Commission Regulation (EC) No 668/2004 of 10 March 2004 amending certain Annexes to Regulation (EC) No 1774/2002 of the European Parliament and of the Council, as regards the importation from third countries of animal by-products (Text with EEA relevance) (repealed)

COMMISSION REGULATION (EC) No 668/2004

of 10 March 2004

amending certain Annexes to Regulation (EC) No 1774/2002 of the European Parliament and of the Council, as regards the importation from third countries of animal by-products

(Text with EEA relevance) (repealed)

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)], as last amended by Commission Regulation (EC) No 808/2003[(2)], and in particular the second paragraph of Article 28, Article 29(3) and Article 32(1),

Whereas:

(1) Regulation (EC) No 1774/2002 provides that certain processed products that may be used as feed material and petfood, dogchews and technical products may be imported into the Community provided that they comply with the relevant requirements of that Regulation.

(2) Following the Scientific Steering Committee's opinion of 10 and 11 May 2001 on the safety of collagen it is appropriate to lay down the specific hygiene conditions to be applied for the processing and marketing of collagen that may be used as feed material. Annex VII to Regulation (EC) No 1774/2002 which sets out specific hygiene requirements for the processing and placing on the market of processed animal protein and other processed products that could be used as feed material should therefore be amended accordingly.

(3) Annex VIII to Regulation (EC) No 1774/2002 sets out requirements for the placing on the market of petfood, dogchews and technical products. It is necessary to amend that Annex, in order to introduce some technical amendments, to include the marking requirements of Article 28 of that Regulation for by-products intended for petfood derived from animals which have been treated with certain substances, and to clarify the import requirements applicable to fat derivatives and to certain processed products associated with the production of petfood, known as ‘flavouring innards’. Annex VIII should therefore be amended accordingly.

(4) Annex X to Regulation (EC) No 1774/2002 sets out a model health certificate for the importation from third countries of certain animal by-products and products derived therefrom. It is necessary to amend that Annex, in order to create additional models of importation certificates and to review the existing models to introduce some technical amendments including animal health considerations. Annex X should therefore be amended accordingly.

(5) 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. In the interests of clarity of Community legislation, those lists should in the near future be consolidated and combined with the lists of countries from which Member States are authorised to import products of various animal species which are already established under Community legislation for public health and animal health purposes. In the meantime, it is already appropriate to clarify and update the references made in Annex XI to those lists and Annex XI should therefore be amended accordingly.

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

HAS ADOPTED THIS REGULATION:

Article 1

Amendments to Regulation (EC) No 1774/2002

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

Article 2

Entry into force and applicability

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 1 May 2004.

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

ANNEX

Annexes I, VII, VIII, X and XI to Regulation (EC) No 1774/2002 are amended as follows:

1. Annex I is amended as follows:

   (a) definition number 40 is replaced by:

   '"petfood plant" means a plant producing petfood or dogchews or flavouring innards and in which certain animal by-products are used in the preparation of such petfood, dogchews or flavouring innards'

   (b) the following definition number 64 is added:

   '"flavouring innard" means a liquid or dehydrated processed product of animal origin used to enhance the palatability values of petfood'

2. Annex VII is amended as follows:

   (a) Chapter II is amended as follows:

      (i) point C(9)(d) is replaced by the following:
      if it is accompanied by a health certificate that conforms to the model set out in Chapter 1 of Annex X.

   (b) Chapter III is amended as follows:

      (i) point C(3)(a) is replaced by the following:
      come from third countries that appear on the list of part V and part VI of Annex XI as appropriate.

      (ii) point C(3)(d) is replaced by the following:
      if it is accompanied by a health certificate that conforms to the model set out in Chapter 4(B) of Annex X.

   (c) Chapter IV is amended as follows:

      (i) point B(2)(e) is replaced by the following:
      are accompanied by a health certificate that conforms to the model set out in Chapter 10(A) of Annex X.

      (ii) point C(3)(d) is replaced by the following:
      is accompanied by a health certificate that conforms to the model set out in Chapter 9 of Annex X.

   (d) Chapter VI is amended as follows:

      (i) point C(4)(d) is replaced by the following:
      are accompanied by a health certificate that conforms to the models set out in Chapter 11 and Chapter 12 of Annex X as appropriate.

   (e) Chapter VII is amended as follows:

      (i) point B(3)(d) is replaced by the following:
      is accompanied by a health certificate that conforms to the model set out in Chapter 12 of Annex X.

(f) Chapter VIII is amended as follows:

(i) point A(1)(b) is replaced by the following:

continuous cooking with steam at 145 °C during 30 minutes at 4 bars.

(ii) point B(2)(d) is replaced by the following:

is accompanied by a health certificate that conforms to the model set out in Chapter 12 of Annex X.

(g) the following Chapters IX and X are added:

'CHAPTER IX

Specific requirements for collagen

The following conditions apply in addition to the general conditions laid down in Chapter I.

A. Processing standards

1. Collagen must be produced by a process ensuring that unprocessed category 3 material is subjected to a treatment involving washing, pH adjustment using acid or alkali followed by one or more rinses, filtration and extrusion. After that treatment collagen may undergo a drying process.

2. The use of preservatives, other than those permitted under Community legislation shall be prohibited.

3. Collagen must be wrapped, packaged, stored and transported under satisfactory hygiene conditions. In particular:

(a) a room must be provided for storing materials for wrapping and packaging;

(b) wrapping and packaging must take place in a room or in a place intended for that purpose; and

(c) wrapping and packages containing collagen must be labelled with the words "collagen suitable for animal consumption".

B. Importation

4. Member States must authorise the importation of collagen if it:

(a) comes from a third country that appears on a Community list set out in Part XI of Annex XI;

(b) comes from a plant that appears on the list referred to in Article 29(4);

(c) has been produced in accordance with this Regulation; and

(d) is accompanied by a health certificate that conforms to the model set out in Chapter 11 of Annex X.'

‘CHAPTER X

Specific requirements for egg products

The following conditions apply in addition to the general conditions laid down in Chapter I.

A. Processing standards

1. Egg products must have been:

(a) submitted to any of processing Methods 1 to 5 or 7; or

(b) submitted to a method and parameters which ensure that the products comply with the microbiological standards set in Chapter I, paragraph 10; or

(c) treated in accordance with Chapter V of the Annex to Council Directive 89/437/EC[(3)] laying down hygiene and health problems affecting the production and the placing on the market of egg products.

B. Importation

2. Member States must authorise the importation of egg products if they:

(a) come from a third country that appears on a Community list set out in Part XVI of Annex XI;

(b) come from a plant that appears on the list referred to in Article 29(4);

(c) have been produced in accordance with this Regulation; and

(d) are accompanied by a health certificate that conforms to the model set out in Chapter 15 of Annex X.’

3. Annex VIII is amended as follows:

(a) Chapter IV is replaced by the following:

CHAPTER IV

Requirements for blood and blood products used for technical purposes, including pharmaceuticals, *in vitro* diagnosis and laboratory reagents, but excluding serum of equidae.

A. Importation

1. Imports of blood are subject to the requirements laid down in Chapter XI.

2. Member States must authorise importation of blood products if they:

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

(b) come from a plant approved by the competent authority of the third country meeting the specific conditions laid down in this Regulation; and

(c) are accompanied by a health certificate that conforms to the model set out in Chapter 4 C of Annex X; and

3. Member States must authorise importation of blood products if they originate in a third country or regions thereof where:
either:

(a) in the case of blood products derived from ruminant animals:

(i) the animals and the products come from a region where no case of foot-and-mouth disease, vesicular stomatitis, rinderpest, peste des petits ruminants, Rift Valley fever, African horse sickness and bluetongue[4] has been recorded for 12 months and in which vaccination has not been carried out against those diseases for at least 12 months in the susceptible species and from which imports of ruminant animals of the specified species are authorised pursuant to Community legislation. The blood from which such products are manufactured must have been collected:

— in slaughterhouses approved in accordance with Community legislation,

— from live animals in facilities approved in accordance with Community legislation; or

— in slaughterhouses approved and supervised by the competent authority of the third country. In this case, the Commission and Member States must be notified of the address and approval number of such slaughterhouse or the certificate shall indicate this information;

or

(ii) the products have undergone one of the following treatments guaranteeing the absence of pathogens of the ruminant diseases referred to in subparagraph (i):

— heat treatment at a temperature of 65 °C for at least three hours, followed by an effectiveness check,

— irradiation at 2,5 megarads or by gamma rays, followed by an effectiveness check,

