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Contents

I Acts whose publication is obligatory

Commission Regulation (EC) No 1283/2006 of 29 August 2006 establishing the standard import values for determining the entry price of certain fruit and vegetables	1
★ Commission Regulation (EC) No 1284/2006 of 29 August 2006 concerning the permanent authorisations of certain additives in feedingstuffs ⁽¹⁾	3
★ Commission Regulation (EC) No 1285/2006 of 29 August 2006 opening the procedure for the allocation of export licences for cheese to be exported to the United States of America in 2007 under certain GATT quotas	8
★ Commission Regulation (EC) No 1286/2006 of 29 August 2006 amending for the 70th time Council Regulation (EC) No 881/2002 imposing certain specific restrictive measures directed against certain persons and entities associated with Usama bin Laden, the Al-Qaida network and the Taliban, and repealing Council Regulation (EC) No 467/2001	14
★ Commission Directive 2006/74/EC of 21 August 2006 amending Council Directive 91/414/EEC to include dichlorprop-P, metconazole, pyrimethanil and triclopyr as active substances ⁽¹⁾	17

Corrigenda

★ Corrigendum to Council Regulation (EC) No 318/2006 of 20 February 2006 on the common organisation of the markets in the sugar sector (OJ L 58, 28.2.2006)	23
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⁽¹⁾ Text with EEA relevance

I

(Acts whose publication is obligatory)

COMMISSION REGULATION (EC) No 1283/2006
of 29 August 2006
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 30 August 2006.

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

Done at Brussels, 29 August 2006.

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 386/2005 (OJ L 62, 9.3.2005, p. 3).

ANNEX

to Commission Regulation of 29 August 2006 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	052	53,4
	068	147,1
	999	100,3
0707 00 05	052	87,4
	999	87,4
0709 90 70	052	85,7
	999	85,7
0805 50 10	388	67,0
	524	48,0
	528	52,6
	999	55,9
0806 10 10	052	82,4
	220	99,2
	624	139,0
	999	106,9
0808 10 80	388	89,1
	400	90,7
	508	81,5
	512	85,8
	528	77,4
	720	82,6
	800	140,1
	804	92,5
0808 20 50	999	92,5
	052	124,2
	388	88,6
0809 30 10, 0809 30 90	999	106,4
	052	119,8
	096	12,8
0809 40 05	999	66,3
	052	96,0
	066	47,1
	098	45,7
	624	149,1
	999	84,5

⁽¹⁾ Country nomenclature as fixed by Commission Regulation (EC) No 750/2005 (OJ L 126, 19.5.2005, p. 12). Code '999' stands for 'of other origin'.

COMMISSION REGULATION (EC) No 1284/2006
of 29 August 2006
concerning the permanent authorisations of certain additives in feedingstuffs
(Text with EEA relevance)

THE COMMISSION OF THE EUROPEAN COMMUNITIES,

Regulation (EC) No 1831/2003. Those applications are therefore to continue to be treated in accordance with Article 4 of Directive 70/524/EEC.

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 9d(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 applications for the authorisation of the additives listed in the Annexes to this Regulation were submitted before the date of application of Regulation (EC) No 1831/2003.
- (4) Initial comments on those applications, as provided for in Article 4(4) of Directive 70/524/EEC, were forwarded to the Commission before the date of application of

- (5) The use of the enzyme preparation of endo-1,3(4)-beta-glucanase produced by *Trichoderma longibrachiatum* (ATCC 2106), endo-1,4-beta-xylanase produced by *Trichoderma longibrachiatum* (ATCC 2105) and alpha-amylase produced by *Bacillus amyloliquefaciens* (DSM 9553) was provisionally authorised for the first time for weaned piglets by Commission Regulation (EC) No 2690/1999⁽³⁾. New data were submitted in support of an application for authorisation without a time-limit of that enzyme preparation. The assessment shows that the conditions laid down in Article 3a of Directive 70/524/EEC for such authorisation are satisfied. Accordingly, the use of that enzyme preparation, as specified in Annex I to this Regulation, should be authorised without a time-limit.
- (6) The use of the enzyme preparation of 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) and endo-1,4-beta-xylanase produced by *Trichoderma viride* (NIBH FERM BP 4842) was provisionally authorised for the first time for turkeys for fattening by Commission Regulation (EC) No 2013/2001⁽⁴⁾. New data were submitted in support of an application for authorisation without a time-limit of that enzyme preparation. The assessment shows that the conditions laid down in Article 3a of Directive 70/524/EEC for such authorisation are satisfied. Accordingly, the use of that enzyme preparation, as specified in Annex II to this Regulation, should be authorised without a time-limit.
- (7) The use of the enzyme preparation of endo-1,4-beta-xylanase produced by *Trichoderma longibrachiatum* (ATCC 2105) was provisionally authorised for the first time for pigs for fattening by Commission Regulation (EC) No 1411/1999⁽⁵⁾. New data were submitted in support of an application for authorisation without a time-limit of that enzyme preparation. The assessment shows that the conditions laid down in Article 3a of Directive 70/524/EEC for such authorisation are satisfied. Accordingly, the use of that enzyme preparation, as specified in Annex III to this Regulation, should be authorised without a time-limit.

⁽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).

⁽³⁾ OJ L 326, 18.12.1999, p. 33.

⁽⁴⁾ OJ L 272, 13.10.2001, p. 24.

