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(Amendment of Regulation (EC) No 1774/2002)

HAY ACUERDO TÉCNICO UNÁNIME.

COMMISSION OF THE EUROPEAN COMMUNITIES

Brussels,
C(2005) final

Draft

COMMISSION REGULATION

of

amending Annexes I, II, VII, VIII, X and XI to Regulation (EC) No 1774/2002 of the European Parliament and of the Council as regards the placing on the market of certain animal by-products

(Text with EEA relevance)

(Memorandum from Mr M. KYPRIANOU)

Draft

COMMISSION REGULATION

of

amending Annexes I, II, VII, VIII, X and XI to Regulation (EC) No 1774/2002 of the European Parliament and of the Council as regards placing on the market of certain animal by-products

(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 1774/2002 of the European Parliament and of the Council of 3 October 2002 laying down health rules concerning animal by-products not intended for human consumption[1] and in particular the second paragraph of Article 28, the first subparagraph of Article 29(3) and Article 32(1),

Whereas:

(1) Regulation (EC) No 1774/2002 lays down animal and public health requirements for the importation into and transit through the Community of certain animal by-products and products derived there from. That Regulation provides for general hygiene requirements for the processing of Category 1, 2 and 3 materials and also sets out requirements for the placing on the market of those animal by-products and products derived there from including model health certificates for their importation into the Community.

(2) Regulation (EC) No 999/2001 of the European Parliament and of the Council of 22 May 2001 laying down rules for the prevention, control and eradication of certain transmissible spongiform encephalopathies[2] applies to the production and placing on the market of live animals and products of animal origin. It prohibits the feeding of processed animal protein to farmed animals which are kept, fattened or bred for the production of food.

(3) While that prohibition in Regulation (EC) No 999/2001 remains in force, less stringent processing requirements as laid down in Chapter II of Annex VII to Regulation (EC) No 1774/2002 than the current processing Method 1 should apply to processed animal

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1 OJ L 273, 10.10.2002, p. 1. Regulationas last amended by Commission Regulation (EC) No 208/2006 ~~416/2005~~ (OJ L 36, 8.2.2006, p. 25 ~~66, 12.3.2005, p. 10~~).

2 OJ L 147, 31.5.2001, p. 1. Regulation as last amended by Commission Regulation (EC) No 688/2006 ~~1974/2005~~ (OJ 120, 5.5.2006, p. 10 ~~No. L 317, 03.12.2005, p. 4~~).

protein derived from porcine blood, as there is no scientific indication of a risk of transmitting TSEs from pigs. It is necessary however for public and animal health reasons to introduce a minimum temperature requirement for the processing of porcine blood. Chapter II of Annex VII to Regulation (EC) No 1774/2002 should therefore be amended accordingly

(4) The relevant health certificate set out in Chapter I of Annex X to Regulation (EC) No 1774/2002 for imports into the Community of processed animal protein, not intended for human consumption, and products other than petfood containing such protein should be amended accordingly in order to provide for equivalent processing possibilities for third countries as for the Community. Chapter I of Annex X to Regulation (EC) No 1774/2002 should therefore be amended accordingly.

(5) Chapter 3 (B) of Annex X to Regulation (EC) No 1774/2002 sets out the model health certificate for the importation into the Community of processed pet food, other than canned pet food. It appears that processed pet food is increasingly produced in third countries by mixing already processed ingredients which have individually been treated in compliance with the requirements of that Regulation. Heat treatment of such ingredients could have adverse effects on their nutritional qualities. Therefore, the importation into the Community of processed petfood consisting of such safe ingredients should be allowed and the model health certificate set out in Chapter 3 (B) of Regulation (EC) No 1774/2002 should be amended accordingly.

(6) Regulation (EC) No 1774/2002 provides that certain animal by-products for the manufacture of petfood may be imported into the Community if they contain material which has been derived from animals treated with certain prohibited substances. However, the importation into the Community of processed petfood, dogchews and flavouring innards containing such material is currently not allowed. Since it appears that the health risk is not higher when importing such material for the production of petfood within the Community than in case such material is an ingredient of imported processed pet food, dogchews and flavouring innards, the importation into the Community of these processed products containing such material should also be allowed. Therefore, the model health certificates set out in Chapters 3(A), 3(B), 3(C) and 3(E) of Annex X to Regulation (EC) No 1774/2002 should be amended accordingly.

(7) Chapter 3(B) of Annex X to Regulation (EC) No 1774/2002 provides that the health certificate for processed pet food, other than canned petfood, must contain a certificate stating that the pet food was packed in new packaging, which bears labels indicating "not intended for human consumption". Council Directive 79/373/EEC of 2 April 1979 on the circulation of compound feedingstuffs[3] already provides for equivalent marking requirements which prevent unintentional or negligent misuse of such products for human consumption. The requirements for the model health certificate in Chapter 3(B) of Annex X to Regulation (EC) No 1774/2002 should take account of that legislation and should, therefore, be amended accordingly.

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[3] OJ L 86, 6.4.1979, p. 30. Directive as last amended by Regulation (EC) No 807/2003 (OJ L 122, 16.5.2003, p. 36). ~~Directive 2002/2/EC of the European Parliament and the Council (OJ L 63, 6.3.2002, p. 23).~~

(8) Annex VIII to Regulation (EC) No 1774/2002 lays down requirements for the placing on the market and importation into the Community of pet food, dogchews and technical products. Chapter II B(4) of that Annex provides that dogchews must have undergone a heat treatment during processing sufficient to destroy pathogenic organisms. Several Member States have requested to allow other treatments, during processing of dogchews, which provide equivalent safety guarantees. Therefore, Chapter II B (4) of Annex VIII to Regulation (EC) should be amended accordingly.

(9) Chapter 3 (C) of Annex X to Regulation (EC) No 1774/2002 sets out the model health certificate for the importation into the Community of dogchews. Since other treatment requirements than heat treatment shall be allowed for the processing of dogchews within the Community, those other treatments should also be allowed for imported dogchews. Some dogchews are produced according to traditional methods from fish. Animal health considerations do not require a heat treatment in such case. Therefore, the model health certificate in Chapter 3 (C) of Annex X to that Regulation should be amended accordingly.

(10) Chapter II A (1) of Annex VIII to Regulation (EC) No 1774/2002 permits the use of parts of slaughtered animals which are fit for human consumption but are not intended for human consumption for commercial reasons in the manufacture of raw petfood within the Community. It appears that parts of such animals which are rejected as unfit for human consumption but which are not affected by any signs of disease communicable to humans or animals may also be safely used in petfood. Chapter II A (1) and Chapter XI of Annex VIII to that Regulation, the latter providing for the respective provisions for the importation into the Community, should therefore be amended accordingly.

(11) Chapter II B (6) of Annex VIII to Regulation (EC) No 1774/2002 lays down microbiological standards applicable to raw petfood produced in the Community. Those standards should also be imposed on imports into the Community of raw petfood for direct sale and on animal by-products to be fed to farmed fur animals intended for dispatch to the Community. Chapter 3 (D) of Annex X of that Regulation should therefore be amended accordingly.

(12) Annex XI to Regulation (EC) No 1774/2002 sets out lists of third countries from which Member States may authorise imports of animal by-products not intended for human consumption. Part XIII of that Annex provides for a list of third countries from which Member States may authorise the imports of serum of equidae. According to that part XIII, this list includes third countries referred to in Annex I to Commission Decision 2004/211/EC of 6 January 2004 establishing the list of third countries and parts of territory thereof from which Member States authorise imports of live equidae and semen, ova and embryos of the equine species, and amending Decisions 93/195/EECand 94/63/EC[4], from which the importation of equidae for slaughter is allowed.

(13) Serum of equidae may, however, be sourced from living animals. Therefore, the animal health status allowing for the inclusion of a third country in that list of third countries from which the importation of equidae for breeding and production is

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[4] OJ L 73, 11.03.2004, p. 1.

allowed should be sufficient to determine whether serum of equidae may be imported from this third country. That would, in particular, allow for the importation of serum of equidae from Mexico. Part XIII of Annex XI to Regulation (EC) No 1774/2002 should therefore be amended accordingly.

(14) It is necessary to amend the existing model health certificates for the importation of hides and skins into the Community, in order to introduce some technical amendments. In the interest of clarity, the specific requirements for those products set out in Chapter VI of Annex VIII, and the model health certificates set out in Chapters 5 (A), 5(B) and 5(C) of Annex X to Regulation (EC) No 1774/2002 should therefore be amended accordingly.

(15) Chapter X of Annex II to Regulation (EC) No 1774/2002 lays down a model commercial document to accompany animal by-products and processed products during transportation. In order to improve the traceability of hides and skins during transportation, it is necessary to amend that model document. Annex II to that Regulation should therefore be amended accordingly.

(16) The European Food Safety Authority (EFSA) in its *Scientific Opinion on Animal health and welfare aspects of avian influenza,* adopted on 13 and 14 September 2005, concluded that feathers should be treated before trade, in order to reduce the risk of the spread of avian influenza. Chapter VIII of Annex VIII to Regulation (EC) No 1774/2002 lays down the permanent Community measures concerning the placing of the market and importation of feathers. In the light of that opinion and the current worldwide epidemiological situation regarding avian influenza, it is appropriate to amend the relevant Community measures for the importation and treatment of feathers prior to importation. Annex VIII to that Regulation should therefore be amended accordingly.

(17) Due to the threat of introduction of the small hive beetle into the Community, which is currently free of that parasite, it is necessary to lay down requirements for the importation of beeswax, intended for technical purposes, and also to amend the definition of apiculture products in Annex I to Regulation (EC) No 1774/2002. Therefore, the relevant definition in Annex I to that Regulation, the import requirements laid down in Chapter IX of Annex VIII and the model health certificate set out in Chapter 13 of Annex X to that Regulation should be amended accordingly.

(18) A transitional period should be provided for after the date of entry into force of this Regulation, in order to allow for the continued importation into the Community of the animal by-products and products derived there from covered by Regulation (EC) No 1774/2002 and accompanied by health certificates in compliance with that Regulation

(19) 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

Annexes I, II, VII, VIII, X and XI to Regulation (EC) No 1774/2002 are amended in accordance with the Annex to this Regulation.

Article 2

For a transitional period of six months from the date of entry into force of this Regulation, the model health certificates set out in Chapter 1, Chapters 3(A) to 3(E), Chapters 5(A), 5(B) and 5(C) and Chapter 16 of Annex X to Regulation (EC) No 1774/2002 which where completed in conformity with the provisions applicable before the date of entry into force of the present Regulation, may continue to accompany the products covered by such certificates, as appropriate.

Article 3

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

It shall apply from [xxxx].

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

Done at Brussels,

For the Commission
Markos KYPRIANOU
Member of the Commission

ANNEX

The Annexes to Regulation (EC) No 1774/2002 are amended as follows:

(1) Annex I is amended as follows:

(a) Point 1 is replaced by the following:

'1. 'apiculture by-products' means honey, beeswax, royal jelly, propolis or pollen not intended for human consumption;'

(b) Point 42 is replaced by the following:

'42. 'processed animal protein' means animal protein derived entirely from Category 3 material, which have been treated in accordance with Chapter II of Annex VII so as to render them suitable for direct use as feed material or for any other use in feedingstuffs, including petfood, or for use as and/or in organic fertilisers or soil improvers; however, it does not include blood products, milk, milk-based products, colostrum, gelatine, hydrolysed proteins and dicalcium phosphate, eggs and egg-products, tricalcium phosphate and collagen;'

(2) In Annex II, Chapter X is replaced by the following:

'CHAPTER X

Commercial document

1. The following commercial document shall accompany animal by-products and processed products during transportation. However, Member States may decide to use a different commercial document, in paper or in electronic form, for animal by-products and processed products transported within the same Member State provided that such commercial document specifies the requirements laid down in Chapter III(2).

2. Where more than one transporter is involved, each transporter shall fill in a declaration as referred to in point 7 of the commercial document, which shall be part of the document.

MODEL COMMERCIAL DOCUMENT
FOR THE TRANSPORTATION WITHIN THE EUROPEAN COMMUNITY OF ANIMAL BY-PRODUCTS AND PROCESSED PRODUCTS

Notes

(a) Commercial documents shall be produced, according to the layout of the model appearing in this Annex. It shall contain, in the numbered order that appears in the model, the attestations that are required for the transportation of animal by-products and processed products derived there from.