— change in pH to pH 5 for two hours, followed by an effectiveness check,

— heat treatment of at least 90 °C throughout their substance, followed by an effectiveness check, or

— any other treatment provided for in accordance with the procedure referred to in Article 33(2);

(iii) by way of derogation from point (ii) above, a Member State may allow import from countries where sero-positive bluetongue animals are present, of blood and blood products intended for technical purposes including pharmaceuticals, *in vitro* diagnosis and laboratory reagents, provided that the approved technical plant of final destination is situated in the same Member State; the consignment must go directly to that plant and all precautions including safe disposal of waste, unused or surplus material must be taken to avoid risks of spreading diseases to animals or humans;

or

(b) in the case of blood products derived from animals belonging to the taxa Proboscidae and Artiodactyla, and their crossbreeds, other than ruminants:

(i) the animals and the products come from a region where no case of foot-and-mouth disease, swine vesicular disease, African horse sickness, classical swine fever, African swine fever, rinderpest, peste des petits ruminants, Newcastle disease or avian influenza has been recorded for 12 months in the susceptible species and in which vaccination has not been carried out against those diseases for at least 12 months;

or

(ii) the products have undergone one of the following treatments guaranteeing the absence of pathogens of the diseases referred to in subparagraph (i):

— heat treatment at a temperature of 65 °C for at least three hours, followed by an effectiveness check,

— irradiation at 2,5 megarads or by gamma rays, followed by an effectiveness check,

— heat treatment of at least 90 °C throughout their substance, followed by an effectiveness check, or

— any other treatment provided for in accordance with the procedure referred to in Article 33(2).

4. The specific conditions relating to imports of products for use *in vitro* diagnosis and laboratory reagents may be laid down, where necessary, under the procedure referred to in Article 33(2).

(b) Chapter V is amended as follows:

(i) point B(2)(a) is replaced by the following:

it comes from equidae born and raised in a third country that appears on the list of part XIII of Annex XI;

(ii) point B(2)(d) is replaced by the following:

it is accompanied by a health certificate that conforms to the model set out in Chapter 4(A) of Annex X

(c) Chapter VI is amended as follows:

(i) point C(5)(b) is replaced by the following:

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:

(i) for at least 12 months before dispatch, has been free from the following diseases:

— classical swine fever,

— African swine fever, and

— rinderpest, and

(ii) 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;

(ii) point C(6)(c) is replaced by the following:

they come either:

(i) 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(B) of Annex XI and they have been treated in accordance with paragraph 2; or

(ii) from animals originating from other regions of a third country or other third countries and they have been treated in accordance with paragraph 2(c) or (d); or

(iii) from ruminant animals and have been treated in accordance with paragraph 2 and 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 (C) of Annex XI. In this case, the certificate referred to in subparagraph (b) is replaced by a declaration conforming to the model laid down in Chapter 5(C) of Annex X, to the effect that or proving that those requirements have been met;

(d) Chapter VII is amended as follows:

(i) the following point B(5)(c) is added:

they come from a third country appearing on the list set out in part XV (A) of Annex XI.

(ii) point B(6)(a) is replaced by the following:

that appear on the lists set out in part XV(B) and (C) of Annex XI as appropriate; and

(e) Chapter VIII is amended as follows:

(i) the following point B(3)(c) is added:

they come from a third country that appears on the list of part VIII of Annex XI as appropriate.

(f) Chapter IX is replaced by:

'CHAPTER IX

Requirements for apiculture products

A. Raw material

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

(a) not come from an area which is subject of a prohibition order associated with an occurrence of:

(i) American foulbrood (*Paenibacillus larvae larvae*), except where the competent authority has assessed the risk to be negligible, issued a specific authorisation for use only in that Member State, and taken all other necessary measures to ensure no spread of that disease;

(ii) acariosis (*Acarapis woodi* (Rennie), except where the area of destination has obtained additional guarantees in accordance with Article 14(2) of Directive 92/65/EEC[5];

(iii) small hive beetle (*Aethina tumida*); or

(iv) Tropilaelaps spp. (*Tropilaelaps* spp); and

(b) meet the requirements provided for in Article 8(a) of Directive 92/65/EEC.

B. Importation

2. As the small hive beetle and *Tropilaelaps* spp. are not present in the Community, the following additional safeguards concerning importation of apiculture products have to be laid down.

3. Member States must authorise the importation of apiculture products intended for use in apiculture if they:

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

(b) (i) are new and have not been in use before and if they have not come into contact with bees or used apiculture products; or

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

(iii) in the case of wax, the material has been refined or rendered before exportation;

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

(g) Chapter X is amended as follows:

(i) the following point 1(d) is added:

they come from a third country appearing on the list set out in part XVII of Annex XI.

(ii) the fourth indent of point 2(a)(iv) is replaced by the following:

ashed for one hour to at least 800 °C to the core before drying, or

(iii) point 2(b) is replaced by the following:

a declaration of the importer that conforms to the model laid down in Chapter 16 of Annex X and that must be in at least one official language of the Member State through which the consignment first enters the Community and in at least one official language of the Member State of destination.

(iv) point 4 is replaced by the following:

4. Following the border check provided for in Directive 97/78/EC, and in accordance with the conditions laid down in Article 8(4) of that Directive, the material must be transported direct to the technical plant.

(h) Chapter XI is replaced by the following:

‘CHAPTER XI

Animal by-products for the manufacture of feed including petfood, and for pharmaceuticals and other technical products

Member States must authorise the importation of animal by-products intended for the manufacture of feed including petfood, and for pharmaceutical products and other technical products if they:

1. come from third countries appearing on the lists set out in part VI and VII(A) and (B) of Annex XI as appropriate;

2. consist only of animal by-products referred to in Article 6(1)(a) to (j) and/or, when intended to be used for petfood, material derived from animals treated as referred to in the second paragraph of Article 28;

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

3. have been deep-frozen at the plant of origin or have been preserved in accordance with Community legislation in such a way to prevent spoiling between dispatch and delivery to the plant of destination;

4. have undergone all precautions to avoid contamination with pathogenic agents;

5. were packed in new packaging preventing any leakage;

6. are accompanied by a certificate that conforms to the models set out in Chapter 8(A), Chapter 8(B) or Chapter 3(D) of Annex X;

7. following the border checks provided for in Directive 97/78/EC, and in accordance with the conditions laid down in Article 8(4) of that Directive, they are transported directly either:

   (a) to a petfood or technical plant, which has given the guarantee that the animal by-products shall be used only for the purpose of producing petfood or technical products as appropriate, as specified by the competent authority if necessary, and shall not leave the plant untreated other than for direct disposal; or

   (b) to an intermediate plant; or

(c) to an authorised and registered user or collection centre, which has given the guarantee that the animal by-products shall be used only for permitted purposes, as specified by the Competent Authority if necessary;

and

8.1. in the case of raw material for petfood production derived from animals which have been treated with certain substances prohibited in accordance with Directive 96/22/EC, as referred to in the second paragraph of Article 28 of this Regulation, it shall:

(a) be marked in the third country before entry into the territory of the Community by a cross of liquefied charcoal or activated carbon, on each outer side of each frozen block, in such a way that the marking covers at least 70 % of the diagonal length of the side of the frozen block and is at least 10 cm in width;

(b) in the case of material which is not frozen, be marked in the third country before entry into the territory of the Community by spraying it with liquefied charcoal or by applying charcoal powder in such a way that the charcoal is clearly visible on the material;

(c) be transported directly to:

(i) the petfood plant of destination in accordance with point 7(a) above;

or

(ii) an intermediate plant in accordance with point 7(b) above and from there directly to the petfood plant referred to under (i), provided that the intermediate plant:

— only handles material covered by this point 8.1, or

— only handles material destined for a petfood plant as referred to under (i);

and

(d) be manipulated to remove the marking provided for in (a) and (b) only in the petfood plant of destination and only immediately prior to use of the material for the manufacture of petfood;

8.2. where a consignment is made up of raw material, which has been treated as referred to in 8.1 above and other non-treated raw material, all the raw materials in the consignment must be marked as laid down in point 8.1(a) and (b) above.

8.3. the marking provided for in point 8.1(a) and (b) and 8.2 shall remain visible from the dispatch and until the delivery to the petfood plant of destination.’

(i) Chapter XII is replaced by the following:

'CHAPTER XII

Rendered fats from category 2 materials for oleochemical purposes

A. Processing standards

1. Rendered fats derived from category 2 material for oleochemical purposes must be produced using methods 1 to 5 as referred to in Annex V, Chapter III.

