, 13.3.2007, p. 4.

⁽³⁾ Opinion of the Scientific Panel on Additives and Products or Substances used in Animal Feed on the safety and efficacy of the enzyme preparation of Natuphos (3-phytase) produced by *Aspergillus niger* for laying hens and turkeys for fattening. Adopted on 17 April 2007. The EFSA Journal (2007) 472, 1-4.

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

Done at Brussels, 1 October 2007.

For the Commission
Markos KYPRIANOU
Member of the Commission

ANNEX

Identification number of the additive	Name of the holder of authorisation	Additive (Trade name)	Composition, chemical formula, description, analytical method	Species or category of animal	Maximum age	Minimum content		Maximum content	Other provisions	End of period of authorisation
						Units of activity/kg of complete feedingstuff with a moisture content of 12 %				
Category of zootechnical additives. Functional group: digestibility enhancers										
4a1600	BASF Aktiengesellschaft	3-phytase EC 3.1.3.8 (Natuphos 5 000 G Natuphos 5 000 L Natuphos 5 000 L Natuphos 10 000 G Natuphos 10 000 L)	Additive composition: 3-phytase produced by <i>Aspergillus niger</i> (CBS 101 672) having a minimum activity of: Solid form: 5 000 FTU ⁽¹⁾ /g Liquid form: 5 000 FTU/ml Characterisation of the active substance: 3-phytase produced by <i>Aspergillus niger</i> (CBS 101 672) Analytical method ⁽²⁾ Colorimetric method measuring inorganic phosphate released by the enzyme from phytate substrate.	Laying hens	—	250 FTU			1. In the directions for use of the additive and premixture, indicate the storage temperature, storage life, and stability to pelleting. 2. Recommended dose per kilogram of complete feedingstuff: 300-400 FTU. 3. For use in feed containing more than 0,23 % phytin bound phosphorus.	22.10.2017
				Turkeys for fattening		250 FTU			1. In the direction of use of the additive and premixtures, indicate the storage life and stability in pelleting. 2. Recommended dose per kilogram of complete feedingstuff: 500 FTU. 3. For use in feed containing more than 0,23 % phytin bound phosphorus.	

^(1) 1 FTU is the amount of enzyme which liberates 1 micromole of inorganic phosphate per minute from sodium phytate at pH 5,5 and 37 °C.
Details of the analytical methods are available at the following address of the Community Reference Laboratory: www.irmm.jrc.be/crl-feed-additives

⁽¹⁾ 1 FTU is the amount of enzyme which liberates 1 micromole of inorganic phosphate per minute from sodium phytate at pH 5,5 and 37 °C.

⁽²⁾ Details of the analytical methods are available at the following address of the Community Reference Laboratory: www.irmm.jrc.be/crl-feed-additives

COMMISSION REGULATION (EC) No 1143/2007

of 1 October 2007

amending Regulation (EC) No 256/2002 as regards the authorisation of the feed additive preparation of *Bacillus cereus* var. *toyoi*, belonging to the group of microorganisms

(Text with EEA relevance)

THE COMMISSION OF THE EUROPEAN COMMUNITIES,

Having regard to the Treaty establishing the European Community,

Having regard to Regulation (EC) No 1831/2003 of the European Parliament and of the Council of 22 September 2003 on additives for use in animal nutrition⁽¹⁾, and in particular Article 9(2) thereof,

Whereas:

(1) Regulation (EC) No 1831/2003 provides for the authorisation of additives for use in animal nutrition and for the grounds and procedures for granting such authorisation.

(2) The preparation *Bacillus cereus* var. *toyoi* (NCIMB 40112/CNCM I-1012), belonging to the group of 'micro-organisms', was authorised in accordance with Council Directive 70/524/EEC⁽²⁾, in particular, by Commission Regulation (EC) No 256/2002⁽³⁾ without a time limit for use for piglets, until two months, and for sows, from one week prior to farrowing until weaning. This additive was subsequently entered in the Community Register of Feed Additives as an existing product, in accordance with Article 10 of Regulation (EC) No 1831/2003.

(3) In accordance with Article 7 of Regulation (EC) No 1831/2003, an application for an amendment of the authorisation of that preparation was submitted to allow its use in feed for sows from service to weaning. That application was accompanied by the particulars and documents required under Article 7(3) of that Regulation.