⁽⁵⁾ OJ L 164, 30.6.1999, p. 56.

(8) The assessment of these applications 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 ⁽¹⁾.

(9) 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 belonging to the group 'Enzymes', as specified in Annex I, is authorised without a time-limit as additive in animal nutrition under the conditions laid down in that Annex.

Article 2

The preparation belonging to the group 'Enzymes', as specified in Annex II, is authorised without a time-limit as additive in animal nutrition under the conditions laid down in that Annex.

Article 3

The preparation belonging to the group 'Enzymes', as specified in Annex III, is authorised without a time-limit as additive in animal nutrition under the conditions laid down in that Annex.

Article 4

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, 29 August 2006.

For the Commission

Markos KYPRIANOU

Member of the Commission

⁽¹⁾ OJ L 183, 29.6.1989, p. 1. Directive as amended by Regulation (EC) No 1882/2003 of the European Parliament and of the Council (OJ L 284, 31.10.2003, p. 1).

ANNEX I

EC No	Additive	Chemical formula, description	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			
Enzymes								
E 1638	Endo-1,3(4)-beta-glucanase EC 3.2.1.6 Endo-1,4-beta-xylanase EC 3.2.1.8 Alpha-amylase EC 3.2.1.1	Preparation of endo-1,3(4)-beta-glucanase produced by <i>Trichoderma longibrachiatum</i> (ATCC 2106), endo-1,4-beta-xylanase produced by <i>Trichoderma longibrachiatum</i> (ATCC 2105) and alpha-amylase produced by <i>Bacillus amyloliquefaciens</i> (DSM 9553) having a minimum activity of: Endo-1,3(4)-beta-glucanase: 250 U ⁽¹⁾ /g Endo-1,4-beta-xylanase: 400 U ⁽²⁾ /g Alpha-amylase: 1 000 U ⁽³⁾ /g	Piglets (weaned)	—	Endo-1,3(4)-beta-glucanase: 250 U Endo-1,4-beta-xylanase: 400 U Alpha-amylase: 1 000 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 kg of complete feedingstuff: — endo-1,3(4)-beta-glucanase: 250 U — endo-1,4-beta-xylanase: 400 U — alpha-amylase: 1 000 U. 3. For use in compound feed containing cereals rich in starch and non-starch polysaccharides (mainly arabinoxylans and beta-glucans), e.g. containing more than 50 % barley. 4. For weaned piglets up to approximately 35 kg.	Without a time-limit
⁽¹⁾ 1 U is the amount of enzyme which liberates 1 micromole of reducing sugars (glucose equivalents) from barley beta-glucan per minute at pH 5,0 and 30 °C. ⁽²⁾ 1 U is the amount of enzyme which liberates 1 micromole of reducing sugars (xylose equivalents) from oat spelt xylan per minute at pH 5,3 and 50 °C. ⁽³⁾ 1 U is the amount of enzyme which hydrolyses 1 micromole of glucosidic linkages from a water insoluble cross-linked starch polymer substrate per minute at pH 6,5 and 37 °C.								

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

⁽²⁾ 1 U is the amount of enzyme which liberates 1 micromole of reducing sugars (xylose equivalents) from oat spelt xylan per minute at pH 5,3 and 50 °C.

⁽³⁾ 1 U is the amount of enzyme which hydrolyses 1 micromole of glucosidic linkages from a water insoluble cross-linked starch polymer substrate per minute at pH 6,5 and 37 °C.

ANNEX II

EC No	Additive	Chemical formula, description	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				
Enzymes									
E 1621	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 Endo-1,4-beta-xylanase E.C. 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) 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: 10 000 U ⁽¹⁾ /g Endo-1,4-beta-glucanase: 120 000 U ⁽²⁾ /g Alpha-amylase: 400 U ⁽³⁾ /g Endo-1,4-beta-xylanase: 210 000 U ⁽⁴⁾ /g	Turkeys for fattening	—	Endo-1,3(4)-beta-glucanase: 500 U Endo-1,4-beta-glucanase: 6 000 U Alpha-amylase: 20 U Endo-1,4-beta-xylanase: 10 500 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 kg of complete feedingsuff: — endo-1,3(4)-beta-glucanase: 1 000-1 500 U — endo-1,4-beta-glucanase: 12 000-18 000 U — alpha-amylase: 40-60 U — endo-1,4-beta-xylanase: 21 000-31 500 U 3. For use in compound feed rich in non-starch polysaccharides (mainly beta-glucans and arabinoxylans), e.g. containing more than 30 % wheat.	Without a time-limit	
⁽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 0,0036 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 liberates 0,0067 micromoles of reducing sugars (xylose equivalents) from birchwood xylan per minute at pH 5,3 and 50 °C.									

⁽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 liberates 0,0067 micromoles of reducing sugars (xylose equivalents) from birchwood xylan per minute at pH 5,3 and 50 °C.

ANNEX III

EC No	Additive	Chemical formula, description	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				
Enzymes									
E 1628	Endo-1,4-beta-xylanase EC 3.2.1.8	Preparation of endo-1,4-beta-xylanase produced by <i>Trichoderma longibrachiatum</i> (ATCC 2105) having a minimum activity of: Powder form: Endo-1,4-beta-xylanase: 8 000 U (1)/g Liquid form: Endo-1,4-beta-xylanase: 8 000 U/ml	Pigs for fattening	—	Endo-1,4-beta-xylanase: 1 000 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 kg of complete feedingstuff: — endo-1,4-beta-xylanase: 1 000-4 000 U 3. For use in compound feed rich in non-starch polysaccharides (mainly arabinoxylans), e.g. containing more than 35 % wheat.	Without a time-limit	

(1) 1 U is the amount of enzyme which liberates 1 micromole of reducing sugars (xylose equivalents) from oat spelt xylan per minute at pH 5.3 and 50 °C.