2. Rendered fats derived from ruminant animals must be purified in such a way that the maximum level of remaining total insoluble impurities does not exceed 0,15 % in weight;

B. Importation of rendered fats

3. Member States must authorise the importation of rendered fats derived from category 2 materials, intended to be processed using a method that at least meets the standards of one of the processes described in Annex VI, Chapter III, if it:

(a) comes from a third country that appears on a Community list set out in part IV of Annex XI;

(b) has been produced in accordance with this Regulation; and

(c) is accompanied by a health certificate that conforms to the model set out in Chapter 10(B) of Annex X.

4. The rendered fats must be conveyed by land and/or sea from the country of origin direct to a border inspection post in the Community.

5. Following the checks provided for in Directive 97/78/EC, and in accordance with the conditions laid down in Article 8(4) of that Directive, the rendered fats must be conveyed to a category 2 oleochemical plant where they are to be processed into fat derivatives.

6. The health certificate referred to in paragraph 3 must state that:

(i) the rendered fats will not be diverted for any use other than further processing by a method that at least meets the standards of one of the processes referred to in Chapter III of Annex VI; and

(ii) the resulting fat derivatives shall only be used in organic fertiliser or soil improvers or other technical uses, other than in cosmetics, pharmaceuticals and medical devices.

7. The health certificate provided for in paragraph 3 must be presented to the competent authority at the border inspection post at the first point of entry of the goods into the Community, and thereafter a copy must accompany the consignment until their arrival at the plant of destination.

8. Following the checks provided for in Directive 97/78/EC, and in accordance with the conditions laid down in Article 8(4) of that Directive, the rendered fats shall be transported directly to the plant of destination.'

(j) The following Chapters XIII and XIV are added:

'CHAPTER XIII

Fat derivatives

A. Processing standards

1. If rendered fat produced from category 2 material is used for the production of fat derivatives a method that at least meets the standards of one of the processes referred to in Chapter III of Annex VI shall be used.

B. Importation

2. Member States shall authorise the importation of fat derivatives only if a health certificate that conforms to the model set out in Chapters 14(A) or 14(B) of Annex X accompanies each consignment.

3. The health certificate referred to in paragraph 2 must state:

(a) whether or not the fat derivatives derive from category 2 or 3 materials;

(b) in the case of fat derivatives produced from category 2 material, that the products:

(i) have been produced using a method that at least meets the standards of one of the processes referred to in Chapter III of Annex VI; and

(ii) shall only be used in organic fertiliser or soil improvers or other technical uses, other than in cosmetics, pharmaceuticals and medical devices.

4. The health certificate provided for in paragraph 2 must be presented to the competent authority at the border inspection post at the first point of entry of the goods into the Community, and thereafter a copy must accompany the consignment until its arrival at the plant of destination.

5. Following the checks provided for in Directive 97/78/EC, and in accordance with the conditions laid down in Article 8(4) of that Directive, the fat derivatives shall be transported directly to the plants of destination.

CHAPTER XIV

Specific requirements for flavouring innards for the manufacture of pet food

The following conditions apply in addition to the requirements for approval laid down in Chapter I.

A. Raw Material

1. Only animal by-products referred to in Article 6(1)(a) to (j) may be used for the production of liquid/dehydrated processed products of animal origin used to enhance the palatability values of pet food.

B. Processing standards

2. Flavouring innards must have been submitted to a treatment method and parameters, which ensure that the product complies with the microbiological standards laid down in Annex VIII, paragraph 6 of Chapter II. After treatment, every precaution must be taken to ensure that the product is not exposed to contamination.

3. The end product must:

(a) be packed in new or sterilised packaging; or

(b) 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.

C. Importation

4. Member States must authorise the importation of flavouring innards if they:

(a) come from third countries that appear on the list set out in part VII(C) of Annex XI;

(b) come from petfood plants approved by the competent authority of the third country meeting the specific conditions laid down in Article 18;

(c) have been produced in accordance with this Regulation; and

(d) are accompanied by a health certificate that conforms to the model set out in Chapter 3(E) of Annex X.'

4. Annex X is replaced by the following:

'ANNEX X

MODEL HEALTH CERTIFICATES FOR THE IMPORTATION FROM THIRD COUNTRIES OF CERTAIN ANIMAL BY-PRODUCTS AND PRODUCTS DERIVED THEREFROM

Notes

a) Veterinary certificates shall be produced by the exporting country, based on the models appearing in this Annex X, according to the layout of the model that corresponds to the animal by-products concerned. They shall contain, in the numbered order that appears in the model, the attestations that are required for any third country and, as the case may be, those supplementary guarantees that are required for the exporting third country or part thereof.

b) The original of each certificate 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.

c) It shall be drawn up in at least one of the official languages of the EU Member State in which the inspection at the border post shall be carried out and of the EU Member State of destination. However, these Member States may allow other languages, if necessary, accompanied by an official translation.

d) If for reasons of identification of the items of the consignment, additional pages are attached to the certificate, these pages shall also be considered as forming part of the original of the certificate by the application of the signature and stamp of the certifying official veterinarian, in each of the pages.

e) When the certificate, including additional schedules referred to in (d), comprises more than one page, each page shall be numbered — (page number) of (total number of pages) — on its bottom and shall bear the code number of the certificate that has been designated by the competent authority on its top.

f) The original of the certificate must be completed and signed by an official veterinarian. In doing so, the competent authorities of the exporting country shall ensure that the principles of certification equivalent to those laid down in Council Directive 96/93/EC are followed (OJ L 13, 16.1.1997, p. 28).

g) The colour of the signature shall be different to that of the printing. The same rule applies to stamps other than those embossed or watermark.

h) The original of the certificate must accompany the consignment at the EU border inspection post.

CHAPTER 1

Health certificate

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

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

1. **Consignor** (name and address in full)	**VETERINARY CERTIFICATE** **For processed animal protein not intended for human consumption, including mixtures and products other than petfood containing such protein, intended for dispatch to the European Community** Reference number (1) ORIGINAL
2. **Consignee** (name and address in full)	3. **Origin of the processed animal protein or product** 3.1. Country: ..... 3.2. Code of territory: .....
	4. **Competent Authority** 4.1. Responsible Ministry: ..... 4.2. Certifying department: ....
5. **Intended destination of the processed animal protein or product** 5.1. EU Member State: ..... 5.2. Name and address of destination:	6. **Place of loading for exportation** ....
7. **Means of transport and consignment identification** 7.1. (Lorry, rail wagon, ship, or aircraft) (2) 7.2. Number of seal (if applicable): ..... 7.3. Registration number(s), ship name or flight number:	7.4. Nature of packaging: .... 7.5. Number of packages: ..... 7.6. Net weight: ..... 7.7. Lot/batch production reference number: .... 7.8. Nature of packaging: ....
8. **Identification of the processed animal protein or product** 8.1. Nature of the processed animal protein or product: ..... 8.2. Processed animal protein of: ..... *(animal species)* 8.3. Address and approval number of the approved establishment of origin:	
9. **Health attestation** I, the undersigned official veterinarian, declare that I have read and understood Regulation (EC) No 1774/2002 (3) and certify that: 9.1. 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:

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

(2) or [heating to at least 80 °C throughout its substance;]]

9.2. the competent authority examined a random sample immediately prior to dispatch and found it to comply with the following standards (4):

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

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

9.3. 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'

9.4. the end product was stored in enclosed storage;

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

Official stamp and signature

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

..... (signature of the official veterinarian) (5)

(stamp) (5)

..... (name, qualifications and title, in capital letters)

Notes

(1) Issued by the competent authority.
(2) Delete as appropriate.
(3) OJ L 273, 10.10.2002, p. 1.
(4) 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.
(5) The signature and the stamp must be in a different colour to that of the printing.