(b) It shall be drawn up in one of the official languages of the Member State of origin or the Member State of destination, as appropriate. However, it may also be drawn up in other official Community languages, if accompanied by an official translation or if previously agreed by the competent authority of the Member State of destination.

(c) The commercial document must be produced at least in triplicate (one original document and two copies). The original document must accompany the consignment to its final destination. The receiver must retain it. The producer must retain one of the copies and the carrier the other.

(d) The original of each commercial document shall consist of a single page, both sides, or, where more text is required it shall be in such a form that all pages needed are part of an integrated whole and indivisible.

(e) If for reasons of identification of the items of the consignment, additional pages are attached to the commercial document, these pages shall also be considered as forming part of the original document by the application of the signature of the person responsible for the consignment, on each of the pages.

(f) When the commercial document, including additional pages referred to in (e), comprises more than one page, each page shall be numbered -*(page number)* of *(total number of pages)*– at the bottom and shall bear the code number of the document that has been designated by the responsible person at the top.

(g) The original of the commercial document must be completed and signed by the responsible person. In doing so, the responsible person shall ensure that the principles of documentation as laid down in Chapter III of Annex II to Regulation (EC) No 1774/2002 are followed. The commercial document must specify:

(i) the date on which the material was taken from the premises;

(ii) the description of the material, including the identification of the material, the animal species for Category 3 material and processed products derived therefrom destined for use as feed material and, if applicable, the ear-tag number of the animal;

(iii) the quantity of the material;

(iv) the place of origin of the material;

(v) the name and the address of the carrier of the material;

(vi) the name and the address of the receiver and, if applicable, its approval number; and

(vii) if appropriate, the approval number of the plant of origin, and the nature and the methods of the treatment.

(h) The colour of the signature of the responsible person shall be different to that of the printing.

(i) The commercial document must be kept for a period of at least two years for presentation to the competent authority to verify the records referred to in Article 9 of Regulation (EC) No 1774/2002.

(j) Where Member States decide to use a commercial document in electronic form, the requirements listed in points (a) to (i) shall be shall be complied with as appropriate for such electronic form.

Commercial document

For the transportation within the European Community of animal by-products and processed products not intended for human consumption in accordance with Regulation (EC) No 1774/2002([1])

EUROPEAN COMMUNITY **Intra trade document**

Part I : Details of consignment presented

I.1. Consignor
Name
Address
Postal code

I.2. Document reference number | I.2.a.Local reference number:

I.3. Central Competent Authority

I.4. Local Competent Authority

I.5. Consignee
Name
Address
Postal code

I.6.

I.7.

I.8.Country of origin ISO code | I.9. Region of origin Code

I.10. Country of destination ISO code | I.11. Region of destination Code

I.12. Place of origin
Establishment
Name Approval number
Address
Postal code

I.13. Place of destination
Establishment Other
Name Approval number
Address
Postal code

I.14. Place of loading
Postal code

I.15. Date and time of departure

I.16. Means of transport
Aeroplane Ship Railway wagon
Road vehicle Other
Identification:

I.17. Transporter
Name Approval number
Address
Postal code Member State

I.18. Description of commodity

I.19. Commodity code (CN code)

I.20. Number/quantity

I.21 Temperature of products
Ambient Chilled Frozen

I.22. Number of packages

I.23. Identification of container/seal number

I.24.Type of packaging

I.25. Commodities certified for
Animal feedingstuff Technical use Other

I.26. Transit through third country
Third country ISO code
Exit point Code
Entry point BIP unit no.:

I.27. Transit through Member States
Member State ISO code
Member State ISO code
Member State ISO code

I.28. Export
Third country ISO code
Exit point Code

I.29.

I.30.

I.31. Identification of the commodities

Species (Scientific name)	Nature of commodity	Category	Treatment type	Approval number of establishments: Manufacturing plant	Batch number

EUROPEAN COMMUNITY

Animal by-products and processed products not intended for human consumption

Part II: Declaration

II.a. Certificate reference number	II..b. Local reference number

II.1. Declaration by the consignor

I, the undersigned, declare that:

II.1.1. A label attached to the container/carton/other packaging material carries the following indication (1):

(a) the Category of the animal by-products (see box reference I.31: Category).....

(b) in the case of processed products, the Category of animal by-products from which the processed products were derived (see box reference I.31: Category):.....

(c) (i).....in the case of Category 3 material, the words "not for human consumption"

(ii) in the case of Category 2 material, other than manure and digestive tract content and processed products derived therefrom, the words "not for animal consumption",

(iii) in the case of Category 2 material intended for feeding of animals referred to in point (c) of Article 23(2) under the conditions provided for in that Article of Regulation (EC) No 1774/2002 (2), the words "for feeding to ..." completed with the name of the specific species of those animal(s) for the feeding of which the material is intended.

(iv) in the case of manure and digestive tract content, the word "manure"; or

(v) in the case of Category 1 material and processed products derived therefrom, the words "for disposal only"

II.1.2. in the case where the packaging is done by the consignor the animal by-products and/or processed products are:

(1) **either** [in sealed new packaging;]

(1) or [transported in bulk in covered leak-proof containers or vehicles or other means of transport that were thoroughly cleaned and dry before use;]

II.1.3. in the case where treatment is applicable,

(a) the hides and skins have been treated in accordance with 'note Part I, box reference I.31: Treatment type, point 2' to this document: and

(b) the consignment has not been in contact with other animal products or live animals presenting a risk

of spreading a serious transmissible disease;

II.1.4. the animal by-products and/or processed products were stored properly prior to loading and dispatch;

II.1.5. all precautions have been taken to avoid contamination of the animal by-products or processed products with pathogenic agents and cross-contamination between various Categories.

Notes

Part I:

- Box reference I.9 and I.11: if appropriate.
- Box reference I.14: complete if different from 'I.1. Consignor'.
- Box reference I.31:

Animal species: For Category 3 material and processed products derived therefrom destined for use as feed material.

Nature of commodity: Enter unprocessed animal by-product or processed product chosen among the following list: apiculture products', 'blood products', 'blood', 'bloodmeal', 'canned petfood', 'digestion residues', 'digestive tract content', 'dogchews', 'fishmeal', 'gelatin', 'greaves', 'hides and skins', 'hydrolysed proteins', 'organic fertilizers', 'petfood', 'processed animal protein', 'processed petfood', 'processed products', 'raw petfood', 'rendered fats'.

Category: Categories 1, 2 or 3. In case of Category 3, specify which letter from 3a to 3k (as under Article 6, paragraph 1 described below).

In the case of animal by-product for use in raw petfood indicate 3a or 3b whether the animal by-products derive from:

Category 3a, Article 6(1)(a) i.e. parts of slaughtered animals, which were fit for human consumption in accordance with Community legislation, but are not intended for human consumption for commercial reasons, or

Category 3b, Article 6(1)(b) i.e. parts of slaughtered animals, which are rejected as unfit for human consumption but are not affected by any signs of diseases communicable to humans or animals and derive from carcasses that are fit for human consumption in accordance with Community legislation;

In the case of hides and skins and processed products derived there from, indicate 3c or 3k whether the animal by-products derive from:

Category 3c, Article 6(1)(c) i.e. hides and skins originating from animals that are slaughtered in a slaughterhouse, after undergoing ante-mortem inspection, an were fit, as a result of such inspection, for slaughter for human consumption in accordance with Community legislation; or

Category 3k, Article 6(1)(k) i.e. hides and skins originating from animals that did not show clinical signs of any disease communicable through that product to humans or animals

Where the consignment is made of more than one Category, indicate the quantity and if applicable the number of containers per Category of materials.

Treatment type: For treated hides and skins, which (a) are not fulfilling the requirements of Regulation (EC) No 853/2004 of 29 April 2004 on the hygiene of foodstuffs (OJ No L226, 25.6.2004, p. 3) or (b) have not undergone the complete process of tanning or (c) are not "wet blue"; or (d) are not "pickled pelts" or (e)are not limed (treated with lime and in brine at a pH of 12 to 13 for et least eight hours): enter treatment among the following: (a) dried; (b) dry-salted or wet-salted for at least 14 days prior to dispatch; (c) salted for seven days in sea salt with the addition of 2 % sodium carbonate; or (d) preserved by a process other than tanning specified in accordance with the procedure referred to in Article 33(2) of Regulation (EC) No 1774/2002.

For Category 3 materials and processed products derived therefrom destined for use as feed: if appropriate describe the nature and the methods of the treatment.

Batch number: enter batch number or ear tag number if applicable.

Part II:

(1) Delete as appropriate.

(2) OJ L 273, 10.10.2002, p. 1.

- The signature must be in a different colour to that of the printing

Signature

Done at .....on .....
(place) (date)

.....
(signature of the responsible person/consignor)

.....
(name, in capital letters)

Declaration by the transporter

I, the undersigned, declare that:

II.2.1. in the case where the packaging is done by the transporter, the animal by-products and/or processed products are:

(1) **either** [in sealed new packaging;]

(1) or [transported in bulk in covered leak-proof containers or vehicles or other means of transport that were thoroughly cleaned and dry before use and cleaned, washed and disinfected after each use]

II.2.2. all precautions have been taken:

– to avoid contamination of the animal by-products or processed products with pathogenic agents and cross-contamination between various Categories during transportation; and

– to ensure transportation under appropriate temperature to avoid risk to animal or public health

Notes

Part II:

(1) Delete as appropriate.

- The signature must be in a different colour to that of the printing
- Note for the transporters: This document must accompany the consignment* from the place of loading for dispatch until it reaches the point of destination.

  * "Consignment" means "a quantity of products of the same type, which may contain different Categories of animal by-products, coming from the same consignor and covered by the same commercial document conveyed by the same means of transport to the same recipient"

Signature

Done at .....on .....
(place) (date)

.....
(signature of the responsible person/transporter)

.....
(name, in capital letters)

(3) In Annex VII, Chapter II, paragraph A(1), the first subparagraph is replaced by the following:

'Mammalian processed animal protein must have been submitted to processing method 1. However, porcine blood may be submitted to any of processing methods 1 to 5 or to processing method 7 provided that in the case of processing method 7, a heat treatment throughout its substance at a minimum temperature of 80 °C has been applied.'

(4) Annex VIII is amended as follows:

(a) Chapter II is amended as follows:

(i) Paragraph A(1) is replaced by the following:

'1. The only animal by-products that may be used to produce petfood and dogchews are those referred to in Article 6 (1) (a) to (j). However, raw petfood may only be manufactured from animal by-products referred to in Article 6 (1) (a) or Article 6 (1) (b).'

(ii) Paragraph B(4) is replaced by the following:

'4. Dogchews must be subjected to a treatment during processing sufficient to destroy pathogenic organisms, including salmonella.

After that treatment, every precaution must be taken to ensure that such dogchews are not exposed to contamination. The dogchews must be packed in new packaging.'

(b) Chapter III is amended as follows:

(i) Paragraph I(A)(3) is replaced by the following:

'3. Unprocessed manure of equidae which is traded must not originate from a holding subject to animal health restrictions pertaining to glanders, vesicular stomatitis, anthrax or rabies in accordance with Article 4 (5) of Directive 90/426/EEC.'

(ii) Paragraph I(B) is replaced by the following:

'The importation of unprocessed manure shall be prohibited.'

(iii) Paragraph II(B)(6)(d) is replaced by the following:

'(d) are accompanied by a health certificate that conforms to the model laid down in Chapter 17 of Annex X.'