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

COMMISSION REGULATION (EC) No 1285/2006**of 29 August 2006****opening the procedure for the allocation of export licences for cheese to be exported to the United States of America in 2007 under certain GATT quotas**

THE COMMISSION OF THE EUROPEAN COMMUNITIES,

HAS ADOPTED THIS REGULATION:

Having regard to the Treaty establishing the European Community,

Having regard to Council Regulation (EC) No 1255/1999 of 17 May 1999 on the common organisation of the market in milk and milk products ⁽¹⁾, and in particular Article 30 thereof,

Whereas:

- (1) Section 2 of Chapter III of Commission Regulation (EC) No 1282/2006 of 17 August 2006 laying down special detailed rules for the application of Council Regulation (EC) No 1255/1999 as regards export licences and export refunds for milk and milk products ⁽²⁾, provides that export licences for cheese exported to the United States of America as part of the quotas under the agreements concluded during multilateral trade negotiations, may be allocated in accordance with a special procedure provided for therein.
- (2) That procedure should be opened for exports during 2007 and the additional rules relating to it should be determined.
- (3) In administering imports the competent authorities in the USA make a distinction between the additional quota granted to the European Community under the Uruguay Round and the quotas resulting from the Tokyo Round. Export licences should be allocated taking into account the eligibility of those products for the USA quota in question as described in the Harmonised Tariff Schedule of the United States of America.
- (4) With a view to exporting the maximum quantity under the quotas for which there is moderate interest, applications covering the whole quota quantity should be allowed.
- (5) The measures provided for in this Regulation are in accordance with the opinion of the Management Committee for Milk and Milk Products,

Article 1

Export licences for products falling within CN code 0406 and listed in Annex I to this Regulation to be exported to the United States of America in 2007 under the quotas referred to in Article 23 of Regulation (EC) No 1282/2006 shall be issued in accordance with Section 2 of Chapter III of that Regulation and with the provisions of this Regulation.

Article 2

1. Applications for licences referred to in Article 24 of Regulation (EC) No 1282/2006 (hereinafter referred to as applications) shall be lodged with the competent authorities from 1 to 7 September 2006 at the latest.

2. Applications shall be admissible only if they contain all the information referred to in Article 24 of Regulation (EC) No 1282/2006 and if they are accompanied by the documents referred to therein.

Where, for the same group of products referred to in column 2 of Annex I to this Regulation the available quantity is divided between the Uruguay Round quota and the Tokyo Round quota, licence applications may cover only one of those quotas and shall indicate the quota concerned, specifying the identification of the group and of the quota indicated in column 3 of that Annex.

Information referred to in Article 24 of Regulation (EC) No 1282/2006 shall be presented in accordance with the model set out in Annex II to this Regulation.

3. As regards the quotas identified by 22-Tokyo and 22-Uruguay in column 3 of Annex I, applications shall cover at least 10 tonnes and shall not exceed the quantity available under the quota concerned as set out in column 4 of that Annex.

As regards the other quotas indicated in column 3 of Annex I, applications shall cover at least 10 tonnes and no more than 40 % of the quantity available under the quota concerned as set out in column 4 of that Annex.

⁽¹⁾ OJ L 160, 26.6.1999, p. 48. Regulation as last amended by Regulation (EC) No 1913/2005 (OJ L 307, 25.11.2005, p. 2).

⁽²⁾ OJ L 234, 29.8.2006, p. 4.

4. Applications shall be admissible only if applicants declare in writing that they have not lodged other applications for the same group of products and the same quota and undertake not to do so.

If an applicant lodges several applications for the same group of products and the same quota in one or more Member States, all his applications shall be deemed inadmissible.

Article 3

1. Member States shall notify the Commission, within five working days after the end of the period for lodging applications, of the applications lodged for each of the groups of products and, where applicable, the quotas indicated in Annex I.

All notifications, including 'nil' notifications, shall be made by fax or e-mail on the model form set out in Annex III.

2. Notification shall comprise for each group and, where applicable, for each quota:

- (a) a list of applicants;
- (b) the quantities applied for by each applicant broken down by the product code of the Combined Nomenclature and by their code in accordance with the Harmonized Tariff Schedule of the United States of America (2006);
- (c) indication that applicant has exported the products concerned to the United States of America in at least one of the preceding three years;

- (d) the name and address of the importer designated by the applicant and the confirmation that the importer is a subsidiary of the applicant.

Article 4

The Commission shall, pursuant to Article 25 of Regulation (EC) No 1282/2006, determine the allocation of licences without delay and shall notify the Member States thereof by 31 October 2006 at the latest.

Member States shall notify the Commission, within five working days after publication of the allocation coefficients, for each group and, where applicable, for each quota, the quantities allocated by applicant, in accordance to Article 25 of Regulation (EC) No 1282/2006.

The notification shall be made by fax or e-mail on the model form set out in Annex IV to this Regulation.

Article 5

The information notified under Article 3 of this Regulation and under Article 24 of Regulation (EC) No 1282/2006 shall be verified by the Member States before the licences are issued and by 15 December 2006 at the latest.