CHAPTER 2 (A)

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 the European Community

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

1. **Consignor** (name and address in full)	**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 the European Community** Reference number (1) ORIGINAL
2. **Consignee** (name and address in full)	3. **Origin of the milk/milk based product** 3.1. Country: ..... 3.2. Code of territory: .....
	4. **Competent Authority** 4.1. Responsible Ministry: ..... 4.2. Certifying department: ....
5. **Intended destination of the milk/milk-based products** 5.1. EU Member State: ..... 5.2. Name and address of the destination:	6. **Place of loading for exportation** ....
7. **Means of transport and consignment identification** (2) 7.1. Lorry, rail wagon, ship, or aircraft) (3) 7.2. Number of seal (if applicable): ..... 7.3. Registration number(s), ship name or flight number:	7.4. Nature of packaging: .... 7.5. Number of packages: ..... 7.6. Net weight: ..... 7.7. Lot/batch production reference number: ....
8. **Identification of the milk/milk based product** 8.1. Description of milk and milk-based product: ..... 8.2. Milk of: ..... *(animal species)* 8.3. Address and registration number of treatment or processing establishment (3):	
9. **Health attestation** I, the undersigned official veterinarian, declare that I have read and understood Regulation (EC) No 1774/2002 (4) and certify that: 9.1. ..... (exporting country), ... (region) (5), has been free from foot-and-mouth disease and rinderpest for 12 months immediately prior to export and has not practised vaccination against foot-and-mouth disease or rinderpest in the 12 months immediately prior to export;	

9.2. the milk and milk-based product referred to in this certificate:

(a) has been prepared from raw milk that comes from animals:

— not showing clinical signs of a diseases that can be transmitted through the milk to humans or animals, and

— belonging to holdings that are not under official restriction due to foot-and-mouth disease or rinderpest, and

(b) has undergone a process involving heating to ..... (temperature) for ..... (time), which ensured a negative reaction to the phosphatase test, followed by, in the case of dried milk or dried milk-based product, a drying process;

9.3. every precaution was taken to avoid contamination of the milk/milk-based product after processing;

9.4. the milk/milk-based product was packed:

(3) *either* [in new containers,]

(3) *or* [in vehicles or bulk containers disinfected prior to loading using a product approved by the competent authority,]

and the containers are marked so as to indicate the nature of the milk/milk-based product and bear labels indicating 'NOT FOR HUMAN CONSUMPTION'.

Official stamp and signature

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

.....
(signature of the official veterinarian) (6)

(stamp) (6)

.....
(name, qualifications and title, in capital letters)

Notes

(1) Issued by the competent authority.
(2) For goods vehicles the registration number should be given. For bulk containers, the container number and the seal number (if applicable) should be included.
(3) Delete as appropriate.
(4) OJ L 273, 10.10.2002, p. 1.
(5) For completion if the authorisation to import into the Community is restricted to certain regions of the third country concerned.
(6) The signature and the stamp must be in a different colour to that of the printing.

CHAPTER 2 (B)

Health certificate

For heat-treated milk-based products with a pH reduced to less than six not intended for human consumption and for dispatch to the European Community

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

1. **Consignor** (name and address in full)	**VETERINARY CERTIFICATE** **For heat-treated milk-based products with a pH reduced to less than six not intended for human consumption and for dispatch to the European Community** Reference number (1) ORIGINAL
2. **Consignee** (name and address in full)	3. **Origin of the milkbased product** 3.1. Country: ..... 3.2. Code of territory: .....
	4. **Competent authority** 4.1. Responsible Ministry: ..... 4.2. Certifying department: ....
5. **Intended destination of the milk-based products** 5.1. EU Member State: ..... 5.2. Name and address of the destination:	6. **Place of loading for exportation** ....
7. **Means of transport and consignment identification** (2) 7.1. (Lorry, railwagon, ship, or aircraft) (3) 7.2. Number of seal (if applicable): ..... 7.3. Registration number(s), ship name or flight number:	7.4. Nature of packaging: .... 7.5. Number of packages: ..... 7.6. Net weight: ..... 7.7. Lot/batch production reference number: ....
8. **Identification of milk based product** 8.1. Description of milk-based product: ..... 8.2. Milk of: ..... *(animal species)* 8.3. Address and registration number of treatment or processing establishment (3):	
9. **Health attestation** I, the undersigned official veterinarian, declare that I have read and understood Regulation (EC) No 1774/2002 (4) and certify that: 9.1. the milk-based product referred to in this certificate: (a) has been prepared from raw milk that comes from animals:	

(i) not showing clinical signs of a diseases that can be transmitted through the milk to humans or animals; and

(ii) belonging to holdings that are not under official restriction due to foot-and-mouth disease or rinderpest;

(b) has undergone a process involving heating to (temperature) ..... for ..... (time), which ensured a negative reaction to the phosphatase test, followed by, in the case of dried milk or dried milk-based product, a drying process; and

(c) has undergone an acidification process whereby its pH has been maintained at less than 6 for at least one hour;

9.2. every precaution was taken to avoid contamination of the milk-based product after processing;

9.3. the milk-based product was packed:

(3) *either* [in new containers,]

(3) *or* [in vehicles or bulk containers disinfected prior to loading using a product approved by the competent authority,]

and the containers are marked so as to indicate the nature of the milk-based product and bear labels indicating 'NOT FOR HUMAN CONSUMPTION'.

Official stamp and signature

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

.....
(signature of the official veterinarian) (5)

(stamp) (5)

.....
(name, qualifications and title, in capital letters)

Notes

(1) Issued by the competent authority.
(2) For goods vehicles the registration number should be given. For bulk containers, the container number and the seal number (if applicable) should be included.
(3) Delete as appropriate.
(4) OJ L 273, 10.10.2002, p. 1.
(5) The signature and the stamp must be in a different colour to that of the printing.

CHAPTER 2 (C)

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 the European Community

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

1. **Consignor** (name and address in full)	**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 the European Community** Reference number (1) ORIGINAL
2. **Consignee** (name and address in full)	3. **Origin of the milk/milk based product** 3.1. Country: ..... 3.2. Code of territory: .....
	4. **Competent Authority** 4.1. Responsible Ministry: ..... 4.2. Certifying department: ....
5. **Intended destination of the milk/milk-based products** 5.1. EU Member State: ..... 5.2. Name and address of the destination:	6. **Place of loading for exportation** ....
7. **Means of transport and consignment identification** (2) 7.1. (Lorry, rail wagon, ship, or aircraft) (3) 7.2. Number of seal (if applicable): ..... 7.3. Registration number(s), ship name or flight number:	7.4. Nature of packaging: .... 7.5. Number of packages: ..... 7.6. Net weight: ..... 7.7. Lot/batch production reference number: ....
8. **Identification of milk/milk based product** 8.1. Description of milk/milk-based product: ..... 8.2. Milk of: ..... *(animal species)* 8.3. Address and registration number of treatment or processing establishment (3):	
9. **Health attestation** I, the undersigned official veterinarian, declare that I have read and understood Regulation (EC) No 1774/2002 (4) and certify that: 9.1. the milk/milk-based product referred to in this certificate: (a) has been prepared from raw milk that comes from animals:	

(i) not showing clinical signs of a diseases that can be transmitted through the milk to humans or animals; and

(ii) belonging to holdings that are not under official restriction due to foot-and-mouth disease or rinderpest; and

(b) has undergone:

(3) *either* (i) a sterilisation process whereby an Fc value equal to or greater that 3 is achieved;]

(3) *or* (ii) an initial process involving heating to ..... (temperature) for ..... (time), which ensured a negative reaction to the phosphatase test, followed by a further process involving heating to ..... (temperature) for ..... (time), which ensured a negative reaction to the phosphatase test, followed, in the case of dried milk, or dried milk-based products, by a drying process;]

9.2. every precaution was taken to avoid contamination of the milk/milk-based product after processing;

9.3. the milk/milk-based product was packed:

(3) *either* [in new containers,]

(3) *or* [in vehicles or bulk containers disinfected prior to loading using a product approved by the competent authority,]

and the containers are marked so as to indicate the nature of the milk/milk-based product and bear labels indicating 'NOT FOR HUMAN CONSUMPTION'.

Official stamp and signature

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

.....
(signature of the official veterinarian) (5)

(stamp) (5)

.....
(name, qualifications and title, in capital letters)

Notes

(1) Issued by the competent authority.
(2) For goods vehicles the registration number should be given. For bulk containers, the container number and the seal number (if applicable) should be included.
(3) Delete as appropriate.
(4) OJ L 273, 10.10.2002, p. 1.
(5) The signature and the stamp must be in a different colour to that of the printing.