(c) Chapter VI is amended as follows:

(i) In A, paragraph (1)(a) is replaced by the following:

'(a) to hides and skins of ungulates complying with the requirements of Regulation (EC) No 853/2004 of the European Parliament and of the Council laying down specific hygiene rules for food of animal origin (*)

(*) OJ L 139, 30.4.2004, p. 55; corrected version (OJ L 226, 25.6.2004, p. 22)'

(ii) In B, paragraph 3 is replaced by the following:

'3. Trade in fresh or chilled hides and skins is subject to the same health conditions as those applicable to fresh meat pursuant to Council Directive 2002/99/EC laying down the animal health rules governing the production, processing, distribution and introduction of products of animal origin for human consumption (**)

(**) *OJ L 18, 23.1.2003, p. 11.*'

(iii) In C, paragraph 5(b), the introductory phrase is replaced by the following:

'(b) they come from a third country or, in the case of regionalisation in accordance with Community legislation, from a part of a third country, appearing on the list set out in Part XIV(A) of Annex XI and which, as appropriate to the species concerned:'

(iv) In C, paragraphs 6(b) to (e) are replaced by the following:

'(b) they come either from:

(i) a third country or, in the case of regionalisation in accordance with Community legislation, from a part of a third country, appearing on the list set out in Part XIV(B) of Annex XI from which imports of fresh meat of the corresponding species are authorised and they have been treated in accordance with paragraph 2(a), (b) and (c) of A; or

(ii) a third country appearing on the list set out in Part XIV(B) of Annex XI and they have been treated in accordance with paragraph 2 (c) or (d) of A; or

(iii) equidae or ruminant animals from a third country appearing on the list set out in Part XIV (C) of Annex XI, which have been treated in accordance with paragraph 2(a), (b) and (c) of A and after treatment have been kept separate for at least 21 days.

(c) in the case of salted hides and skins transported by ship, they have been treated in accordance with paragraphs 2 (b) or (c) of A and have been kept separated after treatment during transportation for at least 14 days in the case of paragraph (b) or seven days in the case of paragraph (c) before importation and the health certificate accompanying the consignment attests such treatment and the duration of the transportation; and

(d) a health certificate conforming to the model health certificate laid down in Chapter 5 (B) of Annex X, or, in the case of hides and skins referred to in paragraph 6(b) (iii) of C of this Annex, an official declaration conforming to the model laid down in Chapter 5 (C) of Annex X, accompanies them.'

(d) Chapter VIII is amended as follows:

(i) In B, paragraph 4, the introductory phrase is replaced by the following:

'Member States must authorise the importation of unprocessed wool and hair, if they are:'

(ii) In B, the following paragraph 5 is added:

'5. The importation of unprocessed feathers and parts of feathers is prohibited.

Member States must authorise the importation of processed feathers and parts of feathers if:

(a) they are treated decorative feathers, treated feathers carried by travellers for their private use or consignments of treated feathers sent to private individuals for non-industrial purposes; or

(b) they are accompanied by a commercial document stating that the feathers or parts of feathers have been treated with a steam current or by another method ensuring the inactivation of pathogens and are securely enclosed in packaging and dry.'

(e) Chapter IX is amended as follows:

(i) In A, paragraph 1, the introductory phrase is replaced by the following:

'1. Apiculture by-products intended exclusively for use in apiculture must:'

(ii) In B, paragraph 3 is replaced by the following:

'3. Member States must authorise the importation of apiculture by-products, other than beeswax, in the form of honeycomb intended for use in apiculture if they:

(a) come from third countries that appear on the list in Part XII of Annex XI;

(b) either:

(i) have been subjected to a temperature of -12°C or lower for at least 24 hours; or

(ii) in the case of wax, the material has been refined or rendered before importation; and

(c) are accompanied by a health certificate that conforms to the model set out in Chapter 13 of Annex X.'

(iii) In B, the following paragraphs 4 and 5 are added:

'4. Member States must authorise the importation of beeswax for technical purposes, other than beeswax in the form of honeycomb, if it:

(i) has been refined or rendered before importation; and

(ii) is accompanied by a commercial document attesting that refinement or rendering.

5. The importation of beeswax in the form of honeycomb shall be prohibited.'

(f) In Chapter XI, point 2, the second subparagraph is replaced by the following:

'however, animal by-products for use in feed for farmed fur animals or for use in raw petfood must consist of animal by-products referred to in Article 6(1)(a) and (b) only;'

(5) Annex X is amended as follows:

(a) The title is replaced by the following:

'ANNEX X

MODEL HEALTH CERTIFICATES FOR THE IMPORTATION FROM THIRD COUNTRIES AND FOR THE TRANSIT THROUGH THE EUROPEAN COMMUNITY OF CERTAIN ANIMAL BY-PRODUCTS AND PRODUCTS DERIVED THEREFROM'

(b) In the Notes to Annex X, the following Note i) is added:

'i) If health certificates are used for consignments in transit, the box No I.5 ("Consignee") of the relevant health certificate shall be completed with the name and address of the border inspection post through which the consignment is intended to leave the European Community.'

(c) Chapter 1 is replaced by the following:

'CHAPTER 1

Health certificate

For processed animal protein not intended for human consumption, including mixtures and products other than petfood containing such protein, for dispatch to [or for transit through] [2] the European Community

COUNTRY **Veterinary certificate to EU**

Part I : Details of dispatched consignment

I.1. Consignor
Name
Address
Tel.N°

I.2. Certificate reference number | I.2.a

I.3. Central Competent Authority

I.4. Local Competent Authority

I.5. Consignee
Name
Address
Postal code
Tel.N°

I.6 Importer
Name
Address
Postal code
Tel.N°

I.7.Country of origin | ISO code | I.8. Region of origin | Code

I.9. Country of destination | ISO code | I.10. Region of destination | Code

I.11. Place of origin
Name | Approval number
Address

I.12. Place of destination
Establishment ☐
Custom warehouse ☐ | Ship supplier ☐
Name | Approval number
Address
Postal code

I.13. Place of loading

I.14. Date of departure

I.15. Means of transport
Aeroplane ☐ | Ship ☐ | Railway wagon ☐
Road vehicle ☐ | Other ☐
Identification:
Documentary references:

I.16. Entry BIP in EU

I.17.

I.18. Description of commodity

I.19. Commodity code (HS code)

I.20.Quantity

I.21 Temperature of product
Ambient ☐ | Chilled ☐ | Frozen ☐

I.22. Number of packages

I.23. Identification of container/Seal number

I.24.Type of packaging

I.25. Commodities certified for:
Animal feedingstuff ☐ | Technical use ☐ | Other ☐

I.26. For transit to 3rd Country vis-à-vis EU ☐
3rd country | ISO code

I.27. For import or admission into EU ☐

I.28. Identification of the commodities

Species (Scientific name)	Nature of commodity	Approval number of establishments: Manufacturing plant	Net weight	Batch number

COUNTRY

Processed animal protein not intended for human consumption including mixtures and products other than petfood containing such protein

Part II: Certification

II.a. Certificate reference number	II.b.

II. Health attestation

I, the **undersigned** official veterinarian, declare that I have read and understood Regulation (EC) No 1774/2002[1] and in particular Article 6 and Annex VII Chapter II thereof and certify that :

II.1. the processed animal protein or product described above contains exclusively processed animal protein not intended for human consumption that :

a) has been prepared and stored in a plant approved, validated and supervised by the competent authority in accordance with Article 17 and where appropriate Article 11 of Regulation (EC) No 1774/2002, and

b) has been prepared exclusively with the following animal by-products :

[2]either [- parts of slaughtered animals, which were fit for human consumption in accordance with Community legislation, but are not intended for human consumption for commercial reasons,]

[2]and/*or* [- parts of slaughtered animals, which were rejected as unfit for human consumption but are not affected by any signs of diseases communicable to humans or animals and derive from carcases that were fit for human consumption in accordance with Community legislation,]

[2]and/or [- hides and skins, hooves and horns, pig bristles and feathers originating from

animals that were slaughtered in a slaughterhouse, underwent ante-mortem inspection and were fit, as a result of such inspection, for slaughter in accordance with Community legislation,]

(2)and/*or* [- blood obtained from animals other than ruminants that were slaughtered in a slaughterhouse, underwent ante-mortem inspection and were fit, as a result of such inspection, for slaughter in accordance with Community legislation,]

(2)and/*or* [- animal by-products derived from the production of products intended for human consumption, including degreased bones and greaves,]

(2)and/*or* [- former foodstuffs of animal origin, or former foodstuffs containing products of animal origin, other than catering waste, which are no longer intended for human consumption for commercial reasons or due to problems of manufacturing or packaging defects or other defects which do not present any risk to humans or animals,]

(2)and/*or* [- fish or other sea animals, except sea mammals, caught in the open sea for the purposes of fishmeal production,]

(2)and/or [- fresh by-products from fish from plants manufacturing fish products for human consumption,]

(2)and/*or* [- shells, hatchery by-products and cracked egg by-products originating from animals which did not show clinical signs of any disease communicable through that product to humans or animals,]

and

c) has been subjected to the following processing standard :

(2)either [heating to a core temperature of more than 133°C for at least 20 minutes without interruption at a pressure (absolute) of at least 3 bars produced by saturated steam, with a particle size prior to processing of not more than 50 millimetres;]

(2)or [in the case of non-mammalian protein other than fishmeal, the processing method ..... as set out in Annex V, Chapter III, of Regulation (EC) 1774/2002;]

(2)or [in the case of fishmeal the processing method ..... as set out in Annex V, Chapter III, of Regulation (EC) 1774/2002;]

(2)or [in the case of porcine blood, the processing method.....as set out in Annex V, Chapter III to Regulation (EC) No 1774/2002, where in case of method 7 a heat treatment of at least 80 °C has been applied throughout its substance]

II.2. the competent authority examined a random sample immediately prior to dispatch and found it to comply with the following standards (3) :

Salmonella: Absence in 25 g: n = 5, c = 0, m = 0, M = 0

Enterobacteriaceae: n = 5, c = 2, m = 10, M = 300 in 1g;

II.3. the end product :

(2)either [was packed in new or sterilised bags,]

(2)or [was transported in bulk in containers or other means of transport that were thoroughly cleaned and disinfected with a disinfectant approved by the competent authority before use,]

which bear labels indicating "NOT FOR HUMAN CONSUMPTION"

II.4. the end product was stored in enclosed storage;

II.5. the product has undergone all precautions to avoid recontamination with pathogenic agents after treatment.

Official veterinarian	
Name (in capitals):	Qualification and title:
Date: Stamp:'	Signature:

(d) Chapter 2(A) is amended as follows:

(i) The title to the health certificate is replaced by the following:

'Health certificate

For milk and milk-based products, which have undergone a single heat treatment and are not intended for human consumption for dispatch to [or for transit through] [2] the European Community';

(ii) In the box in the second column of the model health certificate, the title is replaced by the following:

'VETERINARY CERTIFICATE

For milk and milk-based products, which have undergone a single heat treatment and are not intended for human consumption for dispatch to [or for transit through] [2] the European Community';

(e) Chapter 2(B) is amended as follows:

(i) The title to the health certificate is replaced by the following:

'Health certificate

For heat-treated milk-based products which a pH reduced to less than six not intended for human consumption and for dispatch to [or for transit through] [2] the European Community';

(ii) In the box in the second column of the model health certificate, the title is replaced by the following:

'VETERINARY CERTIFICATE

For heat-treated milk-based products which a pH reduced to less than six not intended for human consumption and for dispatch to [or for transit through] [2] the European Community';

(f) Chapter 2(C) is amended as follows:

(i) The title to the health certificate is replaced by the following:

'Health certificate

For milk and milk-based products which have undergone a sterilisation or a double heat treatment and are not intended for human consumption, for dispatch to [or for transit through] [2] the European Community';

(ii) In the box in the second column of the model health certificate, the title is replaced by the following:

'VETERINARY CERTIFICATE

For milk and milk-based products which have undergone a sterilisation or a double heat treatment and are not intended for human consumption, for dispatch to [or for transit through] [2] the European Community';

(g) Chapters 3(A) to (E) are replaced by the following:

'CHAPTER 3 (A)

Health certificate

For canned petfood intended for dispatch to [or for transit through] [2] the European Community

COUNTRY

Veterinary certificate to EU

Part I : Details of dispatched consignment

I.1. Consignor
☐ Name
Address
Tel.N°

I.2. Certificate reference number | I.2.a

I.3. Central Competent Authority

I.4. Local Competent Authority

I.5. Consignee
Name
Address
Postal code
Tel.N°

I.6 Importer
Name
Address
Postal code
Tel.N°

I.7.Country of origin | ISO code | I.8. Region of origin | Code

I.9. Country of destination | ISO code | I.10. Region of destination | Code

I.11. Place of origin
Name | Approval number
Address

I.12. Place of destination
Establishment ☐
Custom warehouse ☐
Ship supplier ☐
Name | Approval number
Address
Postal code

I.13. Place of loading

I.14. Date of departure

I.15. Means of transport
Aeroplane ☐ Ship ☐ Railway wagon ☐
Road vehicle ☐ Other ☐
Identification:
Documentary references:

I.16. Entry BIP in EU

I.17.