Where it is found that incorrect information has been supplied by an operator to whom a licence has been issued, the licence shall be cancelled and the security forfeited. The Member States shall communicate it to the Commission without any delay.

Article 6

This Regulation shall enter into force on the 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, 29 August 2006.

For the Commission

Mariann FISCHER BOEL

Member of the Commission

ANNEX I

Cheese to be exported to the United States of America in 2007 under certain GATT quotas

Section 2 of Chapter III of Regulation (EC) No 1282/2006 and Regulation (EC) No 1285/2006

Identification of group in accordance with Additional Notes in Chapter 4 of the Harmonised Tariff Schedule of the United States		Identification of group and quota	Quantity available for 2007
Note to	Group		Tonnes
(1)	(2)	(3)	(4)
16	Not specifically provided for (NSPF)	16 — Tokyo	908,877
		16 — Uruguay	3 446,000
17	Blue Mould	17	350,000
18	Cheddar	18	1 050,000
20	Edam/Gouda	20	1 100,000
21	Italian type	21	2 025,000
22	Swiss or Emmenthaler cheese other than with eye formation	22 — Tokyo	393,006
		22 — Uruguay	380,000
25	Swiss or Emmenthaler cheese with eye formation	25 — Tokyo	4 003,172
		25 — Uruguay	2 420,000

ANNEX II
Presentation of information required pursuant to Article 24 of Regulation (EC) No 1282/2006

Identification of group and quota referred to in column 3 of Annex I to Regulation (EC) No 1282/2006:

Name of group indicated in column 2 of Annex I to Regulation (EC) No 1282/2006:

Origin of quota: Uruguay Round: ☐ Tokyo Round: ☐

Name/address of applicant	Product code of the Combined Nomenclature	Quantity applied for in tonnes	Harmonised Tariff Schedule of the USA code	Name/address of designated importer	Information required under Article 24(3) of Regulation (EC) No 1282/2006	
					Exports in at least one of the preceding three years	Importer is subsidiary
					Yes ☐	Yes ☐
					Yes ☐	Yes ☐
Total:						

COMMISSION REGULATION (EC) No 1286/2006**of 29 August 2006****amending for the 70th time Council Regulation (EC) No 881/2002 imposing certain specific restrictive measures directed against certain persons and entities associated with Usama bin Laden, the Al-Qaida network and the Taliban, and repealing Council Regulation (EC) No 467/2001**

THE COMMISSION OF THE EUROPEAN COMMUNITIES,

Having regard to the Treaty establishing the European Community,

Having regard to Council Regulation (EC) No 881/2002 imposing certain specific restrictive measures directed against certain persons and entities associated with Usama bin Laden, the Al-Qaida network and the Taliban, and repealing Council Regulation (EC) No 467/2001 prohibiting the export of certain goods and services to Afghanistan, strengthening the flight ban and extending the freeze of funds and other financial resources in respect of the Taliban of Afghanistan ⁽¹⁾, and in particular Article 7(1), first indent, thereof,

Whereas:

- (1) Annex I to Regulation (EC) No 881/2002 lists the persons, groups and entities covered by the freezing of funds and economic resources under that Regulation.

- (2) On 18 and 23 August 2006, the Sanctions Committee of the United Nations Security Council decided to amend the list of persons, groups and entities to whom the freezing of funds and economic resources should apply. On 25 July, the Sanctions Committee decided a number of amendments of existing entries and one of them should still be included in Annex I. Annex I should therefore be amended accordingly,

HAS ADOPTED THIS REGULATION:

Article 1

Annex I to Regulation (EC) No 881/2002 is hereby amended as set out in the Annex to this Regulation.

Article 2

This Regulation shall enter into force on the 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, 29 August 2006.

For the Commission

Eneko LANDÁBURU

Director-General for External Relations

⁽¹⁾ OJ L 139, 29.5.2002, p. 9. Regulation as last amended by Commission Regulation (EC) No 1228/2006 (OJ L 222, 15.8.2006, p. 6).

ANNEX

Annex I to Regulation (EC) No 881/2002 is amended as follows:

- (1) The following entry is removed from the heading 'Natural persons':

'Ali Ahmed Yusuf (alias Ali Galoul), Krälingegrund 33, S-16362 Spånga, Sweden; date of birth: 20 November 1974; place of birth: Garbaharey, Somalia; nationality: Swedish; passport No: Swedish passport 1041635; national identification No: 741120-1093'.

- (2) The entry 'Hani Al-Sayyid **Al-Sebai** (alias (a) Hani Yousef Al-Sebai, (b) Hani Youssef, (c) Hany Youseff, (d) Hani Yusef, (e) Hani al-Sayyid Al-Sabai, (f) Hani al-Sayyid El Sebai, (g) Hani al-Sayyid Al Siba'i, (h) Hani al-Sayyid El Sabaay, (i) El-Sababt, (j) Abu Tusnin, (k) Abu Akram, (l) Hani El Sayyed Elsebai Yusef, (m) Abu Karim). Address: London, United Kingdom. Date of birth: (a) 1.3.1961, (b) 16.6.1960. Place of birth: Qaylubiyah, Egypt. Nationality: Egyptian' under the heading 'Natural persons' is replaced by:

'Hani Al-Sayyid **Al-Sebai** (alias (a) Hani Yousef **Al-Sebai**, (b) Hani Youssef, (c) Hany Youseff, (d) Hani Yusef, (e) Hani al-Sayyid **Al-Sabai**, (f) Hani al-Sayyid **El Sebai**, (g) Hani al-Sayyid **Al Siba'i**, (h) Hani al-Sayyid **El Sabaay**, (i) El-Sababt, (j) Abu Tusnin, (k) Abu Akram, (l) Hani El Sayyed Elsebai Yusef, (m) Abu Karim, (n) Hani Elsayed Youssef). Address: London, United Kingdom. Date of birth: (a) 1.3.1961, (b) 16.6.1960. Place of birth: Qaylubiyah, Egypt. Nationality: Egyptian.'