CHAPTER 3 (A)

Health certificate

For canned petfood intended for dispatch to the European Community

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

1. **Consignor** (name and address in full)	**VETERINARY CERTIFICATE** **For canned petfood intended for dispatch to the European Community** Reference number (1) ORIGINAL
2. **Consignee** (name and address in full)	3. **Origin of petfood** 3.1. Country: ..... 3.2. Code of territory: .....
	4. **Competent authority** 4.1. Responsible Ministry: ..... 4.2. Certifying department: ....
5. **Destination of petfood** 5.1. EU Member State: ..... 5.2. Name and address of the destination:	6. **Place of loading for exportation** ....
7. **Means of transport and consignment identification** (2) 7.1. (Lorry, rail wagon, ship, or aircraft) (3) 7.2. Number of seal (if applicable): ..... 7.3. Registration number(s), ship name or flight number:	7.4. Nature of packaging: .... 7.5. Number of packages: ..... 7.6. Net weight: ..... 7.7. Lot/batch production reference number: ....
8. **Identification of petfood** 8.1. The petfood was produced from raw material of the following species: .... *(animal species)* 8.2. Address and registration number of the approved establishment:	
9. **Health attestation** I, the undersigned official veterinarian, declare that I have read and understood Regulation (EC) No 1774/2002 (4) and certify that the petfood described above: 9.1. 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; 9.2. has been prepared exclusively with the following animal by-products:	

(3) *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;]

(3) *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;]

(3) *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;]

(3) *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;]

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

(3) *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;]

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

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

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

(3) *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;]

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

9.4. 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 9.1;

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

Official stamp and signature

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

.....
(stamp) (5) (signature of the official veterinarian) (5)

.....
(name, qualifications and title, in capital letters)

Notes

(1) Issued by the competent authority.
(2) For goods vehicles the registration number should be given. For bulk containers, the container number and the seal number (if applicable) should be included.
(3) Delete as appropriate.
(4) OJ L 273, 10.10.2002, p. 1.
(5) The signature and the stamp must be in a different colour to that of the printing.

CHAPTER 3 (B)

Health certificate

For processed petfood other than canned petfood, intended for dispatch to the European Community

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

1. **Consignor** (name and address in full)	**VETERINARY CERTIFICATE** **For processed petfood other than canned petfood, intended for dispatch to the European Community** Reference number (1) ORIGINAL
2. **Consignee** (name and address in full)	3. **Origin of petfood** 3.1. Country: ..... 3.2. Code of territory: ..... 4. **Competent authority** 4.1. Responsible Ministry: ..... 4.2. Certifying department: ....
5. **Destination of petfood** 5.1. EU Member State: ..... 5.2. Name and address of the destination:	6. **Place of loading for exportation** ....
7. **Means of transport and consignment identification** (2) 7.1. (Lorry, rail wagon, ship, or aircraft) (3) 7.2. Number of seal (if applicable): ..... 7.3. Registration number(s), ship name or flight number:	7.4. Nature of packaging: .... 7.5. Number of packages: ..... 7.6. Net weight: ..... 7.7. Lot/batch production reference number: ....
8. **Identification of petfood** 8.1. The petfood was produced from raw material of the following species: .... *(animal species)* 8.2. Address and registration number of the approved establishment:	
9. **Health attestation** I, the undersigned official veterinarian, declare that I have read and understood Regulation (EC) No 1774/2002 (4) and certify that the petfood described above: 9.1. 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; 9.2. has been prepared exclusively with the following animal by-products:	

(3) *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;]

(3) *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;]

(3) *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;]

(3) *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;]

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

(3) *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;]

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

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

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

(3) *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;]

9.3. was subjected to a heat treatment of at least 90 °C throughout its substance;

9.4. was analysed by a 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 (5):

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

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

9.5. has undergone all precautions to avoid contamination with pathogenic agents after treatment;

9.6. was packed in new packaging, which bear labels indicating 'NOT FOR HUMAN CONSUMPTION'

Official stamp and signature

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

.....
(stamp) (6) (signature of the official veterinarian) (6)

.....
(name, qualifications and title, in capital letters)

Notes

(1) Issued by the competent authority.
(2) For goods vehicles the registration number should be given. For bulk containers, the container number and the seal number (if applicable) should be included.
(3) Delete as appropriate.
(4) OJ L 273, 10.10.2002, p. 1.
(5) 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.
(6) The signature and the stamp must be in a different colour to that of the printing.

CHAPTER 3 (C)

Health certificate

For dogchews intended for dispatch to the European Community

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

1. **Consignor** (name and address in full)	**VETERINARY CERTIFICATE** **For dogchews intended for dispatch to the European Community** Reference number (1) ORIGINAL
2. **Consignee** (name and address in full)	3. **Origin of dogchews** 3.1. Country: ..... 3.2. Code of territory: ..... 4. **Competent authority** 4.1. Responsible Ministry: ..... 4.2. Certifying department: ....
5. **Destination of dogchews** 5.1. EU Member State: ..... 5.2. Name and address of the destination:	6. **Place of loading for exportation** ....
7. **Means of transport and consignment identification** (2) 7.1. (Lorry, rail wagon, ship, or aircraft) (3) 7.2. Number of seal (if applicable): ..... 7.3. Registration number(s), ship name or flight number:	7.4. Nature of packaging: .... 7.5. Number of packages: ..... 7.6. Net weight: .....
8. **Identification of dogchews** 8.1. The dogchews were produced from raw material of the following species: .... *(animal species)* 8.2. Address and registration number of the approved establishment:	
9. **Health attestation** I, the undersigned official veterinarian, declare that I have read and understood Regulation (EC) No 1774/2002 (4) and certify that the dogchews described above: 9.1. 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; 9.2. have been prepared exclusively with the following animal by-products: (3) *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;]	

(3) *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;]

(3) *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;]

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

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

9.3. have been subjected:

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

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

9.4. 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 (5):

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

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

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

9.6. were packed in new packaging.

Official stamp and signature

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

.....
(stamp) (6) (signature of the official veterinarian) (6)

.....
(name, qualifications and title, in capital letters)

Notes

(1) Issued by the competent authority.
(2) For goods vehicles the registration number should be given. For bulk containers, the container number and the seal number (if applicable) should be included.
(3) Delete as appropriate.
(4) OJ L 273, 10.10.2002, p. 1.
(5) 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.
(6) The signature and the stamp must be in a different colour to that of the printing.

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 the European Community

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

1. **Consignor** (name and address in full)	**VETERINARY CERTIFICATE** **For raw petfood for direct sale or animal by-products to be fed to farmed fur animals, intended for dispatch to the European Community** Reference number (1) ORIGINAL
2. **Consignee** (name and address in full)	3. **Origin of raw petfood/animal by-products** (2) 3.1. Country: ..... 3.2. Code of territory: .....
	4. **Competent authority** 4.1. Responsible Ministry: ..... 4.2. Certifying department: ....
5. **Destination of raw petfood/animal by-products** (2) 5.1. EU Member State: ..... 5.2. Name and address of the destination:	6. **Place of loading for exportation** ....
7. **Means of transport and consignment identification** (3) 7.1. (Lorry, rail wagon, ship, or aircraft) (2) 7.2. Number of seal (if applicable): ..... 7.3. Registration number(s), ship name or flight number:	7.4. Nature of packaging: .... 7.5. Number of packages: ..... 7.6. Net weight: ..... 7.7. Lot/batch production reference number: ....
8. **Identification of raw petfood/animal by-products** (2) 8.1. The raw petfood/animal by-products (2) derive from the following species: .... *(animal species)* 8.2. Address and registration number of the approved establishment:	
9. **Health attestation** I, the undersigned official veterinarian, declare that I have read and understood Regulation (EC) No 1774/2002 (4) and certify that the raw petfood or animal by-product described above: 9.1. consist of animal by-products that satisfy the health requirements below; 9.2. 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 (5), 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 94/984/EC (6), 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 (7), , 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 (8) on animal welfare;

9.3. consist only of the following animal by-products:

(a) in the case of animal by-product for use in feed for farmed fur animals:

(a) (i) 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

(a) (ii) 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 carcases that are fit for human consumption in accordance with Community legislation;

(b) in the case of animal by-products for use in raw petfood:

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

9.4. 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;

9.5. 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;

9.6. in the case of raw petfood, 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.

Official stamp and signature

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

(stamp) (9)

.....
(signature of the official veterinarian) (9)

.....
(name, qualifications and title, in capital letters)

Notes

(1) Issued by the competent authority.
(2) Delete as appropriate.
(3) For goods vehicles the registration number should be given. For bulk containers, the container number and the seal number (if applicable) should be included.
(4) OJ L 273, 10.10.2002, p. 1.
(5) [SANCO/10167/2002 Rev 21 on Community health conditions on imports of animals and fresh meat including minced meat from third countries and amending Decisions 79/542/EEC, 2000/572/EC and 2000/585/EC.]
(6) Commission Decision 94/984/EC laying down animal health conditions and veterinary certificates for the importation of fresh poultry meat from third countries.
(7) Commission Decision 2000/585/EC of 7 September 2000 laying down animal and public health conditions and veterinary certifications for import of wild and farmed game meat and rabbit meat from third countries.
(8) Council Directive 93/119/EC of 22 December 1993 on the protection of animals at the time of slaughter or killing.
(9) The signature and the stamp must be in a different colour to that of the printing.