I.18. Description of commodity

I.19. Commodity code (HS code)
23.09.10

I.20.Quantity

I.21 Temperature of product
Ambient ☐ Chilled ☐ Frozen ☐

I.22. Number of packages

I.23. Identification of container/Seal number

I.24.Type of packaging

I.25. Commodities certified for:
Animal feedingstuff ☐ Technical use ☐ Other ☐

I.26. For transit to 3rd Country vis-à-vis EU ☐
3rd country | ISO code

I.27. For import or admission into EU ☐

I.28. Identification of the commodities
Species (Scientific name) | Approval number of establishments Manufacturing plant | Net weight | Batch number

Part II: Certification

II.a. Certificate reference number	II.b.

II. Health attestation

I, the undersigned official veterinarian, declare that I have read and understood Regulation (EC) No 1774/2002[1] and in particular Article 6 and Annex VIII Chapter II thereof and certify that the petfood described above :

II.1. has been prepared and stored in a plant approved and supervised by the competent authority in accordance with Article 18 and where appropriate Article 11 of Regulation (EC) No 1774/2002;

II.2. has been prepared exclusively with the following animal by-products :

[2] *either* [- parts of slaughtered animals, which were fit for human consumption in accordance with Community legislation, but are not intended for human consumption for commercial reasons;]

[2] *and/or* [- parts of slaughtered animals, which were rejected as unfit for human consumption but are not affected by any signs of diseases communicable to humans or animals and derive from carcases that were fit for human consumption in accordance with Community legislation;]

[2] *and/or* [- hides and skins, hooves and horns, pig bristles and feathers originating from animals that were slaughtered in a slaughterhouse, underwent ante-mortem inspection and were fit, as a result of such inspection, for slaughter in accordance with Community legislation;]

[2] *and/or* [- blood obtained from animals other than ruminants that were slaughtered in a slaughterhouse, underwent ante-mortem inspection and were fit, as a result of such inspection, for slaughter in accordance with Community legislation;]

[2] *and/or* [- animal by-products derived from the production of products intended for human consumption, including degreased bones and greaves;]

[2] *and/or* [- former foodstuffs of animal origin, or former foodstuffs containing products of animal origin,

other than catering waste, which are no longer intended for human consumption for commercial reasons or due to problems of manufacturing or packaging defects or other defects which do not present any risk to humans or animals;]

(2) *and/or* [- raw milk originating from animals that do not show clinical signs of any disease communicable through that product to humans or animals;]

(2) *and/or* [- fish or other sea animals, except sea mammals, caught in the open sea for the purposes of fishmeal production;]

(2) *and/or* [- fresh by-products from fish from plants manufacturing fish products for human consumption;]

(2) *and/or* [- shells, hatchery by-products and cracked egg by-products originating from animals which did not show clinical signs of any disease communicable through that product to humans or animals;]

(2) *and/or* [material from animals which have been treated with certain substances which are prohibited pursuant to Directive 96/22/EC, the import of the material being permitted in accordance with Article 28 of Regulation (EC) No 1774/2002].

II.3. has been subjected to heat treatment to a minimum Fc value of 3 in hermetically sealed containers;

II.4. was analysed by a random sampling of at least five containers from each processed batch by laboratory diagnostic methods to ensure adequate heat treatment of the whole consignment as foreseen under point II.1;

II.5. has undergone all precautions to avoid contamination with pathogenic agents after treatment.

Notes

Part I:

- Box reference I.6: Importer: this box is to be filled in only if it is a certificate for transit commodity.
- Box reference I.12: Place of destination: this box is to be filled in only if it is a certificate for transit commodity.
- Box reference I.15: Registration number (railway wagons or container and lorries), flight number (aircraft) or name (ship) and information is to be provided in the event of unloading and reloading.
- Box reference I.23: for bulk containers, the container number and the seal number (if applicable) should be given.
- Box reference I.26 and I.27: fill in according to whether it is a transit or an import certificate.

Part II:

(1) OJ L 273, 10.10.2002, p. 1.

(2) Delete as appropriate.

- The signature and the stamp must be in a different colour to that of the printing.
- Note for the importer: This certificate is only for veterinary purposes and has to accompany the consignment until it reaches the border inspection post.

Official veterinarian

Name (in capitals): Qualification and title:

Date: Signature:

Stamp:

CHAPTER 3 (B)

Health certificate

For processed petfood other than canned petfood, intended for dispatch to [or for transit through] [2] the European Community

COUNTRY **Veterinary certificate to EU**

Part I : Details of dispatched consignment

I.1. Consignor
Name
Address
Tel.N°

I.2. Certificate reference number | I.2.a

I.3. Central Competent Authority

I.4. Local Competent Authority

I.5. Consignee
Name
Address
Postal code
Tel.N°

I.6 Importer
Name
Address
Postal code
Tel.N°

I.7.Country of origin | ISO code | I.8. Region of origin | Code

I.9. Country of destination | ISO code | I.10. Region of destination | Code

I.11. Place of origin
Name | Approval number
Address

I.12. Place of destination
Establishment ☐ Custom warehouse ☐ Ship supplier ☐
Name | Approval number
Address
Postal code

I.13. Place of loading

I.14. Date of departure

I.15. Means of transport
Aeroplane ☐ Ship ☐ Railway wagon ☐
Road vehicle ☐ Other ☐
Identification:
Documentary references:

I.16. Entry BIP in EU

I.17.

I.18. Description of commodity

I.19. Commodity code (HS code) **23.09.10**

I.20.Quantity

I.21 Temperature of product
Ambient ☐ Chilled ☐ Frozen ☐

I.22. Number of packages

I.23. Identification of container/Seal number

I.24.Type of packaging

I.25. Commodities certified for:
Animal feedingstuff ☐ Technical use ☐ Other ☐

I.26. For transit to 3rd Country vis-à-vis EU ☐
3rd country | ISO code

I.27. For import or admission into EU ☐

I.28. Identification of the commodities

Species (Scientific name)	Approval number of establishments: Manufacturing plant	Net weight	Batch number

Part II: Certification

II.a.	Certificate reference number	II.b.

II. Health attestation

I, the undersigned official veterinarian, declare that I have read and understood Regulation (EC) No 1774/2002[1] and in particular Article 6 and Annex VIII Chapter II thereof and certify that the petfood described above :

II.1. has been prepared and stored in a plant approved and supervised by the competent authority in accordance with Article 18 and where appropriate Article 11 of Regulation (EC) No 1774/2002;

II.2. has been prepared exclusively with the following animal by-products :

[2] either [- parts of slaughtered animals, which were fit for human consumption in accordance with Community legislation, but are not intended for human consumption for commercial reasons;]

[2] *and/or* [- parts of slaughtered animals, which were rejected as unfit for human consumption but are not affected by any signs of diseases communicable to humans or animals and derive from carcases that were fit for human consumption in accordance with Community legislation;]

[2] *and/or* [- hides and skins, hooves and horns, pig bristles and feathers originating from animals that were slaughtered in a slaughterhouse, underwent ante-mortem inspection and were fit, as a result of such inspection, for slaughter in accordance with Community legislation;]

[2] and/*or* [- blood obtained from animals other than ruminants that were slaughtered in a slaughterhouse, underwent ante-mortem inspection and were fit, as a result of

such inspection, for slaughter in accordance with Community legislation;]

(2) and/*or* [- animal by-products derived from the production of products intended for human consumption, including degreased bones and greaves;]

(2) *and/or* [- former foodstuffs of animal origin, or former foodstuffs containing products of animal origin, other than catering waste, which are no longer intended for human consumption for commercial reasons or due to problems of manufacturing or packaging defects or other defects which do not present any risk to humans or animals;]

(2) and/*or* [- raw milk originating from animals that do not show clinical signs of any disease communicable through that product to humans or animals;]

(2) *and/or* [- fish or other sea animals, except sea mammals, caught in the open sea for the purposes of fishmeal production;]

(2) *and/*or [- fresh by-products from fish from plants manufacturing fish products for human consumption;]

(2) *and/or* [shells, hatchery by-products and cracked egg by-products originating from animals which did not show clinical signs of any disease communicable through that product to humans or animals;]

(2) *and/or* [material from animals which have been treated with certain substances which are prohibited pursuant to Directive 96/22/EC, the import of the material being permitted in accordance with Article 28 of Regulation (EC) No 1774/2002].

II.3.

(2) either [was subjected to a heat treatment of at least 90 °C throughout its substance;]

(2) ***or*** [was produced exclusively using products of animal origin which had been

(a) in the case of milk and milk based products,

(i) produced and derived in the region .....of .....(country) listed in the Annex to Commission Decision 2004/438/EC(6) and which has been free from foot-and-mpouth disease and rinderpest for 12 months immediately prior to export and has not practised vaccination against rinderpest during that period

(ii) produced from raw milk derived from animals which at the time of milking did not show clinical signs of any disease transmissible through milk to humans or animals, and which had been kept for at least 30 days prior to production on holdings that were not subject to official restrictions due to foot-and-mouth disease or rinderpest;

(iii) are

(3)*either* [milk or milk products, excluding whey, that have undergone one of the treatments or combinations thereof described in point iv]

(3)*or* [comprised entirely of whey with a pH below 6, which was collected not earlier than 16 hours after clotting from milk subjected to one of the treatments described in point iv]

(iv) have been subject to one of the following treatments:

(3)*either* [High Temperature Short Time pasteurisation at 72°C for at least 15 seconds, or an equivalent pasteurisation achieving a negative reaction to a phosphatase test, followed by:

(3)*either* a second High Temperature Short Time pasteurisation at 72°C for at least 15 seconds or an equivalent pasteurisation which itself achieves a negative reaction to a phosphatase test]

(3)*or* additional heating to 72°C or higher, combined with a drying process]

(3)*or* a process by which the pH is reduced and kept for at least one hour at a level below 6]

(3)*or* [sterilisation at a level of at least $F_0 3$]

(3)*or* [Ultra High Temperature treatment at 132°C for at least one second followed by

(3)*either* additional heating to 72°C or higher, combined with a drying process]

Official veterinarian

Name (in capitals): Qualification and title:

Date:
Stamp: Signature:

CHAPTER 3 (C)

Health certificate

For dogchews intended for dispatch to [or for transit through] [2] the European Community

COUNTRY — **Veterinary certificate to EU**

Part I : Details of dispatched consignment

I.1. Consignor
☐ Name
Address
Tel.N°

I.2. Certificate reference number | I.2.a

I.3. Central Competent Authority

I.4. Local Competent Authority

I.5. Consignee
Name
Address
Postal code
Tel.N°

I.6 Importer
Name
Address
Postal code
Tel.N°

I.7.Country of origin | ISO code | I.8. Region of origin | Code

I.9. Country of destination | ISO code | I.10. Region of destination | Code

I.11. Place of origin
Name | Approval number
Address

I.12. Place of destination
Establishment ☐
Custom warehouse ☐ | Ship supplier ☐
Name | Approval number
Address
Postal code

I.13. Place of loading

I.14. Date of departure

I.15. Means of transport
Aeroplane ☐ | Ship ☐ | Railway wagon ☐
Road vehicle ☐ | Other ☐
Identification:
Documentary references:

I.16. Entry BIP in EU

I.17.

I.18. Description of commodity

I.19. Commodity code (HS code) **42.05.00**

I.20.Quantity

I.21 Temperature of product
Ambient ☐ | Chilled ☐ | Frozen ☐

I.22. Number of packages

I.23. Identification of container/Seal number

I.24.Type of packaging

I.25. Commodities certified for:
Animal feedingstuff ☐ | Technical use ☐ | Other ☐

I.26. For transit to 3rd Country vis-à-vis EU ☐
3rd country | ISO code

I.27. For import or admission into EU ☐

I.28. Identification of the commodities
Species (Scientific name) | Approval number of establishments Manufacturing plant | Net weight | Batch number

COUNTRY

Dogchews

Part II: Certification

II.a.	Certificate reference number	II.b.