- (3) The entry 'Sajid Mohammed **Badat** (alias (a) Abu Issa, (b) Saajid **Badat**, (c) Sajid **Badat**, (d) Muhammed **Badat**, (e) Sajid Muhammad **Badat**, (f) Saajid Mohammad **Badet**, (g) Muhammed **Badet**, (h) Sajid Muhammad **Badet**). Date of birth: (a) 28.3.1979, (b) 8.3.1976. Place of birth: Gloucester, United Kingdom. Passport No: (a) United Kingdom passport number 703114075, (b) United Kingdom passport number 026725401. Other information: Currently in custody in the United Kingdom. Previous address is Gloucester, United Kingdom' under the heading 'Natural persons' is replaced by:

'Sajid Mohammed **Badat** (alias (a) Abu Issa, (b) Saajid **Badat**, (c) Sajid **Badat**, (d) Muhammed **Badat**, (e) Sajid Muhammad **Badat**, (f) Saajid Mohammad **Badet**, (g) Muhammed **Badet**, (h) Sajid Muhammad **Badet**, (i) Sajid Mahomed **Badat**). Date of birth: (a) 28.3.1979, (b) 8.3.1976. Place of birth: Gloucester, United Kingdom. Passport No: (a) 703114075 (United Kingdom passport), (b) 026725401 (United Kingdom passport). Other information: Currently in custody in the United Kingdom. Previous address is Gloucester, United Kingdom.'

- (4) The entry 'Shamil BASAYEV (alias Abdullakh Shamil Abu-Idris); place of birth: Dyshni-Vedeno, Chechnya, Russian Federation; date of birth: 14 January 1965; Russian passport No 623334 (January 2002)' under the heading 'Natural persons' is replaced by:

'Shamil Salmanovich **Basayev** (alias Abdullakh Shamil Abu-Idris). Date of birth: 14.1.1965. Place of birth: Dyshni-Vedeno, Vedensk district, Chechen-Ingush Autonomous Soviet Socialist Republic, Soviet Union (Russian Federation). Nationality: Russian. Passport No: 623334 (Russian passport, January 2002). National identification No: IY-OZH No 623334 (issued on 9 June 1989 by the Vedensk district). Other information: International arrest warrant issued by the Russian Government.'

- (5) The entry 'Al Sayyid Ahmed Fathi **Hussein Eliwah** (alias (a) Al Sayyid Ahmed Fathi Hussein Eliwa, (b) Al Sayyid Ahmed Fathi Hussein Alaiwah, (c) Al Sayyid Ahmed Fathi Hussein Elaiwa, (d) Al Sayyid Ahmed Fathi Hussein Ilewah, (e) Al Sayyid Ahmed Fathi Hussein Alaywah, (f) El Sayed Ahmad Fathi Hussein Elaiwa, (g) Hatim, (h) Hisham, (i) Abu Umar). Date of birth: (a) 30.7.1964, (b) 30.1.1964. Place of birth: (a) Suez, Egypt, (b) Alexandria, Egypt. Nationality: Egyptian. Other information: lives in the United Kingdom' under the heading 'Natural persons' is replaced by:

'Al Sayyid Ahmed Fathi **Hussein Eliwah** (alias (a) Al Sayyid Ahmed Fathi Hussein Eliwa, (b) Al Sayyid Ahmed Fathi Hussein Alaiwah, (c) Al Sayyid Ahmed Fathi Hussein Elaiwa, (d) Al Sayyid Ahmed Fathi Hussein Ilewah, (e) Al Sayyid Ahmed Fathi Hussein Alaywah, (f) El Sayed Ahmad Fathi Hussein Elaiwa, (g) Hatim, (h) Hisham, (i) Abu Umar). Date of birth: (a) 30.7.1964, (b) 30.1.1964. Place of birth: (a) Suez, Egypt, (b) Alexandria, Egypt. Nationality: Egyptian. Passport No: RP0185179 (United Kingdom passport in the name of Al-Sayyid Ilewah, which was issued on 11.9.2001 and expires on 11.9.2011). Other information: lives in the United Kingdom.'