CHAPTER 3 (E)

Health certificate

For flavouring innards for use in the manufacture of petfood, intended for dispatch to the European Community

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

1. **Consignor** (name and address in full).....	**VETERINARY CERTIFICATE** **For flavouring innards for use in the manufacture of petfood, intended for dispatch to the European Community** Reference number (1) ORIGINAL
2. **Consignee** (name and address in full)	3. **Origin of the flavouring innards products** 3.1. Country: ..... 3.2. Code of territory: .....
5. **Destination of the flavouring innards products** 5.1. EU Member State: ..... 5.2. Name and address of the destination:	4. **Competent Authority** 4.1. Responsible Ministry: ..... 4.2. Certifying department: ..... 6. **Place of loading for exportation**
7. **Means of transport and consignment identification** (2) 7.1. (Lorry, rail wagon, ship, or aircraft) (3) 7.2. Number of seal (if applicable): ..... 7.3. Registration number(s), ship name or flight number: .	7.4. Nature of packaging: ..... 7.5. Number of packages: ..... 7.6. Net weight: ..... 7.7. Lot/batch production reference number:
8. **Identification of the flavouring innards products** 8.1. Nature of flavouring innards products: ..... 8.2. Species of animals from which the flavouring innards products derive: ..... 8.3. Address and registration number of the approved establishment:	
9. **Health attestation** I, the undersigned official veterinarian, declare that I have read and understood Regulation (EC) No 1774/2002 (4) and certify that the flavouring innards products described above: 9.1. consist of animal by-products that satisfy the animal health requirement below;	

9.2. 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;

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

(3) *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;]

(3) *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;]

(3) *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;]

(3) *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;]

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

(3) *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;]

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

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

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

(3) *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;]

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

9.5. have been examined by the competent authority taking a random sample immediately prior to dispatch and found it to comply with the following standards (5):

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

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

9.6. the end product was:

(3) *either* [packed in new or sterilised bags,]

(3) *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';

9.7. the end product was stored in enclosed storage;

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

Official stamp and signature

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

(stamp) (6)

..... (signature of the official veterinarian) (6)

..... (name, qualifications and title, in capital letters)

Notes

(1) Issued by the competent authority.
(2) For goods vehicles the registration number should be given. For bulk containers, the container number and the seal number (if applicable) should be included.
(3) Delete as appropriate.
(4) OJ L 273, 10.10.2002, p. 1.
(5) 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.
(6) The signature and the stamp must be in a different colour to that of the printing.

CHAPTER 4 (A)

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 the European Community

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

1. **Consignor** (name and address in full)	**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 the European Community** Reference number (1) ORIGINAL
2. **Consignee** (name and address in full)	3. **Origin of the serum** 3.1. Country: ..... 3.2. Code of territory: .....
	4. **Competent authority** 4.1. Responsible Ministry: ..... 4.2. Certifying department: ....
5. **Destination of the serum** 5.1. EU Member State: ..... 5.2. Name and address of the destination:	6. **Place of loading for exportation** ....
7. **Means of transport and consignment identification** (2) 7.1. (Lorry, rail wagon, ship, or aircraft) (3) 7.2. Number of seal (if applicable): ..... 7.3. Registration number(s), ship name or flight number:	7.4. Nature of packaging: .... 7.5. Number of packages: .... 7.6. Net weight: .....
8. **Identification of the serum** 8.1. Serum of: ..... *(animal species)* 8.2. Address and veterinary control number of the registered establishment of collection:	
9. **Health attestation** I, the undersigned official veterinarian, declare that I have read and understood Regulation (EC) No 1774/2002 (4) and certify that the serum of equidae described above: 9.1. consist of serum from equidae that satisfy the health requirements below; 9.2. consist exclusively of serum of equidae not intended for human nor animal consumption; 9.3. comes from a country where the following diseases are compulsorily notifiable: African horse sickness, dourine, glanders, equine encephalomyelitis (all types including VEE), equine infectious anemia, vesicular stomatitis, rabies, anthrax;	

9.4. was obtained, under the supervision of a veterinarian, from equidae which, at the time of collection, were free from clinical signs of infectious disease or were obtained from equidae that passed ante-mortem inspection at the time of slaughter;

9.5. was obtained from equidae that have remained since birth in the territory or, in case of official regionalisation according to Community legislation, in parts of the territory of a third country in which:

(a) Venezuelan equine encephalomyelitis has not occurred during the last two years;

(b) dourine has not occurred during the last six months; and

(c) glanders has not occurred during the last six months;

9.6. was obtained from equidae that had never been present on a holding that had been subject to prohibition for animal health reasons or where:

(3) *either* [(a) in the case of equine encephalomyelitis, the date on which all the equidae suffering from the disease were slaughtered was at least six months before the date of collection;

(b) in the case of infectious anaemia, all the infected animals had been slaughtered and the remaining animals showed a negative reaction to two Coggins tests carried out three months apart;

(c) in the case of vesicular stomatitis, the prohibition was lifted at least six months before the date of collection;

(d) in the case of rabies, the last recorded case was at least a month before the date of collection; and

(e) in the case of anthrax, the last recorded case was at least 15 days before the date of collection;]

(3) *or* [all the animals of species susceptible to the disease located on the holding were slaughtered and the premises disinfected, at least 30 days before the date of collection (or, in the case of anthrax, at least 15 days before);]

9.7. has undergone all precautions to avoid contamination with pathogenic agents during production, handling and packaging;

9.8. was packed in sealed impermeable containers clearly labelled 'serum from equidae' and bearing the registration number of the establishment of collection.

Official stamp and signature

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

(stamp) (5)

..... (signature of the official veterinarian) (5)

..... (name, qualifications and title, in capital letters)

Notes

(1) Issued by the competent authority.
(2) For goods vehicles the registration number should be given. For bulk containers, the container number and the seal number (if applicable) should be included.
(3) Delete as appropriate.
(4) OJ L 273, 10.10.2002, p. 1.
(5) The signature and the stamp must be in a different colour to that of the printing.

CHAPTER 4 (B)

Health certificate

For blood products not intended for human consumption that could be used as feed material, intended for dispatch to the European Community

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

1. **Consignor** (name and address in full)	**VETERINARY CERTIFICATE** **For blood products not intended for human consumption that could be used as feed material, intended for dispatch to the European Community** Reference number (1) ORIGINAL
2. **Consignee** (name and address in full)	3. **Origin of the blood products** 3.1. Country: ..... 3.2. Code of territory: .....
	4. **Competent Authority** 4.1. Responsible Ministry: ..... 4.2. Certifying department: ....
5. **Destination of the blood products** 5.1. EU Member State: ..... 5.2. Name and address of the destination:	6. **Place of loading for exportation** ....
7. **Means of transport and consignment identification** (2) 7.1. (Lorry, rail wagon, ship, or aircraft) (3) 7.2. Number of seal (if applicable): ..... 7.3. Registration number(s), ship name or flight number:	7.4. Nature of packaging: .... 7.5. Number of packages: ..... 7.6. Net weight: ..... 7.7. Lot/batch production reference number: ....
8. **Identification of the blood products** 8.1. Nature of the blood products: ..... 8.2. Species of animals from which the blood products derive: .... 8.3. Address and registration number of the approved establishment:	
9. **Health attestation** I, the undersigned official veterinarian, declare that I have read and understood Regulation (EC) No 1774/2002 (4) and certify that the blood products described above: 9.1. consist of blood products that satisfy the health requirements below; 9.2. consist exclusively of blood products not intended for human consumption;	

9.3. have 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;

9.4. have been prepared (derived) exclusively with the following animal by-products:

(3) *either* [blood of slaughtered animals, which is fit for human consumption in accordance with Community legislation, but is not intended for human consumption for commercial reasons;]

(3) *and/or* [blood of slaughtered animals, which is rejected as unfit for human consumption but is not affected by any signs of diseases communicable to humans or animals, derived from carcases that are fit for human consumption in accordance with Community legislation;]

9.5. have been submitted:

(3) *either* [to processing in accordance with processing method ..... (5) as set out in Annex V, Chapter III of Regulation (EC) No 1774/2002/EC,]

(3) *or* [to a method and parameters which ensure that the product complies with the microbiological standards set in Chapter I, paragraph 10 of Regulation (EC) No 1774/2002/EC,]

in order to kill pathogenic agents;

9.6. have been examined by the competent authority taking a random sample immediately prior to dispatch and found it to comply with the following standards (6):

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

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

9.7. the end product was:

(3) *either* [packed in new or sterilised bags,]

(3) *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';

9.8. the end product was stored in enclosed storage;

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

Official stamp and signature

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

(stamp) (7)

.....
(signature of the official veterinarian) (7)

.....
(name, qualifications and title, in capital letters)

Notes

(1) Issued by the competent authority.
(2) For goods vehicles the registration number should be given. For bulk containers, the container number and the seal number (if applicable) should be included.
(3) Delete as appropriate.
(4) OJ L 273, 10.10.2002, p. 1.
(5) Insert method 1 to 5 or 7 as applicable
(6) 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.
(7) The signature and the stamp must be in a different colour to that of the printing.