(4) The European Food Safety Authority (the Authority) concluded in its opinion of 7 March 2007 that the preparation *Bacillus cereus* var. *toyoi* NCIMB 40112/CNCM I-1012 does not have an adverse effect on animal health, human health or the environment⁽⁴⁾. It further concluded that the preparation does not present any other risk which would, in accordance with Article 5(2) of Regulation (EC) No 1831/2003, exclude authorisation. The Authority does not consider that there is a need for specific requirements of post market monitoring. It also verified the report on the method of analysis of the feed additive in feed submitted by the Community Reference Laboratory set up by Regulation (EC) No 1831/2003.

(5) The assessment of that preparation shows that the conditions provided for in Article 5 of Regulation (EC) No 1831/2003 are satisfied.

(6) Regulation (EC) No 256/2002 should therefore be amended accordingly.

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

HAS ADOPTED THIS REGULATION:

Article 1

Annex III to Regulation (EC) No 256/2002 is replaced by the text in the Annex to this Regulation.

Article 2

This Regulation shall enter into force on the 20th day following its publication in the *Official Journal of the European Union*.

⁽¹⁾ OJ L 268, 18.10.2003, p. 29. Regulation as amended by Commission Regulation (EC) No 378/2005 (OJ L 59, 5.3.2005, p. 8).

⁽²⁾ OJ L 270, 14.12.1970, p. 1. Directive repealed by Regulation (EC) No 1831/2003.

⁽³⁾ OJ L 41, 13.2.2002, p. 6.

⁽⁴⁾ Opinion of the Scientific Panel on Additives and Products or Substances used in Animal Feed on the safety and efficacy of the product Toyocerin (*Bacillus cereus* var. *Toyoi*) as feed additive for sows in accordance with Regulation (EC) No 1831/2003. Adopted on 7 March 2007. The EFSA Journal (2007) 458, 1-9.

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

Done at Brussels, 1 October 2007.

For the Commission
Markos KYPRIANOU
Member of the Commission

ANNEX

'ANNEX III

Microorganisms

EC No	Additive	Chemical formula, description	Species or category of animal	Maximum age	Minimum content	Maximum content	Other provisions	End of period of authorisation
					CFU/kg of complete feedingstuff			
Microorganisms								
E 1701	<i>Bacillus cereus</i> var. <i>toyoi</i> NCIMB 40112/ CNCM I-1012	Preparation of <i>Bacillus cereus</i> var. <i>toyoi</i> containing a minimum of 1×10^{10} CFU/g additive	Piglets	Two months	1×10^9	1×10^9	In the directions for use of the additive and premixture, indicate the storage temperature, storage life and stability to pelleting.	Without a time limit
			Sows	From service to weaning	$0,5 \times 10^9$	2×10^9	In the directions for use of the additive and premixture, indicate the storage temperature, storage life and stability to pelleting.	Without a time limit

COMMISSION REGULATION (EC) No 1144/2007**of 1 October 2007****amending Regulation (EC) No 1831/96 opening and providing for the administration of Community tariff quotas bound under GATT for certain fruit and vegetables and processed fruit and vegetable products from 1996**

THE COMMISSION OF THE EUROPEAN COMMUNITIES,

Having regard to the Treaty establishing the European Community,

Having regard to Council Regulation (EC) No 2200/96 of 28 October 1996 on the common organisation of the market in fruit and vegetables ⁽¹⁾, and in particular Article 34(1) thereof,

Whereas:

(1) The Agreement in the form of an Exchange of Letters between the European Community and the Argentine Republic pursuant to Article XXIV:6 and Article XXVIII of the General Agreement on Tariffs and Trade (GATT) 1994 ⁽²⁾, approved by Council Decision 2006/930/EC ⁽³⁾, provides for the increasing of the existing GATT tariff quota for apples.

(2) Commission Regulation (EC) No 1831/96 of 23 September 1996 opening and providing for the admin-

istration of Community tariff quotas bound under GATT for certain fruit and vegetables and processed fruit and vegetable products from 1996 ⁽⁴⁾ should be amended accordingly.

(3) The measures provided for in this Regulation are in accordance with the opinion of the Management Committee for Fresh Fruit and Vegetables,

HAS ADOPTED THIS REGULATION:

Article 1

In Annex III to Regulation (EC) No 1831/96, the quota volume (tonnes) for Order No 09.0061 for Apples, fresh, other than cider apples is replaced by '696'.

Article 2

This Regulation shall enter into force on the third 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 October 2007.