- (6) The entry 'Abdelghani **Mzoudi** (*alias* (a) Abdelghani Mazwati, (b) Abdelghani Mazuti). Address: op de Wisch 15, 21149 Hamburg, Germany. Date of birth: 6.12.1972. Place of birth: Marrakesh (Morocco). Nationality: Moroccan. Passport No: (a) Moroccan passport No F 879567, issued 29.4.1992 in Marrakesh, Morocco, valid until 28.4.1997, renewed until 28.2.2002; (b) Moroccan passport No M271392, issued 4.12.2000 by the Moroccan Embassy in Berlin, Germany. National identification No: Moroccan personal ID No E 427689, issued 20 March 2001 by the Moroccan Consulate General in Düsseldorf, Germany. Additional information: (a) remanded in custody in Germany (June 2003), (b) last registered at this address' under the heading 'Natural persons' is replaced by:

'Abdelghani **Mzoudi** (*alias* (a) Abdelghani Mazwati, (b) Abdelghani Mazuti). Address: op de Wisch 15, 21149 Hamburg, Germany. Date of birth: 6.12.1972. Place of birth: Marrakesh (Morocco). Nationality: Moroccan. Passport No: (a) F 879567 (Moroccan passport which was issued in Marrakesh, Morocco, on 29.4.1992, valid until 28.4.1997 and renewed until 28.2.2002), (b) M271392 (Moroccan passport issued on 4.12.2000 by the Moroccan Embassy in Berlin, Germany). National identification No: E 427689 (Moroccan personal ID, issued on 20.3.2001 by the Moroccan Consulate General in Düsseldorf, Germany). Other information: (a) last registered at this address, (b) after release from custody, he left Germany for Morocco in June 2005.'

- (7) The entry 'Mansour Thaer, date of birth: 21 March 1974; place of birth: Baghdad, Iraq' under the heading 'Natural persons' is replaced by:

'Mansour **Thaer**. Date of birth: 21.3.1974. Place of birth: Baghdad, Iraq. Other information: Deported from Germany to Jordan in February 2005.'

COMMISSION DIRECTIVE 2006/74/EC**of 21 August 2006****amending Council Directive 91/414/EEC to include dichlorprop-P, metconazole, pyrimethanil and triclopyr as active substances****(Text with EEA relevance)**

THE COMMISSION OF THE EUROPEAN COMMUNITIES,

Member State was Ireland and all relevant information was submitted on 21 November 2003.

Having regard to the Treaty establishing the European Community,

Having regard to Council Directive 91/414/EEC of 15 July 1991 concerning the placing of plant protection products on the market ⁽¹⁾, and in particular Article 6(1) thereof,

Whereas:

(1) Commission Regulations (EC) No 451/2000 ⁽²⁾ and (EC) No 703/2001 ⁽³⁾ lay down the detailed rules for the implementation of the second stage of the programme of work referred to in Article 8(2) of Directive 91/414/EEC and establish a list of active substances to be assessed, with a view to their possible inclusion in Annex I to Directive 91/414/EEC. That list includes dichlorprop-P, metconazole, pyrimethanil and triclopyr.

(2) For those active substances the effects on human health and the environment have been assessed in accordance with the provisions laid down in Regulations (EC) No 451/2000 and (EC) No 703/2001 for a range of uses proposed by the notifier. Moreover, those Regulations designate the rapporteur Member States which have to submit the relevant assessment reports and recommendations to the European Food Safety Authority (EFSA) in accordance with Article 8(1) of Regulation (EC) No 451/2000. For dichlorprop-P the rapporteur Member State was Denmark and all relevant information was submitted on 5 November 2003. For metconazole the rapporteur Member State was Belgium and all relevant information was submitted on 27 January 2004. For pyrimethanil the rapporteur Member State was Austria and all relevant information was submitted on 15 April 2004. For triclopyr the rapporteur

(3) The assessment reports have been peer reviewed by the Member States and the EFSA and presented to the Commission on 14 December 2005 for triclopyr and on 13 January 2006 for dichlorprop-P, metconazole and pyrimethanil in the format of the EFSA Scientific Reports ⁽⁴⁾. These reports have been reviewed by the Member States and the Commission within the Standing Committee on the Food Chain and Animal Health and finalised on 23 May 2006 in the format of the Commission review reports for dichlorprop-P, metconazole, pyrimethanil and triclopyr.

(4) It has appeared from the various examinations made that plant protection products containing dichlorprop-P, metconazole, pyrimethanil and triclopyr may be expected to satisfy, in general, the requirements laid down in Article 5(1)(a) and (b) of Directive 91/414/EEC, in particular with regard to the uses which were examined and detailed in the Commission review reports. It is therefore appropriate to include these active substances in Annex I, in order to ensure that in all Member States the authorisations of plant protection products containing these active substances can be granted in accordance with the provisions of that Directive.

(5) Without prejudice to that conclusion, it is appropriate to obtain further information on certain specific points concerning dichlorprop-P, pyrimethanil and triclopyr. Article 6(1) of Directive 91/414/EEC provides that inclusion of a substance in Annex I may be subject to conditions. Therefore it is appropriate to require that dichlorprop-P, pyrimethanil and triclopyr should be subjected to further testing for confirmation of the risk assessment for some issues and that such studies should be presented by the notifiers.

⁽¹⁾ OJ L 230, 19.8.1991, p. 1. Directive as last amended by Commission Directive 2006/64/EC (OJ L 206, 27.7.2006, p. 107).

⁽²⁾ OJ L 55, 29.2.2000, p. 25. Regulation as last amended by Regulation (EC) No 1044/2003 (OJ L 151, 19.6.2003, p. 32).

⁽³⁾ OJ L 98, 7.4.2001, p. 6.

⁽⁴⁾ EFSA Scientific Report (2006) 52, 1-67, Conclusion regarding the Peer review of the pesticide risk assessment of the active substance dichlorprop-P (finalised: 13 January 2006).

EFSA Scientific Report (2006) 64, 1-71, Conclusion regarding the Peer review of the pesticide risk assessment of the active substance metconazole (finalised: 13 January 2006).