CHAPTER 4 (C)

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 the European Community

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

1. **Consignor** (name and address in full)	**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 the European Community** Reference number (1) ORIGINAL
2. **Consignee** (name and address in full)	3. **Origin of the blood products** 3.1. Country: ..... 3.2. Code of territory: .....
	4. **Competent Authority** 4.1. Responsible Ministry: ..... 4.2. Certifying department: ....
5. **Destination of the blood products** 5.1. EU Member State: ..... 5.2. Name and address of the destination:	6. **Place of loading for exportation** ....
7. **Means of transport and consignment identification** (2) 7.1. (Lorry, rail wagon, ship, or aircraft) (3) 7.2. Number of seal (if applicable): ..... 7.3. Registration number(s), ship name or flight number:	7.4. Nature of packaging: .... 7.5. Number of packages: ..... 7.6. Net weight: ..... 7.7. Lot/batch production reference number: ....
8. **Identification of the blood products** 8.1. Nature of the blood products: ..... 8.2. Species of animals from which the blood products derive: .... 8.3. Address and registration number of the approved establishment:	
9. **Health attestation** I, the undersigned official veterinarian, declare that I have read and understood Regulation (EC) No 1774/2002 (4) and certify that the blood products described above: 9.1. consist of blood products that satisfy the health requirements below; 9.2. consist exclusively of blood products not intended for human or animal consumption;	

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

(3) *either* [- blood of slaughtered animals, which is fit for human consumption in accordance with Community legislation, but is not intended for human consumption for commercial reasons;]

(3) *and/or* [- blood of slaughtered animals, which is rejected as unfit for human consumption but is not affected by any signs of diseases communicable to humans or animals, derived from carcasses that are fit for human consumption in accordance with Community legislation;]

(3) *and/or* [- blood obtained from animals other than ruminants that are 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;]

(3) *and/or* [- blood and blood products derived from the production of products intended for human consumption;]

(3) *and/or* [- blood and blood products originating from animals that did not show clinical signs of any disease communicable through that product to humans or animals;]

(3) *either* [9.4. in the case of blood products derived from ruminant animals they originate in a third country or regions where:

(3) *either* [the animals and products come from a region where no case of foot-and-mouth disease, vesicular stomatitis, rinderpest, peste des petits ruminants, Rift Valley fever and bluetongue (5) has been recorded for 12 months and in which vaccination has not been carried out against those diseases for at least 12 months and from which imports of ruminant animals are authorised pursuant to Community legislation. The blood from which such products are manufactured must have been collected:

(3) *either* [in slaughterhouses approved in accordance with Community legislation,]

(3) *or* [from live animals in facilities approved in accordance with Community legislation,]

(3) *or* [in slaughterhouses approved and supervised by the competent authority of the third country. In this case the Commission and Member States must be notified of the address and approval number of such slaughterhouse and the certificate shall indicate this information,]]

(3) *or* [the products have undergone one of the following treatments, guaranteeing the absence of pathogens of the ruminant diseases foot-and-mouth disease, vesicular stomatitis, rinderpest, peste des petits ruminants, Rift Valley fever and bluetongue (5):

(3) *either* [heat treatment at a temperature of 65 °C for at least three hours, followed by an effectiveness check,]

(3) *or* [irradiation at 2,5 megarads or by gamma rays, followed by an effectiveness check,]

(3) *or* [change in pH to pH 5 for two hours, followed by an effectiveness check,]

(3) *or* [heat treatment of at least 90 °C throughout their substance, followed by an effectiveness check,]]

(3) *or* [sero-positive bluetongue animals are present, and the blood and blood products are intended for technical purposes including pharmaceuticals, in vitro diagnosis and laboratory reagents, to be processed in the approved plants [approval number] in [Member State] (6)]]

(3) *or* [9.4. in the case of blood products derived from animals excluding ruminants they originate in a third country or regions where:

(3) *either* [the animals and the products come from a region where no case of foot-and-mouth disease, swine vesicular disease, classical swine fever, African swine fever, Newcastle disease or avian influenza has been recorded for 12 months in the susceptible species and in which vaccination has not been carried out against those diseases for at least 12 months. The health certificate shall follow the model according to the species of animal from which the blood products are derived;]

(3) *or* [the products have undergone a heat treatment at a temperature of 65 °C for at least three hours, followed by an effectiveness check, guaranteeing the absence of pathogens of the following diseases:

foot-and-mouth disease, swine vesicular disease, classical swine fever, African swine fever, Newcastle disease or avian influenza in the susceptible species;]]

9.5. the end product was:

(3) *either* [packed in new or sterilised bags,]

(3) *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 OR ANIMAL CONSUMPTION';

9.6. the end product was stored in enclosed storage;

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

Official stamp and signature

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

(stamp) (7)

.....
(signature of the official veterinarian) (7)

.....
(name, qualifications and title, in capital letters)

Notes

(1) Issued by the competent authority.
(2) For goods vehicles the registration number should be given. For bulk containers, the container number and the seal number (if applicable) should be included.
(3) Delete as appropriate.
(4) OJ L 273, 10.10.2002, p. 1.
(5) In the case of countries in which bluetongue sero-positive ruminant animals are present, blood products have been treated or the animals have been tested seronegative.
(6) This must be the same Member State of first entry of the products into the Community.
(7) The signature and the stamp must be in a different colour to that of the printing.

CHAPTER 5 (A)

Health certificate

For fresh or chilled hides and skins of ungulates, intended for dispatch to the European Community

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

1. **Consignor** (name and address in full)	**VETERINARY CERTIFICATE** **For fresh or chilled hides and skins of ungulates, intended for dispatch to the European Community** Reference number (1) ORIGINAL
2. **Consignee** (name and address in full)	3. **Origin of the hides and skins** 3.1. Country: ..... 3.2. Code of territory: .....
	4. **Competent Authority** 4.1. Responsible Ministry: ..... 4.2. Certifying department: ....
5. **Destination of the hides and skins** 5.1. EU Member State: ..... 5.2. Name and address of the destination:	6. **Place of loading for exportation** ....
7. **Means of transport and consignment identification** (2) 7.1. (Lorry, rail wagon, ship, or aircraft) (3) 7.2. Number of seal (if applicable): ..... 7.3. Registration number(s), ship name or flight number:	7.4. Nature of packaging: .... 7.5. Number of packages: ..... 7.6. Net weight: ..... 7.7. Number(s) of the seal(s) on the container(s), road vehicle(s), railway wagon(s) or bale(s): ....
8. **Identification of the hides and skins** 8.1. Hides and skins of: ..... *(animal species)* 8.2. Address and veterinary control number of the registered and supervised establishment:	
9. **Health attestation** I, the undersigned official veterinarian, declare that I have read and understood Regulation (EC) No 1774/2002 (4) and certify that the hides and skins described above: 9.1. have been obtained from animals that were: (a) slaughtered in a slaughterhouse, underwent ante mortem inspection and were fit, as a result of such inspection, for slaughter in accordance with Community legislation; (b) not affected by any signs of diseases communicable to humans or animals; and	

(c) not killed to eradicate any epizootic disease;

9.2. 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:

▶(1) (a) has been free, for at least 12 months before dispatch, from the following diseases: ◀

(3) *either* [classical swine fever, and

- African swine fever,]

(3) and/*or* [-rinderpest,]

and

(b) has been free for at least 24 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;

9.3. have been obtained from:

(3) *either* [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;]

(3) *or* [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;]

(3) *or* [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;]

(3) *or* [animals that have passed the ante-mortem health inspection at the slaughterhouse during the 24 hours before slaughter and have shown no evidence of [foot-and-mouth disease], [rinderpest], [classical swine fever], [African swine fever] or [swine vesicular disease] (3);]

9.4. have undergone all precautions to avoid recontamination with pathogenic agents.