II. Health attestation

I, the undersigned official veterinarian, declare that I have read and understood Regulation (EC) No 1774/2002[1] and in particular Article 6 and Annex VIII Chapter II thereof and certify that the dogchews described above :

II.1. have been prepared and stored in a plant approved and supervised by the competent authority in accordance with Article 18 and where appropriate Article 11 of Regulation (EC) No 1774/2002;

II.2. have been prepared exclusively with the following animal by-products :

[2] *either* [- parts of slaughtered animals, which were fit for human consumption in accordance with Community legislation, but are not intended for human consumption for commercial reasons;]

[2] *and/or* [- parts of slaughtered animals, which were rejected as unfit for human consumption but are not affected by any signs of diseases communicable to humans or animals and derive from carcasses that were fit for human consumption in accordance with Community legislation;]

[2] *and/or* [- hides and skins originating from animals that were slaughtered in a slaughterhouse, underwent ante-mortem inspection and were fit, as a result of such inspection, for slaughter in accordance with Community legislation;]

[2] *and/or* [- animal by-products derived from the production of products intended for human consumption, including degreased bones and greaves;]

[2] *and/or* [- fresh by-products from fish from plants manufacturing fish products for

human consumption;]

(2) *and/or* [- material from animals which have been treated with certain substances which are prohibited pursuant to Directive 96/22/EC, the import of the material being permitted in accordance with Article 28 of Regulation (EC) No 1774/2002].

II.3. have been subjected

(2) *either* [in the case of dogchews made from hides and skins of ungulates or from fish, to a treatment sufficient to destroy pathogenic organisms (including salmonella); and the dogchews are dry];

(2) *or* [in the case of dogchews made from animal by-products other than hides and skins of ungulates or from fish, to a heat treatment of at least 90°C throughout their substance;]

II.4. were examined by random sampling of at least five samples from each processed batch taken during or after storage at the processing plant and complies with the following standards (3):

Salmonella: absence in 25g: n = 5, c = 0, m = 0, M = 0,

Enterobacteriaceae: n = 5, c = 2, m = 10, M = 300 in 1 gram;

II.5. have undergone all precautions to avoid contamination with pathogenic agents after treatment;

II.6. were packed in new packaging

Notes

Part I:

- Box reference I.6: Importer: this box is to be filled in only if it is a certificate for transit commodity.
- Box reference I.12: Place of destination: this box is to be filled in only if it is a certificate for transit commodity.
- Box reference I.15: Registration number (railway wagons or container and lorries), flight number (aircraft) or name (ship) and information is to be provided in the event of unloading and reloading.
- Box reference I.23: for bulk containers, the container number and the seal number (if applicable) should be given.
- Box reference I.26 and I.27: fill in according to whether it is a transit or an import certificate.

Part II:

(1) OJ L 273, 10.10.2002, p.1.

(2) Delete as appropriate.

(3) Where:

n = number of samples to be tested;

m = threshold value for the number of bacteria; the result is considered satisfactory if the number of bacteria in all samples does not exceed m;

M = maximum value for the number of bacteria; the result is considered unsatisfactory if the number of bacteria in one or more samples is M or more; and

c = number of samples the bacterial count of which may be between m and M, the sample still being considered acceptable if the bacterial count of the other samples is m or less

- The signature and the stamp must be in a different colour to that of the printing.
- Note for the importer: This certificate is only for veterinary purposes and has to accompany the consignment until it reaches the border inspection post.

Official veterinarian

Name (in capitals):

Qualification and title:

Date:

Stamp:

Signature:

CHAPTER 3 (D)

Health certificate

For raw petfood for direct sale or animal by-products to be fed to farmed fur animals, intended for dispatch to [or for transit through] [2] the European Community

COUNTRY

Veterinary certificate to EU

Part I : Details of dispatched consignment

I.1. Consignor
Name
Address
Tel.N°

I.2. Certificate reference number | I.2.a

I.3. Central Competent Authority

I.4. Local Competent Authority

I.5. Consignee
Name
Address
Postal code
Tel.N°

I.6 Importer
Name
Address
Postal code
Tel.N°

I.7.Country of origin | ISO code | I.8. Region of origin | Code

I.9. Country of destination | ISO code | I.10. Region of destination | Code

I.11. Place of origin
Name | Approval number
Address

I.12. Place of destination
Establishment ☐ Custom warehouse ☐ Ship supplier ☐
Name | Approval number
Address
Postal code

I.13. Place of loading

I.14. Date of departure

I.15. Means of transport
Aeroplane ☐ Ship ☐ Railway wagon ☐
Road vehicle ☐ Other ☐
Identification:
Documentary references:

I.16. Entry BIP in EU

I.17.

I.18. Description of commodity

I.19. Commodity code (HS code)

I.20.Quantity

I.21 Temperature of product
Ambient ☐ Chilled ☐ Frozen ☐

I.22. Number of packages

I.23. Identification of container/Seal number

I.24.Type of packaging

I.25. Commodities certified for:
Animal feedingstuff ☐ Technical use ☐ Other ☐

I.26. For transit to 3rd Country vis-à-vis EU ☐
3rd country | ISO code

I.27. For import or admission into EU ☐

I.28. Identification of the commodities

Species (Scientific name)	Nature of commodity	Approval number of establishments: Manufacturing plant	Net weight	Batch number

COUNTRY

Raw petfood for direct sale or animal by products to be fed to farmed fur animals

Part II: Certification

II.a. Certificate reference number	II.b.

II. Health attestation

I, the undersigned official veterinarian, declare that I have read and understood Regulation (EC) No 1774/2002[1] and in particular Article 6 and Annex VIII Chapter II thereof and certify that the raw petfood or animal by-product described above :

II.1. consist of animal by-products that satisfy the health requirements below;

II.2. consist of animal by-products :

a) derived from meat which satisfies the relevant animal and public health requirements laid down in :

- Council Decision 79/542/EEC[2] and provided the animals from which the meat is derived come from a territory or part of a territory .....(ISO code) as listed in that Decision which has been free of foot and mouth disease, rinderpest, classical swine fever, African swine fever and swine vesicular disease for the last 12 months and where no vaccination has taken place during that time (only as relevant for the susceptible species),
- and/or Commission Decision 2006/XXX/EC[3], and provided the animals from which the meat is derived come from a territory or part of a territory .....(ISO code) as listed in that Decision which has been free from Newcastle disease and Avian Influenza for the last 12 months,
- and/or Commission Decision 2000/585/EC[4] , and provided the animals from which the meat is derived come from a territory or part of a territory .....(ISO code) as listed which has been free from foot and mouth disease, rinderpest, classical swine

fever, African swine fever, swine vesicular disease, Newcastle disease and Avian Influenza for the last 12 months and where no vaccination has taken place during that time (only as relevant for the susceptible species),

b) derived from animals that, at the slaughterhouse, have passed the ante-mortem health inspection during the 24 hours before the slaughter and have shown no evidence of the diseases referred in the Decisions above for which the animals are susceptible, and

c) derived from animals that have been treated in the slaughterhouse before and at the time of slaughter or killing in accordance with the relevant provisions of Council Directive 93/119/EC[5] on animal welfare;

II.3. consist only of the following animal by-products:

a) parts of slaughtered animals, which were fit for human consumption in accordance with Community legislation, but are not intended for human consumption for commercial reasons, and

b) parts of slaughtered animals, which are rejected as unfit for human consumption but are not affected by any signs of diseases communicable to humans or animals and derive from carcasses that are fit for human consumption in accordance with Community legislation;

and

II.4. have been obtained and prepared without contact with other material not complying with the conditions required in the Decisions above, and it has been handled so as to avoid contamination with pathogenic agents;

II.5. have been packed in final packaging which bear labels indicating "RAW PETFOOD - NOT FOR HUMAN CONSUMPTION" or "ANIMAL BY-PRODUCTS FOR FEED FOR FUR ANIMALS - NOT FOR HUMAN CONSUMPTION" and then in leak-proof and officially sealed boxes/containers or in new packaging preventing any leakage and officially sealed boxes/containers which bear labels indicating "RAW PETFOOD - NOT FOR HUMAN CONSUMPTION" or "ANIMAL BY-PRODUCTS FOR FEED FOR FUR ANIMALS - NOT FOR HUMAN CONSUMPTION", the name and the address of the establishment of destination;

II.6. in the case of raw petfood:

a) have been prepared and stored in a plant approved and supervised by the competent authority in accordance with Article 18 and where appropriate Article 11 of Regulation (EC) No 1774/2002 and

b) were examined by random sampling of at least five samples from each batch taken during storage (before dispatch) and complies with the following standards ([6]):

Salmonella: absence in 25 g: n=5, c=0, m=0, M=0

Enterobacteriacae: n=5, c=2, m=10, M=300 in 1 gram.**Notes**

Part I:

- Box reference I.6: Importer: this box is to be filled in only if it is a certificate for transit commodity.
- Box reference I.12: Place of destination: this box is to be filled in only if it is a certificate for transit commodity.
- Box reference I.15: Registration number (railway wagons or container and lorries), flight number (aircraft) or name (ship) and information is to be provided in the event of unloading and reloading.
- Box reference I.19: use the appropriate HS code: 05.11.91; 05.11.99 or 23.09.90.
- Box reference I.23: for bulk containers, the container number and the seal number (if applicable) should be given.
- Box reference I.26 and I.27: fill in according to whether it is a transit or an import certificate.
- Box reference I.28: Nature of commodity: select raw petfood or animal by-product.

Part II:

(1) OJ L 273, 10.10.2002, p.1.

(2) Council Decision 79/542/EEC of 21 December 1976 drawing up a list of third countries or parts of third countries, and laying down animal and public health and veterinary certification conditions, for importation into the Community of certain live animals and their fresh meat.

(3) Commission Decision 2006/XXX/EC (NOTE: Decision 2006*XXX/EC is reference to the now draft combined poultry import decision, voted in favour in December SCoFCAH)*

Official veterinarian

Name (in capitals): Qualification and title:

Date:
Stamp: Signature:

CHAPTER 3 (E)

Health certificate

For flavouring innards for use in the manufacture of petfood, intended for dispatch to [or for transit through] [2] the European Community

COUNTRY **Veterinary certificate to EU**

Part I : Details of dispatched consignment

I.1. Consignor
Name
Address
Tel.N°

I.2. Certificate reference number | I.2.a

I.3. Central Competent Authority

I.4. Local Competent Authority

I.5. Consignee
Name
Address
Postal code
Tel.N°

I.6 Importer
Name
Address
Postal code
Tel.N°

I.7.Country of origin | ISO code | I.8. Region of origin | Code

I.9. Country of destination | ISO code | I.10. Region of destination | Code

I.11. Place of origin
Name | Approval number
Address

I.12. Place of destination
Establishment ☐
Custom warehouse ☐ | Ship supplier ☐
Name | Approval number
Address
Postal code

I.13. Place of loading

I.14. Date of departure

I.15. Means of transport
Aeroplane ☐ Ship ☐ Railway wagon ☐
Road vehicle ☐ Other ☐
Identification:
Documentary references:

I.16. Entry BIP in EU

I.17.

I.18. Description of commodity

I.19. Commodity code (HS code)

I.20.Quantity

I.21 Temperature of product
Ambient ☐ Chilled ☐ Frozen ☐

I.22. Number of packages

I.23. Identification of container/Seal number

I.24.Type of packaging

I.25. Commodities certified for:
Animal feedingstuff ☐ Technical use ☐ Other ☐

I.26. For transit to 3rd Country vis-à-vis EU ☐
3rd country | ISO code

I.27. For import or admission into EU ☐

I.28. Identification of the commodities

Species (Scientific name)	Nature of commodity	Approval number of establishments Manufacturing plant	Net weight	Batch number

Part II: Certification

II.a.	Certificate reference number	II.b.