EFSA Scientific Report (2006) 61, 1-70, Conclusion regarding the Peer review of the pesticide risk assessment of the active substance pyrimethanil (finalised: 13 January 2006).

EFSA Scientific Report (2005) 56, 1-103, Conclusion regarding the peer review of the pesticide risk assessment of the active substance triclopyr (finalised: 14 December 2005).

- (6) A reasonable period should be allowed to elapse before an active substance is included in Annex I in order to permit Member States and the interested parties to prepare themselves to meet the new requirements which will result from the inclusion.
- (7) Without prejudice to the obligations defined by Directive 91/414/EEC as a consequence of including an active substance in Annex I, Member States should be allowed a period of six months after inclusion to review existing authorisations of plant protection products containing dichlorprop-P, metconazole, pyrimethanil and triclopyr to ensure that the requirements laid down by Directive 91/414/EEC, in particular in its Article 13 and the relevant conditions set out in Annex I, are satisfied. Member States should vary, replace or withdraw, as appropriate, existing authorisations, in accordance with the provisions of Directive 91/414/EEC. By way of derogation from the above deadline, a longer period should be provided for the submission and assessment of the complete Annex III dossier of each plant protection product for each intended use in accordance with the uniform principles laid down in Directive 91/414/EEC.
- (8) The experience gained from previous inclusions in Annex I to Directive 91/414/EEC of active substances assessed in the framework of Commission Regulation (EEC) No 3600/92⁽¹⁾ has shown that difficulties can arise in interpreting the duties of holders of existing authorisations in relation to access to data. In order to avoid further difficulties it therefore appears necessary to clarify the duties of the Member States, especially the duty to verify that the holder of an authorisation demonstrates access to a dossier satisfying the requirements of Annex II to that Directive. However, this clarification does not impose any new obligations on Member States or holders of authorisations compared to the directives which have been adopted until now amending Annex I.
- (9) It is therefore appropriate to amend Directive 91/414/EEC accordingly.
- (10) The measures provided for in this Directive are in accordance with the opinion of the Standing Committee on the Food Chain and Animal Health,

HAS ADOPTED THIS DIRECTIVE:

Article 1

Annex I to Directive 91/414/EEC is amended as set out in the Annex to this Directive.

Article 2

Member States shall adopt and publish by 30 November 2007 at the latest the laws, regulations and administrative provisions necessary to comply with this Directive. They shall forthwith communicate to the Commission the text of those provisions and a correlation table between those provisions and this Directive.

They shall apply those provisions from 1 December 2007.

When Member States adopt those provisions, they shall contain a reference to this Directive or be accompanied by such a reference on the occasion of their official publication. Member States shall determine how such reference is to be made.

Article 3

1. Member States shall in accordance with Directive 91/414/EEC, where necessary, amend or withdraw existing authorisations for plant protection products containing dichlorprop-P, metconazole, pyrimethanil and triclopyr as active substances by 30 November 2007.

By that date they shall in particular verify that the conditions in Annex I to that Directive relating to dichlorprop-P, metconazole, pyrimethanil and triclopyr are met, with the exception of those identified in part B of the entry concerning that active substance, and that the holder of the authorisation has, or has access to, a dossier satisfying the requirements of Annex II to that Directive in accordance with the conditions of Article 13 of that Directive.

2. By way of derogation from paragraph 1, for each authorised plant protection product containing dichlorprop-P, metconazole, pyrimethanil and triclopyr as either the only active substance or as one of several active substances all of which were listed in Annex I to Directive 91/414/EEC by 31 May 2007 at the latest, Member States shall re-evaluate the product in accordance with the uniform principles provided for in Annex VI to Directive 91/414/EEC, on the basis of a dossier satisfying the requirements of Annex III to that Directive and taking into account part B of the entry in Annex I to that Directive concerning dichlorprop-P, metconazole, pyrimethanil and triclopyr respectively. On the basis of that evaluation, they shall determine whether the product satisfies the conditions set out in Article 4(1)(b), (c), (d) and (e) of Directive 91/414/EEC.

⁽¹⁾ OJ L 366, 15.12.1992, p. 10). Regulation as last amended by Regulation (EC) No 2266/2000 (OJ L 259, 13.10.2000, p. 27).

Following that determination Member States shall:

- (a) in the case of a product containing dichlorprop-P, metconazole, pyrimethanil and triclopyr as the only active substance, where necessary, amend or withdraw the authorisation by 31 May 2011 at the latest; or
- (b) in the case of a product containing dichlorprop-P, metconazole, pyrimethanil and triclopyr as one of several active substances, where necessary, amend or withdraw the authorisation by 31 May 2011 or by the date fixed for such an amendment or withdrawal in the respective Directive or Directives which added the relevant substance or substances to Annex I to Directive 91/414/EEC, whichever is the latest.

Article 4

This Directive shall enter into force on 1 June 2007.

Article 5

This Directive is addressed to the Member States.

Done at Brussels, 21 August 2006.