For the Commission

Mariann FISCHER BOEL

Member of the Commission

⁽¹⁾ OJ L 297, 21.11.1996, p. 1. Regulation as last amended by Commission Regulation (EC) No 47/2003 (OJ L 7, 11.1.2003, p. 64).

⁽²⁾ OJ L 355, 15.12.2006, p. 92.

⁽³⁾ OJ L 355, 15.12.2006, p. 91.

⁽⁴⁾ OJ L 243, 24.9.1996 p. 5. Regulation as amended by Regulation (EC) No 973/2006 (OJ L 176, 30.6.2006, p. 63).

II

(Acts adopted under the EC Treaty/Euratom Treaty whose publication is not obligatory)

DECISIONS

COUNCIL DECISION

of 18 September 2007

on the conclusion of the Agreement between the European Community and New Zealand on certain aspects of air services

(2007/633/EC)

THE COUNCIL OF THE EUROPEAN UNION,

later date, in accordance with Council Decision 2006/466/EC ⁽²⁾.

Having regard to the Treaty establishing the European Community, and in particular Article 80(2), in conjunction with Article 300(2), first sentence of the first subparagraph, and Article 300(3), first subparagraph,

(4) The Agreement should be approved,

Having regard to the proposal from the Commission,

HAS DECIDED AS FOLLOWS:

Having regard to the opinion of the European Parliament ⁽¹⁾,

Article 1

The Agreement between the European Community and New Zealand on certain aspects of air services is hereby approved on behalf of the Community.

Whereas:

Article 2

The President of the Council is authorised to designate the person empowered to make the notification provided in Article 8(1) of the Agreement.

(1) By a Decision of 5 June 2003 the Council has authorised the Commission to open negotiations with third countries on the replacement of certain provisions in existing bilateral agreements by a Community agreement.

(2) The Commission has negotiated, on behalf of the Community, an Agreement with New Zealand on certain aspects of air services in accordance with the mechanisms and directives in the Annex to the Decision of 5 June 2003.

Done at Brussels, 18 September 2007.

(3) The Agreement has been signed on behalf of the Community subject to its possible conclusion at a

For the Council
The President
R. PEREIRA

⁽¹⁾ Not yet published in the Official Journal.

⁽²⁾ OJ L 184, 6.7.2006, p. 25.

III

(Acts adopted under the EU Treaty)

ACTS ADOPTED UNDER TITLE V OF THE EU TREATY

COUNCIL JOINT ACTION 2007/634/CFSP

of 1 October 2007

**amending Joint Action 2007/113/CFSP amending and extending the mandate of the European Union
Special Representative for Central Asia**

THE COUNCIL OF THE EUROPEAN UNION,

external actions of the European Union in the region
without prejudice to Community competence;

Having regard to the Treaty on European Union and, in
particular, Articles 14, 18(5) and 23(2) thereof,

Whereas:

(1) On 15 February 2007, the Council adopted Joint Action 2007/113/CFSP ⁽¹⁾ amending and extending the mandate of the European Union Special Representative (EUSR) for Central Asia.

(2) At its meeting of 21 and 22 June 2007, the European Council adopted an EU strategy for a new partnership with Central Asia. As the EUSR has been assigned a role in monitoring the implementation of that strategy, his mandate should be adapted accordingly.

(3) On 27 July 2007, on the basis of a mid-term review of Joint Action 2007/113/CFSP, the Political and Security Committee recommended making further adjustments to the EUSR's mandate.

(4) Joint Action 2007/113/CFSP should be amended accordingly,

HAS ADOPTED THIS JOINT ACTION:

Article 1

Article 3(1) of Joint Action 2007/113/CFSP shall be replaced by the following:

'1. In order to achieve the policy objectives, the EUSR's mandate shall be to:

(a) promote overall political coordination of the European Union in Central Asia and ensure consistency of the

(b) monitor, on behalf of the High Representative and in accordance with his mandate, together with the Commission and the Presidency, and without prejudice to Community competence, the implementation process of the EU Strategy for a New Partnership with Central Asia, make recommendations and report to relevant Council bodies on a regular basis;

(c) assist the Council in further developing a comprehensive policy towards Central Asia;

(d) follow closely political developments in Central Asia by developing and maintaining close contacts with governments, parliaments, judiciary, civil society and mass media;

(e) encourage Kazakhstan, the Kyrgyz Republic, Tajikistan, Turkmenistan and Uzbekistan to cooperate on regional issues of common interest;

(f) develop appropriate contacts and cooperation with the main interested actors in the region, and all relevant regional and international organisations, including the Shanghai Cooperation Organisation (SCO), the Eurasian Economic Community (EURASEC), the Conference on Interaction and Confidence-Building Measures in Asia (CICA), the Collective Security Treaty Organisation (CSTO), the Central Asia Regional Economic Cooperation Program (CAREC) and the Central Asian Regional Information and Coordination Centre (CARICC);

⁽¹⁾ OJ L 46, 16.2.2007, p. 83.