II. Health attestation

I, the undersigned official veterinarian, declare that I have read and understood Regulation (EC) No 1774/2002[1] and in particular Article 6 and Annex VIII Chapter XIV thereof and certify that the flavouring innards products described above :

II.1. Consist of animal by-products that satisfy the animal health requirement below;

II.2. Have been prepared and stored in a plant approved, validated and supervised by the competent authority in accordance with Article 18 and where appropriate Article 11 of Regulation (EC) No 1774/2002;

II.3. have been prepared including the following animal by-products which are exclusively :

[2] *either* [- parts of slaughtered animals, which were fit for human consumption in accordance with Community legislation, but are not intended for human consumption for commercial reasons;]

[2] *and/or* [- parts of slaughtered animals, which were rejected as unfit for human consumption but are not affected by any signs of diseases communicable to humans or animals and derive from carcases that were fit for human consumption in accordance with Community legislation;]

[2] *and/or* [- hides and skins, hooves and horns, pig bristles and feathers originating from animals that were slaughtered in a slaughterhouse, underwent ante-mortem inspection and were fit, as a result of such inspection, for slaughter in accordance with Community legislation;]

[2] *and/or* [- blood obtained from animals other than ruminants that were slaughtered in a slaughterhouse, underwent ante-mortem inspection and were fit, as a result of such inspection, for slaughter in accordance with Community legislation;]

[2] *and/or* [- *animal* by-products derived from the production of products intended for human consumption, including degreased bones and greaves;]

[2] *and/or* [former foodstuffs of animal origin, or former foodstuffs containing products of animal origin, other than catering waste, which are no longer intended for human consumption for commercial reasons or due to problems of manufacturing or packaging defects or other defects which do not present any risk to humans or animals;]

[2] *and/or* [- raw milk originating from animals that do not show clinical signs of any disease communicable through that product to humans or animals,]

[2] *and/or* [- *fish* or other sea animals, except sea mammals, caught in the open sea for the purposes of fishmeal production;]

[2] *and/or* [- fresh by-products from fish from plants manufacturing fish products for human consumption;]

[2] *and/or* [- *shells*, hatchery by-products and cracked egg by-products originating from animals which did not show clinical signs of any disease communicable through that product to humans or animals;]

[2] *and/or* [- *material* from animals which have been treated with certain substances which are prohibited pursuant to Directive 96/22/EC, the import of the material being permitted in accordance with Article 28 of Regulation (EC) No 1774/2002]

II.4. have been subjected to processing in accordance with Annex VIII, Chapter XIV of Regulation 1774/2002/EC , in order to kill pathogenic agents;

II.5. have been examined by the competent authority taking a random sample immediately prior to dispatch and found it to comply with the following standards [3]:

Salmonella: absence in 25g: n = 5, c = 0, m = 0, M = 0,

Enterobacteriaceae: n = 5, c = 2, m = 10, M = 300 in 1 gram;

II.6. the end product was :

[2] *either* [*packed* in new or sterilised bags,]

[2] *or* [*transported* in bulk in containers or other means of transport that were thoroughly cleaned and disinfected with a disinfectant approved by the competent authority before use,]

and *which* bear labels indicating "NOT FOR HUMAN CONSUMPTION";

II.7. the end product was stored in enclosed storage;

II.8. the product has undergone all precautions to avoid contamination with pathogenic agents after treatment.

Part I:

- Box reference I.6: Importer: this box is to be filled in only if it is a certificate for transit commodity.
- Box reference I.12: Place of destination: this box is to be filled in only if it is a certificate for transit commodity.
- Box reference I.15: Registration number (railway wagons or container and lorries), flight number (aircraft) or name (ship) and information is to be provided in the event of unloading and reloading.
- Box reference I.19: use the appropriate HS code: 05.04 or 05.11.91.
- Box reference I.23: for bulk containers, the container number and the seal number (if applicable) should be given.
- Box reference I.26 and I.27: fill in according to whether it is a transit or an import certificate.
- Box reference I.28: define the innard product by choosing among the following list.(?)

Part II:

(1) OJ L 273, 10.10.2002, p.1.

(2)Delete as appropriate.

(3) Where:

n = number of samples to be tested;

m = threshold value for the number of bacteria; the result is considered satisfactory if the number of bacteria in all samples does not exceed m;

M = maximum value for the number of bacteria; the result is considered unsatisfactory if the number of bacteria in one or more samples is M or more; and

c = number of samples the bacterial count of which may be between m and M, the sample still being considered acceptable if the bacterial count of the other samples is m or less

- The signature and the stamp must be in a different colour to that of the printing.
- Note for the importer: This certificate is only for veterinary purposes and has to accompany the consignment until it reaches the border inspection post.

Official veterinarian

Name (in capitals): Qualification and title:

Date: Signature:

Stamp:

(h) Chapter 4(A) is amended as follows:

(i) The title to the health certificate is replaced by the following:

'Health certificate

For the import of serum from equidae to be used for technical purposes including pharmaceuticals, in vitro diagnosis and laboratory reagents, intended for dispatch to [or for transit through] [2] the European Community';

(ii) In the box in the second column of the model health certificate, the title is replaced by the following:

'VETERINARY CERTIFICATE

For the import of serum from equidae to be used for technical purposes including pharmaceuticals, in vitro diagnosis and laboratory reagents, intended for dispatch to [or for transit through] [2] the European Community';

(i) Chapter 4(B) is amended as follows:

(i) The title to the health certificate is replaced by the following:

'Health certificate

For blood products not intended for human consumption that could be used as feed material, intended for dispatch to [or for transit through] [2] the European Community';

(ii) In the box in the second column of the model health certificate, the title is replaced by the following:

'VETERINARY CERTIFICATE

For blood products not intended for human consumption that could be used as feed material, intended for dispatch to [or for transit through] [2] the European Community';

(j) Chapter 4(C) is amended as follows:

(i) The title to the health certificate is replaced by the following:

'Health certificate

For blood products to be used for technical purposes, including pharmaceuticals, in vitro diagnosis and laboratory reagents, but excluding serum of equidae, intended for dispatch to [or for transit through] [2] the European Community';

(ii) In the box in the second column of the model health certificate, the title is replaced by the following:

'VETERINARY CERTIFICATE

For blood products to be used for technical purposes, including pharmaceuticals, in vitro diagnosis and laboratory reagents, but excluding serum of equidae, intended for dispatch to [or for transit through] [2] the European Community';

(k) Chapters 5(A), 5(B) and 5(C) are replaced by the following:

'CHAPTER 5 (A)

Health certificate

For fresh or chilled hides and skins of ungulates, intended *for dispatch to [or for transit through] [2] the European Community*

COUNTRY

Veterinary certificate to EU

Part I : Details of dispatched consignment

I.1. Consignor
☐ Name
Address
Tel.N°

I.2. Certificate reference number | I.2.a

I.3. Central Competent Authority

I.4. Local Competent Authority

I.5. Consignee
Name
Address
Postal code
Tel.N°

I.6 Importer
Name
Address
Postal code
Tel.N°

I.7.Country of origin | ISO code | I.8. Region of origin | Code

I.9. Country of destination | ISO code | I.10. Region of destination | Code

I.11. Place of origin
Name | Approval number
Address

I.12. Place of destination
Establishment ☐ Custom warehouse ☐ Ship supplier ☐
Name | Approval number
Address
Postal code

I.13. Place of loading

I.14. Date of departure

I.15. Means of transport
Aeroplane ☐ Ship ☐ Railway wagon ☐
Road vehicle ☐ Other ☐
Identification:
Documentary references:

I.16. Entry BIP in EU

I.17. No.(s) of CITES

I.18. Description of commodity

I.19. Commodity code (HS code)

I.20.Quantity

I.21 Temperature of product
Ambient ☐ Chilled ☐ Frozen ☐

I.22. Number of packages

I.23. Identification of container/Seal number

I.24.Type of packaging

I.25. Commodities certified for:
Animal feedingstuff ☐ Technical use ☐ Other ☐

I.26. For transit to 3rd Country vis-à-vis EU ☐
3rd country | ISO code

I.27. For import or admission into EU ☐

I.28. Identification of the commodities
Species (Scientific name) | Approval number of establishments Manufacturing plant | Net weight

COUNTRY

Fresh or chilled hides and skins of Ungulates

Part II: Certification

II.a.	Certificate reference number	II.b.

II. Health attestation

I, the undersigned official veterinarian, declare that I have read and understood Regulation (EC) No 1774/2002[1] and in particular Article 6 and Annex VIII Chapter VI thereof and certify that the hides and skins described above :

II.1. have been obtained from animals that [2]:

a) were slaughtered and their carcasses are fit for human consumption in accordance with Community legislation or

b) were slaughtered in a slaughterhouse, after undergoing ante-mortem inspection, and were fit, as a result of such inspection, for slaughter for human consumption in accordance with Community legislation;

II.2. originate from a country or, in the case of regionalisation in accordance with Community legislation, from a part of a country from which imports of all categories of fresh meat of the corresponding species are authorised and which :

a) for at least 12 months before dispatch, has been free from the following diseases[3]:

[- classical swine fever, and African swine fever,]

[- rinderpest,]

and

b) has been free for at least 12 months before dispatch from foot-and-mouth disease and where, for 12 months before dispatch, no vaccination has been carried out against foot-

and-mouth disease [3];

II.3. have been obtained from :

[animals that have remained in the territory of the country of origin for at least three months before being slaughtered or since birth in the case of animals less that three months old;]

[in the case of hides and skins from bi-ungulates, animals that come from holdings in which there has been no outbreak of foot-and-mouth disease in the previous 30 days, and around which within a radius of 10 km there has been no case of foot-and-mouth disease for 30 days;]

[in the case of hides and skins from swine, animals that come from holdings in which there has been no outbreak of swine vesicular disease in the previous 30 days, or of classical or African swine fever in the previous 40 days, and around which within a radius of 10 km there has been no case of these diseases for 30 days;]

[animals that have shown no evidence of [foot-and-mouth disease], [rinderpest], [classical swine fever], [African swine fever] or [swine vesicular disease][5] during ante-mortem health inspection at the slaughterhouse during the 24 hours before slaughter;]

II.4. have undergone all precautions to avoid recontamination with pathogenic agents.

Notes

Part I:

- Box reference I.6: Importer: this box is to be filled in only if it is a certificate for transit commodity.
- Box reference I.12: Place of destination: this box is to be filled in only if it is a certificate for transit commodity.
- Box reference I.15: Registration number (railway wagons or container and lorries), flight number (aircraft) or name (ship) and information is to be provided in the event of unloading and reloading.
- Box reference I.19: use the appropriate HS code: 41.01; 41.02 or 41.03.
- Box reference I.23: for bulk containers, the container number and the seal number (if applicable) should be given.
- Box reference I.26 and I.27: fill in according to whether it is a transit or an import certificate.

Part II:

(1) OJ L 273, 10.10.2002, p.1.

(2)Delete as appropriate.

(3) Delete diseases not applicable to the species concerned.

- The signature and the stamp must be in a different colour to that of the printing.
- Note for the importer: This certificate is only for veterinary purposes and has to accompany the consignment until it reaches the border inspection post.

Official veterinarian

Name (in capitals): Qualification and title:

Date:
Stamp: Signature:

CHAPTER 5 (B)

Health certificate

For treated hides and skins of ungulates, intended for dispatch to [or for transit through] [2] the European Community

COUNTRY

Veterinary certificate to EU

Part I : Details of dispatched consignment

I.1. Consignor
Name
Address
Tel.N°

I.2. Certificate reference number | I.2.a

I.3. Central Competent Authority

I.4. Local Competent Authority

I.5. Consignee
Name
Address
Postal code
Tel.N°

I.6 Importer
Name
Address
Postal code
Tel.N°

I.7.Country of origin | ISO code | I.8. Region of origin | Code

I.9. Country of destination | ISO code | I.10. Region of destination | Code

I.11. Place of origin
Name | Approval number
Address

I.12. Place of destination
Establishment ☐
Custom warehouse ☐
Ship supplier ☐
Name | Approval number
Address
Postal code

I.13. Place of loading

I.14. Date of departure

I.15. Means of transport
Aeroplane ☐ Ship ☐ Railway wagon ☐
Road vehicle ☐ Other ☐
Identification:
Documentary references:

I.16. Entry BIP in EU

I.17. No.(s) of CITES

I.18. Description of commodity

I.19. Commodity code (HS code)

I.20.Quantity

I.21 Temperature of product
Ambient ☐ Chilled ☐ Frozen ☐

I.22. Number of packages

I.23. Identification of container/Seal number

I.24.Type of packaging

I.25. Commodities certified for:
Animal feedingstuff ☐ Technical use ☐ Other ☐

I.26. For transit to 3rd Country vis-à-vis EU ☐
3rd country | ISO code

I.27. For import or admission into EU ☐

I.28. Identification of the commodities
Species (Scientific name) | Approval number of establishments Manufacturin plant | g | Net weight

COUNTRY

Treated hides and skins of Ungulates

Part II: Certification				
		II.a.	Certificate reference number	II.b.