For the Commission

Markos KYPRIANOU

Member of the Commission

ANNEX

The following entries shall be added at the end of the table in Annex I to Directive 91/414/EEC:

No	Common name, identification numbers	IUPAC name	Purity ⁽¹⁾	Entry into force	Expiration of inclusion	Specific provisions
135	Dichlorprop-P CAS No 15165-67-0 CIPAC No 476	(R)-2-(2,4-dichlorophenoxy) propanoic acid	≥ 900 g/kg	1 June 2007	31 May 2017	PART A Only uses as herbicide may be authorised. PART B For the implementation of the uniform principles of Annex VI, the conclusions of the review report on dichlorprop-P, and in particular Appendices I and II thereof, as finalised in the Standing Committee on the Food Chain and Animal Health on 23 May 2006 shall be taken into account. In this overall assessment Member States: — must pay particular attention to the protection of birds, mammals, aquatic organisms and non-target plants. Conditions of authorisation should include risk mitigation measures, where appropriate. The Member States concerned shall request the submission of further studies to confirm the results on animal metabolism and the risk assessment on acute and short-term exposure for birds and on acute exposure for herbivorous mammals. They shall ensure that the notifiers at whose request dichlorprop-P has been included in this Annex provide such studies to the Commission within two years of the entry into force of this Directive.
136	Metconazole CAS No 125116-23-6 (unstated stereo-chemistry) CIPAC No 706	(1RS,5RS:1RS,5SR)-5-(4-chlorobenzyl)-2,2-dimethyl-1-(1H-1,2,4-triazol-1-ylmethyl) cyclopentanol	≥ 940 g/kg (sum of <i>cis</i> - and <i>trans</i> -isomers)	1 June 2007	31 May 2017	PART A Only uses as fungicide may be authorised. PART B For the implementation of the uniform principles of Annex VI, the conclusions of the review report on metconazole, and in particular Appendices I and II thereof, as finalised in the Standing Committee on the Food Chain and Animal Health on 23 May 2006 shall be taken into account. In this overall assessment: — Member States must pay particular attention to the protection of aquatic organisms, birds and mammals. Conditions of authorisation should include risk mitigation measures, where appropriate, — Member States must pay particular attention to the operator safety. Conditions of authorisation should include protective measures, where appropriate.

No	Common name, identification numbers	IUPAC name	Purity (1)	Entry into force	Expiration of inclusion	Specific provisions
137	Pyrimethanil CAS No 53112-28-0 CIPAC No not allocated	N-(4,6- dimethylpyrimidin-2-yl) aniline	≥ 975 g/kg (the manufacturing impurity cyanamide is considered to be of toxicological concern and must not exceed 0,5 g/kg in the technical material)	1 June 2007	31 May 2017	PART A Only uses as fungicide may be authorised. PART B For the implementation of the uniform principles of Annex VI, the conclusions of the review report on pyrimethanil, and in particular Appendices I and II thereof, as finalised in the Standing Committee on the Food Chain and Animal Health on 23 May 2006 shall be taken into account. In this overall assessment Member States: — must pay particular attention to the protection of aquatic organisms. Conditions of authorisation should include risk mitigation measures, where appropriate, such as buffer zones, — must pay particular attention to the operator safety and ensure that conditions of use prescribe the application of adequate personal protective equipment. The Member States concerned shall request the submission of further studies to confirm the risk assessment to fish. They shall ensure that the notifiers at whose request pyrimethanil has been included in this Annex provide such studies to the Commission within two years from the entry into force of this Directive.

No	Common name, identification numbers	IUPAC name	Purity ⁽¹⁾	Entry into force	Expiration of inclusion	Specific provisions
138	Triclopyr CAS No 055335-06-3 CIPAC No 376	3,5,6-trichloro-2-pyridyloxyacetic acid	≥ 960 g/kg (as Triclopyr butoxyethyl ester)	1 June 2007	31 May 2017	PART A Only uses as herbicide may be authorised. PART B In assessing applications to authorise plant protection products containing triclopyr for uses other than spring applications in pasture and grassland, Member States shall pay particular attention to the criteria in Article 4(1)(b), and shall ensure that any necessary data and information is provided before such an authorisation is granted. For the implementation of the uniform principles of Annex VI, the conclusions of the review report on triclopyr, and in particular Appendices I and II thereof, as finalised in the Standing Committee on the Food Chain and Animal Health on 23 May 2006 shall be taken into account. In this overall assessment Member States: — must pay particular attention to the protection of groundwater under vulnerable conditions. Conditions of authorisation should include risk mitigation measures and monitoring programmes should be initiated in vulnerable zones, where appropriate, — must pay particular attention to the safety of operators and ensure that conditions of use prescribe the application of adequate personal protective equipment, — must pay particular attention to the protection of birds, mammals, aquatic organisms and non-target plants. Conditions of authorisation should include risk mitigation measures, where appropriate. The Member States concerned shall request the submission of further studies to confirm the acute and long-term risk assessment for birds and mammals and the risk to aquatic organisms from exposure to the metabolite 6-chloro-2-pyridinol. They shall ensure that the notifiers at whose request triclopyr has been included in this Annex provide such studies to the Commission within two years from the entry into force of this Directive.'

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

CORRIGENDA**Corrigendum to Council Regulation (EC) No 318/2006 of 20 February 2006 on the common organisation of the markets in the sugar sector**

(Official Journal of the European Union L 58 of 28 February 2006)

On page 16 in Article 40(2)(f)(i):

for: '(i) amendments to the definition provided for in Article 2(11);',

read: '(i) amendments to the definition provided for in Article 2(12);'

and on Page 24 in Annex III, 'National and regional quotas', in the column headed 'Isoglucose', against 'Total':

for: '507 680',

read: '507 681'.