- (g) contribute to the implementation of the European Union human rights policy and European Union Guidelines on Human Rights, in particular with regard to women and children in conflict-affected areas, especially by monitoring and addressing developments in this regard;
- (h) contribute, in close cooperation with the OSCE, to conflict prevention and resolution by developing contacts with the authorities and other local actors (NGOs, political parties, minorities, religious groups and their leaders);
- (i) provide input to the formulation of energy security aspects and anti-narcotics aspects of the CFSP with respect to Central Asia.’.

Article 2

This Joint Action shall enter into force on the day of its adoption.

Article 3

This Joint Action shall be published in the *Official Journal of the European Union*.

Done at Luxembourg, 1 October 2007.

For the Council

The President

M. LINO

COUNCIL COMMON POSITION 2007/635/CFSP**of 1 October 2007****extending Common Position 2004/694/CFSP on further measures in support of the effective implementation of the mandate of the International Criminal Tribunal for the former Yugoslavia (ICTY)**

THE COUNCIL OF THE EUROPEAN UNION,

Having regard to the Treaty on European Union, and in particular Article 15 thereof,

Whereas:

- (1) On 11 October 2004, the Council adopted Common Position 2004/694/CFSP ⁽¹⁾ on further measures in support of the effective implementation of the mandate of the International Criminal Tribunal for the former Yugoslavia (ICTY), in order to freeze all funds and economic resources belonging to all persons who have been publicly indicted by the ICTY for war crimes but who are not in the custody of the ICTY.
- (2) Common Position 2004/694/CFSP applies until 10 October 2007.
- (3) The Council considers it necessary to renew Common Position 2004/694/CFSP for a further period of 12 months.
- (4) The implementing Community measures are set out in Council Regulation (EC) No 1763/2004 of 11 October 2004 imposing certain restrictive measures in support of

effective implementation of the mandate of the International Criminal Tribunal for the former Yugoslavia (ICTY) ⁽²⁾,

HAS ADOPTED THIS COMMON POSITION:

Article 1

Common Position 2004/694/CFSP shall be extended until 10 October 2008.

Article 2

This Common Position shall take effect on the date of its adoption.

*Article 3*This Common Position shall be published in the *Official Journal of the European Union*.

Done at Luxembourg, 1 October 2007.

For the Council
The President
M. LINO

⁽¹⁾ OJ L 315, 14.10.2004, p. 52. Common Position as last amended by Decision 2007/449/CFSP (OJ L 169, 29.6.2007, p. 75).

⁽²⁾ OJ L 315, 14.10.2004, p. 14. Regulation as last amended by Commission Regulation (EC) No 789/2007 (OJ L 175, 5.7.2007, p. 27).

CORRIGENDA

Corrigendum to Council Regulation (EC) No 1890/2005 of 14 November 2005 imposing a definitive anti-dumping duty and collecting definitively the provisional duty imposed on imports of certain stainless steel fasteners and parts thereof originating in the People's Republic of China, Indonesia, Taiwan, Thailand and Vietnam and terminating the proceeding on imports of certain stainless steel fasteners and parts thereof originating in Malaysia and the Philippines

(Official Journal of the European Union L 302 of 19 November 2005)

On page 13, in the Annex, under the list of companies:

in the 16th row:

for: 'KUOLIEN SCREW INDUSTRIAL CO. LTD, Kwanmiao',

read: 'KUOLIEN SCREW INDUSTRIAL CO. LTD, Kaohsiung';

in the 17th row:

for: 'KWANTEX RESEARCH INC, Taipei',

read: 'KWANTEX RESEARCH INC, Tainan';

in the 21st row:

for: 'M & W FASTENER CO. LTD, Kaoshiung',

read: 'M & W FASTENER CO. LTD, Kaohsiung';

in the 27th row:

for: 'SUMEEKO INDUSTRIES CO. LTD, Kaoshiung',

read: 'SUMEEKO INDUSTRIES CO. LTD, Kaohsiung'.