II. Health attestation

I, the undersigned official veterinari n, declare that Iahave read and understood Regulation (EC) No 1774/2002[1] and in particular Article 6 and Annex VIII Chapter VI thereof and certify that the hides and skins described above :

9.1. have been obtained from animals that [2]:

a) were slaughtered and their carcasses are fit for human consumption in accordance with Community legislation or

b) were slaughtered in a slaughterhouse, after undergoing ante-mortem inspection, and were fit, as a result of such inspection, for slaughter for human consumption in accordance with Community legislation or

c) did not show any clinical signs of any disease communicable to humans or animals, and

d) were not killed to eradicate any epizootic disease;

[2] *either* [9.2 come from animals originate from a third country or, in the case of regionalisation in accordance with Community legislation, from a part of a third country listed in part 1 of Annex II to Decision 79/542/EEC[3] from which imports of fresh meat of the corresponding species are authorised and have been:

[2] *either* [dried;]

[2] *or* [dry-*salted* or wet-salted for at least 14 days prior to dispatch;]

(2) *or* [dry-*salted* or wet-salted on the following date ..... and according to the declaration of the transporter, the hides and skins will be transported by ship and the duration of transport will be such that they will have undergone a minimum of 14 days of salting before they reach the EC border inspection post]

(2) *or* [*salted* for seven days in sea salt with the addition of 2% of sodium carbonate;]

(2) *or* [salted in sea salt with the addition of 2% of sodium carbonate on the following date ..... and according to the declaration of the transporter, the hides and skins will be transported by ship and the duration of transport will be such that they will have undergone a minimum of 7 days of salting before they reach the EC border inspection post]]

(2) *or* [9.2. come from animals originate from a third country or, in the case of regionalisation in accordance with Community legislation, from a part of a third country listed in part 1 of Annex II to Decision 79/542/EEC from which imports of fresh meat of the corresponding species are **NOT** authorised and have been::

(2) *either* [salted for seven days in sea salt with the addition of 2% of sodium carbonate;]

(2) *or* [salted in sea salt with the addition of 2% of sodium carbonate on the following date ..... and according the declaration of the transporter, the hides and skins will be transported by ship and the duration of transport will be such that they will have undergone a minimum of 7 days of salting before they reach the EC border inspection post]

(2) *or* [dried for 42 days at a temperature of at least 20°C;]

9.3. the consignment has not been in contact with other animal products or with live animals presenting a risk of spreading a serious transmissible disease.

Notes

Part I:

- Box reference I.6: Importer: this box is to be filled in only if it is a certificate for transit commodity.
- Box reference I.12: Place of destination: this box is to be filled in only if it is a certificate for transit commodity.
- Box reference I.15: Registration number (railway wagons or container and lorries), flight number (aircraft) or name (ship) and information is to be provided in the event of unloading and reloading.
- Box reference I.19: use the appropriate HS code: 41.01; 41.02 or 41.03.
- Box reference I.23: for bulk containers, the container number and the seal number (if applicable) should be given.
- Box reference I.26 and I.27: fill in according to whether it is a transit or an import certificate.

Part II:

(1) OJ L 273, 10.10.2002, p.1.

Official veterinarian

Name (in capitals):

Qualification and title:

Date:

Stamp:

Signature:

CHAPTER 5 (C)

Official declaration

For treated hides and skins of ruminants and of equidae that are intended for dispatch to [or for transit through] [2] the European Community and have been kept separate for 21 days or will undergo transport for 21 uninterrupted days before importation

COUNTRY — **Veterinary certificate to EU**

Part I : Details of dispatched consignment

I.1. Consignor
☐ Name
Address
Tel.N°

I.2. Certificate reference number | I.2.a

I.3. Central Competent Authority

I.4. Local Competent Authority

I.5. Consignee
Name
Address
Postal code
Tel.N°

I.6 Importer
Name
Address
Postal code
Tel.N°

I.7.Country of origin | ISO code | I.8. Region of origin | Code

I.9. Country of destination | ISO code | I.10. Region of destination | Code

I.11. Place of origin
Name | Approval number
Address

I.12. Place of destination
Establishment ☐
Custom warehouse ☐
Ship supplier ☐
Name | Approval number
Address
Postal code

I.13. Place of loading

I.14. Date of departure

I.15. Means of transport
Aeroplane ☐ Ship ☐ Railway wagon ☐
Road vehicle ☐ Other ☐
Identification:
Documentary references:

I.16. Entry BIP in EU

I.17. No.(s) of CITES

I.18. Description of commodity

I.19. Commodity code (HS code)

I.20.Quantity

I.21 Temperature of product
Ambient ☐ Chilled ☐ Frozen ☐

I.22. Number of packages

I.23. Identification of container/Seal number

I.24.Type of packaging

I.25. Commodities certified for:
Animal feedingstuff ☐ Technical use ☐ Other ☐

I.26. For transit to 3rd Country vis-à-vis EU ☐
3rd country | ISO code

I.27. For import or admission into EU ☐

I.28. Identification of the commodities
Species (Scientific name) | Approval number of establishments Manufacturing plant | Net weight

COUNTRY

Treated hides and skins of Ruminants and of Equidae that have been kept separate for 21 days or will undergo transport for 21 uninterrupted days before importation

Part II: Declaration

II.a.	Certificate reference number	II.b.

II. Declaration

I, the undersigned declare that the hides and skins described above :

II.1. have been obtained from animals that (2) :

a) were slaughtered and their carcasses are fit for human consumption in accordance with Community legislation or

b) were slaughtered in a slaughterhouse, after undergoing ante-mortem inspection, and were fit , as a result of such inspection, for slaughter for human consumption in accordance with Community legislation or

c) did not show any clinical signs of any disease communicable to humans or animals, and

d) were not killed to eradicate any epizootic disease;

II.2. have been:

(2) *either* [dried;]

(2) *or* [dry-*salted* or wet-salted for at least 14 days prior to dispatch;]

(2) *or* [*salted* for seven days in sea salt with the addition of 2% of sodium carbonate;]

II.3. have not been in contact with other animal products or with live animals presenting a risk or spreading a serious transmissible disease;

(2) *either* [II.4. have been kept separate immediately before dispatch for 21 days under official supervision

after the treatment described under point (9.2).]

(2) *or* [II.4. following the declaration of the transporter, the duration of the transport period is foreseen to be at least 21 days.]

Notes

Part I:

- Box reference I.6: Importer: this box is to be filled in only if it is a certificate for transit commodity.
- Box reference I.12: Place of destination: this box is to be filled in only if it is a certificate for transit commodity.
- Box reference I.15: Registration number (railway wagons or container and lorries), flight number (aircraft) or name (ship) and information is to be provided in the event of unloading and reloading.
- Box reference I.19: use the appropriate HS code: 41.01; 41.02 or 41.03.
- Box reference I.23: for bulk containers, the container number and the seal number (if applicable) should be given.
- Box reference I.26 and I.27: fill in according to whether it is a transit or an import certificate.

Part II:

(1) OJ L 273, 10.10.2002, p.1.

(2)Delete as appropriate.

- The signature and the stamp must be in a different colour to that of the printing.
- Note for the importer: This declaration is only for veterinary purposes and has to accompany the consignment until it reaches the border inspection post.

Official

Name (in capitals): Qualification and title:

Date: Signature:
Stamp:'

(l) Chapter 6(A) is amended as follows:

(i) The title to the health certificate is replaced by the following:

'Health certificate

For treated game trophies of birds and ungulates, being solely bones, horns, hooves, claws, antlers, teeth, hides or skins, for dispatch to [or for transit through] [2] the European Community';

(ii) In the box in the second column of the model health certificate, the title is replaced by the following:

'VETERINARY CERTIFICATE

For treated game trophies of birds and ungulates, being solely bones, horns, hooves, claws, antlers, teeth, hides or skins, for dispatch to [or for transit through] [2] the European Community';

(m) Chapter 6(B) is amended as follows:

(i) The title to the health certificate is replaced by the following:

'Health certificate

For game trophies of birds and ungulates consisting of entire parts not having been treated, intended for dispatch to [or for transit through] [2] the European Community';

(ii) In the box in the second column of the model health certificate, the title is replaced by the following:

'VETERINARY CERTIFICATE

For game trophies of birds and ungulates consisting of entire parts not having been treated, intended for dispatch to [or for transit through] [2] the European Community';

(n) Chapter 7(A) is amended as follows:

(i) The title to the health certificate is replaced by the following:

'Health certificate

For pig bristles from third countries or regions thereof that are free from African swine fever, intended for dispatch to [or for transit through] [2] the European Community';

(ii) In the box in the second column of the model health certificate, the title is replaced by the following:

'VETERINARY CERTIFICATE

For pig bristles from third countries or regions thereof that are free from African swine fever, intended for dispatch to [or for transit through] [2] the European Community';

(o) Chapter 7(B) is amended as follows:

(i) The title to the health certificate is replaced by the following:

'Health certificate

For pig bristles from third countries or regions thereof that are not free from African swine fever, intended for dispatch to [or for transit through] [2] the European Community';

(ii) In the box in the second column of the model health certificate, the title is replaced by the following:

'VETERINARY CERTIFICATE

For pig bristles from third countries or regions thereof that are not free from African swine fever, intended for dispatch to [or for transit through] [2] the European Community';

(p) Chapter 8(A) is amended as follows:

(i) The title to the health certificate is replaced by the following:

'Health certificate

For animal by-products(1) for the manufacture of petfood, intended for dispatch to [or for transit through] [2] the European Community';

(ii) In the box in the second column of the model health certificate, the title is replaced by the following:

'VETERINARY CERTIFICATE

For animal by-products(1) for the manufacture of petfood, intended for dispatch to [or for transit through] [2] the European Community;

(q) Chapter 8(B) is amended as follows:

(i) The title to the health certificate is replaced by the following:

'Health certificate

For animal by-products for the manufacture of technical products (including pharmaceutical products)(1), intended for dispatch to [or for transit through] [2] European Community';

(ii) In the box in the second column of the model health certificate, the title is replaced by the following:

'VETERINARY CERTIFICATE

For animal by-products for the manufacture of technical products (including pharmaceutical products) (1), intended for dispatch to [or for transit through] [2] the European Community';

(iii) Note 1 is replaced by the following:

'1. Excluding raw blood, raw milk, hides and skins of ungulates or ruminants and pig bristles (see relevant specific certificates for the import of these products) as well as wool, hair, feathers or parts of feathers.'

(r) Chapter 9 is amended as follows:

(i) The title to the health certificate is replaced by the following:

'Health certificate

For fish oil not intended for human consumption to used be as feed material or for technical purposes, intended for dispatch to [or for transit through] [2] the European Community';

(ii) In the box in the second column of the model health certificate, the title is replaced by the following:

'VETERINARY CERTIFICATE

For fish oil not intended for human consumption to be used as feed material or for technical purposes, intended for dispatch to [or for transit through] [2] the European Community';

(s) Chapter 10(A) is amended as follows:

(i) The title to the health certificate is replaced by the following:

'Health certificate

For rendered fats not intended for human consumption to be used as feed material or for technical purposes, intended for dispatch to [or for transit through] [2] the European Community';

(ii) In the box in the second column of the model health certificate, the title is replaced by the following:

'VETERINARY CERTIFICATE

For rendered fats not intended for human consumption to be used as feed material or for technical purposes, intended for dispatch to [or for transit through] [2] the European Community';

(t) Chapter 10(B) is amended as follows:

(i) The title to the health certificate is replaced by the following:

'Health certificate

For rendered fats not intended for human consumption to be used for technical purposes, intended for dispatch to [or for transit through] [2] the European Community';

(ii) In the box in the second column of the model health certificate, the title is replaced by the following:

'VETERINARY CERTIFICATE

For rendered fats not intended for human consumption to be used for technical purposes, intended for dispatch to [or for transit through] [2] the European Community';

(u) Chapter 11 is amended as follows:

(i) The title to the health certificate is replaced by the following:

'Health certificate

For gelatine and collagen not intended for human consumption to be used as feed material or for technical purposes, intended for dispatch to [or for transit through] [2] the European Community';

(ii) In the box in the second column of the model health certificate, the title is replaced by the following:

'VETERINARY CERTIFICATE

For gelatine and collagen not intended for human consumption to be used as feed material or for technical purposes, intended for dispatch to [or for transit through] [2] the European Community';

(v) Chapter 11 is amended as follows:

(i) The title to the health certificate is replaced by the following:

'Health certificate

For hydrolysed protein, dicalcium phosphate and tricalcium phospaste not intended for human consumption to be used as feed material or for technical purposes, intended for dispatch to [or for transit through] [2] the European Community';

(ii) In the box in the second column of the model health certificate, the title is replaced by the following:

'VETERINARY CERTIFICATE

For hydrolysed protein, dicalcium phosphate and tricalcium phospaste not intended for human consumption to be used as feed material or for technical purposes, intended for dispatch to [or for transit through] [2] the European Community';

(u) Chapter 13 is replaced by the following:

'CHAPTER 13

Health certificate

For apiculture by-products, intended for dispatch to [or for transit through] [2] the European Community

COUNTRY

Veterinary certificate to EU

Part I : Details of dispatched consignment

I.1. Consignor
Name
Address
Tel.N°

I.2. Certificate reference number | I.2.a

I.3. Central Competent Authority

I.4. Local Competent Authority

I.5. Consignee
Name
Address
Postal code
Tel.N°

I.6 Importer
Name
Address
Postal code
Tel.N°

I.7.Country of origin | ISO code | I.8. Region of origin | Code

I.9. Country of destination | ISO code | I.10. Region of destination | Code

I.11. Place of origin
Name | Approval number
Address

I.12. Place of destination
Establishment ☐ Custom warehouse ☐ Ship supplier ☐
Name | Approval number
Address
Postal code

I.13. Place of loading

I.14. Date of departure

I.15. Means of transport
Aeroplane ☐ Ship ☐ Railway wagon ☐
Road vehicle ☐ Other ☐
Identification:
Documentary references:

I.16. Entry BIP in EU

I.17.

I.18. Description of commodity

I.19. Commodity code (HS code)

I.20.Quantity

I.21 Temperature of product
Ambient ☐ Chilled ☐ Frozen ☐

I.22. Number of packages

I.23. Identification of container/Seal number

I.24.Type of packaging

I.25. Commodities certified for:
Animal feedingstuff ☐ Technical use ☐ Other ☐

I.26. For transit to 3rd Country vis-à-vis EU ☐
3rd country | ISO code

I.27. For import or admission into EU ☐

I.28. Identification of the commodities

Species (Scientific name)	Nature of commodity	Approval number of establishments Manufacturing plant	Net weight

COUNTRY

Apiculture products

Part

II.a. Certificate reference number	II.b.

II. Health attestation

I, the undersigned official veterinarian, declare that I have read and understood Regulation (EC) No 1774/2002[(1)] and in particular Article 6 and Annex VIII Chapter IX thereof and certify that the apiculture by-products described above :

II.1. consist of apiculture products that satisfy the health requirements below;

II.2.

[(2)] *either* [are new and have not been in use before and have not come into contact with bees or used apiculture products;]

[(2)] *or* [have *been* subjected to a temperature of -12°C or lower for at least 24 hours;]

[(2)] *or* [in the *case* of wax, has been refined or rendered;]

II.3. come from an area which is not subject to any restrictions associated with :

a) American foul brood (*Paenibacillus larvae larvae*),

b) Acariosis (Acarapis woodi (*Rennie*)),

c) Small hive beetle (*Aethina tumida*), and

d) Tropilaelaps mites (*Tropilaelaps spp*),

and where the diseases mentioned above are officially notifiable.

Notes

Part I:

- Box reference I.6: Importer: this box is to be filled in only if it is a certificate for transit commodity.
- Box reference I.12: Place of destination: this box is to be filled in only if it is a certificate for transit commodity.
- Box reference I.15: Registration number (railway wagons or container and lorries), flight number (aircraft) or name (ship) and information is to be provided in the event of unloading and reloading.
- Box reference I.19: use the appropriate HS code: 05.11.99 and specify the commodity as listed under note Box reference I.28.
- Box reference I.23: for bulk containers, the container number and the seal number (if applicable) should be given.
- Box reference I.26 and I.27: fill in according to whether it is a transit or an import certificate.
- Box reference I.28: Nature of commodity: means honey, beeswax, royal jelly, propolis or pollen used in bee-keeping;

Part II:

(1) OJ L 273, 10.10.2002, p.1.

(2)Delete as appropriate.

- The signature and the stamp must be in a different colour to that of the printing.
- Note for the importer: This certificate is only for veterinary purposes and has to accompany the consignment until it reaches the border inspection post.

Official veterinarian

Name (in capitals): Qualification and title:

Date: Signature:

Stamp:

(v) Chapter 14(A) is amended as follows:

(i) The title to the health certificate is replaced by the following:

'Health certificate

For fat derivatives not intended for human consumption to be used for technical purposes, intended for dispatch to [or for transit through] [2] the European Community';

(ii) In the box in the second column of the model health certificate, the title is replaced by the following:

'VETERINARY CERTIFICATE

For fat derivatives not intended for human consumption to be used for technical purposes, intended for dispatch to [or for transit through] [2] the European Community';

(w) Chapter 14(B) is amended as follows:

(i) The title to the health certificate is replaced by the following:

'Health certificate

For fat derivatives not intended for human consumption to be used as feed or for technical purposes, intended for dispatch to [or for transit through] [2] the European Community';

(ii) In the box in the second column of the model health certificate, the title is replaced by the following:

'VETERINARY CERTIFICATE

For fat derivatives not intended for human consumption to be used as feed or for technical purposes, intended for dispatch to [or for transit through] [2] the European Community';

(x) Chapter 15 is amended as follows:

(i) The title to the health certificate is replaced by the following:

'Health certificate

For egg products not intended for human consumption that could be used as feed material, intended for dispatch to [or for transit through] [2] the European Community';

(ii) In the box in the second column of the model health certificate, the title is replaced by the following:

'VETERINARY CERTIFICATE

For egg products not intended for human consumption that could be used as feed material, intended for dispatch to [or for transit through] [2] the European Community';

(y) Chapter 16 is replaced by the following:

'CHAPTER 16

Model Declaration

Declaration by the importer of bones and bone products (excluding bone meal), horns and horn products (excluding horn meal) and hooves and hoof products (excluding hoof meal) intended for use other than as feed material, organic fertilisers or soil improvers for dispatch to the Europeam Communities

Note for the importer: This declaration is only for veterinary purposes and has to accompany the consignment until it reaches the border inspection post.

I, the undersigned, declare that the following products[5]:

(a) bones and bone products (excluding bone meal);

(b) horns and horn products (excluding horn meal);

(c) hooves and hoof products (excluding hoof meal);

are intended to be imported by me into the Community, and I declare that these products will not be diverted at any stage for any use in food, feed material, organic fertilizers or soil improvers and will be conveyed directly for the purpose of further processing or treatment to::

Name:..... Address:.....

The importer:

Name:..... Address:.....

Done at .....on.....

(place) *(date)*

Signature.....

Reference number as indicated on the common veterinary entry document (CVED) provided for in Annex III to Commission Regulation (EC) 136/2004:

.....

5 Delete as appropriate.

Official stamp of the border inspection post of entry into the EC [2]

Signature: .....

(Signature of the official veterinarian of the border inspection post)[6]

Name: .....

(Name in capital letters

6 The signature and the stamp must be in a different colour to that of the printing.

(g) The following Chapter 17 is added:

CHAPTER 17

Health certificate

For processed manure and processed manure products intended for dispatch to [or for transit through] [2] the European Community

COUNTRY **Veterinary certificate to EU**

Part I : Details of dispatched consignment

I.1. Consignor
Name
Address
Tel.N°

I.2. Certificate reference number | I.2.a

I.3. Central Competent Authority

I.4. Local Competent Authority

I.5. Consignee
Name
Address
Postal code
Tel.N°

I.6 Importer
Name
Address
Postal code
Tel.N°

I.7.Country of origin | ISO code | I.8. Region of origin | Code

I.9. Country of destination | ISO code | I.10. Region of destination | Code

I.11. Place of origin
Name | Approval number
Address

I.12. Place of destination
Establishment ☐ Custom warehouse ☐ Ship supplier ☐
Name | Approval number
Address
Postal code

I.13. Place of loading

I.14. Date of departure

I.15. Means of transport
Aeroplane ☐ Ship ☐ Railway wagon ☐
Road vehicle ☐ Other ☐
Identification:
Documentary references:

I.16. Entry BIP in EU

I.17.

I.18. Description of commodity

I.19. Commodity code (HS code)

I.20.Quantity

I.21 Temperature of product
Ambient ☐ Chilled ☐ Frozen ☐

I.22. Number of packages

I.23. Identification of container/Seal number

I.24.Type of packaging

I.25. Commodities certified for:
Animal feedingstuff ☐ Technical use ☐ Other ☐

I.26. For transit to 3rd Country vis-à-vis EU ☐
3rd country | ISO code

I.27. For import or admission into EU ☐

I.28. Identification of the commodities

Species (Scientific name)	Nature of commodity	Approval number of establishments: Manufacturing plant	Net weight

COUNTRY

Processed manure and processed manure products

Part II: Certification

II.a. Certificate reference number	II.b.

II. Health attestation

I, the undersigned official veterinarian, declare that I have read and understood Regulation (EC) No 1774/2002[1] and in particular Article 5 and Annex VIII Chapter VI thereof and certify that the processed manure or processed manure products described above :

II.1. come from a technical plant, a biogas plant or a composting plant approved by the competent authority of the third country meeting the special conditions laid down in Regulation (EC) No 1774/2002;

II.2.[2] have been subjected to:

[a heat treatment process of at least 70 °C for at least 60 minutes] or

[an equivalent treatment validated and authorized by the importing Member State in accordance with the specific conditions laid down in Regulation (EC) No 1774/2002 as follows .....
.....
.....;]

II.3. are:

(a) free from *Salmonella* (no *salmonella* in 25 g treated product);

(b) free from *Escherichia coli* or from *enterobacteriacea* (based on the aerobic count: less than 1000 cfu per gram of treated product); and

(c) have been subjected to reduction in spore-forming bacteria and toxic formation.

II.4. the manure described above is securely enclosed in:

(a) well-sealed and insulated containers, or

(ab) properly sealed packs (plastic bags or 'big bags').

Notes

Part I:

- Box reference I.6: Importer: this box is to be filled in only if it is a certificate for transit commodity.
- Box reference I.12: Place of destination: this box is to be filled in only if it is a certificate for transit commodity.
- Box reference I.15: Registration number (railway wagons or container and lorries), flight number (aircraft) or name (ship) and information is to be provided in the event of unloading and reloading.
- Box reference I.23: for bulk containers, the container number and the seal number (if applicable) should be given.
- Box reference I.26 and I.27: fill in according to whether it is a transit or an import certificate.
- Box reference I.31: Nature of commodity: enter if processed manure or if processed manure products.

Part II:

(1) OJ L 273, 10.10.2002, p.1.

(2)Delete as appropriate.

- The signature and the stamp must be in a different colour to that of the printing.
- Note for the importer: This certificate is only for veterinary purposes and has to accompany the consignment until it reaches the border inspection post.

Official veterinarian

Name (in capitals):

Qualification and title:

Date:

Signature:

Stamp:

(6) Annex XI is amended as follows:

(a) Part IX is replaced by the following:

'Part IX

List of third countries from which Member States may authorise imports of processed manure and processed manure products for the treatment of soil
(Health Certificate Chapter 17)

For processed manure and processed manure products, third countries listed in:

(a) Part 1 of Annex II to Decision 79/542/EEC;

(b) Annex I to Commission Decision 2004/211/EC; or

(c) Annex II to Commission Decision 2006/XXX/EC.'

(NOTE: Decision 2006/XXX/EC is reference to the now draft combined poultry import decision, voted in favour in December SCoFCAH)

(b) Part XIII is replaced by the following:

'Part XIII

List of third countries from which Member States may authorise imports of serum of equidae (Health Certificate Chapter 4(A))

Third countries or parts of third countries listed in Annex I to Commission Decision 2004/211/EC(8), from which the importation of equidae for breeding and production is allowed.'