Official stamp and signature

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

.....
(stamp) (5) (signature of the official veterinarian) (5)

.....
(name, qualifications and title, in capital letters)

Notes

(1) Issued by the competent authority.
(2) For goods vehicles the registration number should be given. For bulk containers, the container number and the seal number (if applicable) should be included.
(3) Delete as appropriate.
(4) OJ L 273, 10.10.2002, p. 1.
(5) The signature and the stamp must be in a different colour to that of the printing.

CHAPTER 5 (B)

Health certificate

For treated hides and skins of ungulates, intended for dispatch to the European Community

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

1. **Consignor** (name and address in full)
.....
.....
.....
.....
.....
.....
.....
.....

2. **Consignee** (name and address in full)
.....
.....
.....
.....
.....
.....

VETERINARY CERTIFICATE
For treated hides and skins of ungulates, intended for dispatch to the European Community

Reference number (1) ORIGINAL

3. **Origin of the hides and skins**
3.1. Country: .....
3.2. Code of territory: .....

4. **Competent authority**
4.1. Responsible Ministry: .....
4.2. Certifying department: .....
.....

5. **Destination of the hides and skins**
5.1. EU Member State: .....
5.2. Name and address of the destination: .....
.....
.....
.....

6. **Place of loading for exportation**
.....
.....
.....
.....

7. **Means of transport and consignment identification** (2)
7.1. (Lorry, rail wagon, ship, or aircraft) (3)
7.2. Number of seal (if applicable): .....
7.3. Registration number(s), ship name or flight number: .
.....
.....
.....

7.4. Nature of packaging: .....
.....
7.5. Number of packages: .....
7.6. Net weight: .....
7.7. Number(s) of the seal(s) on the container(s), road vehicle(s), railway wagon(s) or bale(s):
.....
.....
.....

8. **Identification of the hides and skins**
8.1. Hides and skins of: ..... *(animal species)*
8.2. Address and veterinary control number of the registered and supervised establishment: .....
.....

9. **Health attestation**
I, the undersigned official veterinarian, declare that I have read and understood Regulation (EC) No 1774/2002 (4) and certify that the hides and skins described above:
9.1. have been obtained from animals that:
(a) did not show any clinical signs of any disease communicable to humans or animals, and
(b) were not killed to eradicate any epizootic disease;

(3) *either* [9.2. come from animals originating in a country or a part of a country not subject, pursuant to Community legislation, to restrictions as a result of an outbreak of a serious transmissible disease to which the animals of the species concerned are susceptible and have been:

(3) *either* [dried;]

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

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

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

(3) *or* [9.2. have been :

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

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

(3) *or* [9.2. were salted on ..... (date) before being transported by ship;]

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.

Official stamp and signature

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

(stamp) (5)

..... (signature of the official veterinarian) (5)

..... (name, qualifications and title, in capital letters)

Notes

(1) Issued by the competent authority.
(2) For goods vehicles the registration number should be given. For bulk containers, the container number and the seal number (if applicable) should be included.
(3) Delete as appropriate.
(4) OJ L 273, 10.10.2002, p. 1.
(5) The signature and the stamp must be in a different colour to that of the printing.

CHAPTER 5 (C)

Official declaration

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

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

1. **Consignor** (name and address in full)	**Official declaration** **For treated hides and skins of ungulates ruminants that are intended for dispatch to the European Community and have been kept separate for 21 days or will undergo transport for 21 uninterrupted days before importation** Reference number (1) ORIGINAL
2. **Consignee** (name and address in full)	3. **Origin of the hides and skins** 3.1. Country: ..... 3.2. Code of territory: .....
	4. **Competent Authority** 4.1. Responsible Ministry: ..... 4.2. Certifying department: ....
5. **Destination of the hides and skins** 5.1. EU Member State: ..... 5.2. Name and address of the destination:	6. **Place of loading for exportation** ....
7. **Means of transport and consignment identification** (2) 7.1. (lorry, rail wagon, ship, or aircraft) (3) 7.2. Number of seal (if applicable): ..... 7.3. Registration number(s), ship name or flight number:	7.4. Nature of packaging: .... 7.5. Number of packages: ..... 7.6. Net weight: ..... 7.7. Number(s) of the seal(s) on the container(s), road vehicle(s), railway wagon(s) or bale(s): ....
8. **Identification of the hides and skins** 8.1. Hides and skins of: ..... *(animal species)* 8.2. Address and veterinary control number of the establishment:	
9. **Health attestation** I, the undersigned official veterinarian, declare that I have read and understood Regulation (EC) No 1774/2002 (4) and certify that the hides and skins described above: 9.1. have been obtained from animals that: (a) did not show any clinical signs of any disease communicable to humans or animals, and (b) were not killed to eradicate any epizootic disease; 9.2. have been:	

(3) *either* [dried;]

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

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

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

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

(3) *either* [9.4. have been kept separate immediately before dispatch for 21 days under official supervision after the treatment described under point (9.2)]

(3) *or* [9.4. following the declaration of the transporter, the duration of the transport period is foreseen to be at least 21 days]

Official stamp and signature

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

.....
(signature of the official veterinarian) (5)

(stamp) (5)

.....
(name, qualifications and title, in capital letters)

Notes

(1) Issued by the competent authority.
(2) For goods vehicles the registration number should be given. For bulk containers, the container number and the seal number (if applicable) should be included.
(3) Delete as appropriate.
(4) OJ L 273, 10.10.2002, p. 1.
(5) The signature and the stamp must be in a different colour to that of the printing.

CHAPTER 6 (A)

Health certificate

For treated game trophies of birds and ungulates, being solely bones, horns, hooves, claws, antlers, teeth, hides or skins, for dispatch to the European Community

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

1. **Consignor** (name and address in full)	**Health certificate** **For treated game trophies of birds and ungulates, being solely bones, horns, hooves, claws, antlers, teeth, hides or skins, for dispatch to the European Community** Reference number (1) ORIGINAL
2. **Consignee** (name and address in full)	3. **Origin of game trophies** 3.1. Country: ..... 3.2. Code of territory: .....
	4. **Competent Authority** 4.1. Responsible Ministry: ..... 4.2. Certifying department: ....
5. **Destination of the game trophies** 5.1. EU Member State: ..... 5.2. Name and address of the destination:	6. **Place of loading for exportation** ....
7. **Means of transport and consignment identification** (2) 7.1. (Lorry, rail wagon, ship, or aircraft) (3) 7.2. Number of seal (if applicable): ..... 7.3. Registration number(s), ship name or flight number:	7.4. Nature of packaging: .... 7.5. Number of parts or packages: .... 7.6. Reference number of Cites certificate: ....
8. **Identification of game trophies** 8.1. Game trophies of: ..... *(animal species)* 8.2. Nature of the game trophies: (a) solely [bones], [horns], [hooves], [claws], [antlers], [teeth] (3): ..... (b) solely [hides] or [skins] (3):	
9. **Health attestation** I, the undersigned official veterinarian, declare that I have read and understood Regulation (EC) No 1774/2002 (4) and certify that the game trophies described above: 9.1. have been packaged, immediately after treatment, without being in contact with other products of animal origin likely to contaminate them, in individual, transparent and closed packages so as to avoid any subsequent contamination;	

(3) *either* [9.2. in the case of game trophies consisting solely of hides or skin:

(3) *either* [have been dried]

(3) *or* [have been dry-salted or wet-salted for a minimum of 14 days before dispatch]

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

(3) *or* [9.2. in the case of game trophies consisting solely of bone, horns, hooves, claws, antlers or teeth:

(a) have been immersed in boiling water for an appropriate time so as to ensure that any matter other than bone, horns, hooves, claws, antlers or teeth is removed, and

(b) have been disinfected with a product authorised by the competent authority, in particular with hydrogen peroxide where parts consisting of bone are concerned]

Official stamp and signature

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

(stamp) (5)

..... (signature of the official veterinarian) (5)

..... (name, qualifications and title, in capital letters)

Notes

(1) Issued by the competent authority.
(2) For goods vehicles the registration number should be given. For bulk containers, the container number and the seal number (if applicable) should be included.
(3) Delete as appropriate.
(4) OJ L 273, 10.10.2002, p. 1.
(5) The signature and the stamp must be in a different colour to that of the printing.

CHAPTER 6 (B)

Health certificate

For game trophies of birds and ungulates consisting of entire parts not having been treated, intended for dispatch to the European Community

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

1. **Consignor** (name and address in full)	**Health certificate** **For game trophies of birds and ungulates consisting of entire parts not having been treated, intended for dispatch to the European Community** Reference number (1) ORIGINAL
2. **Consignee** (name and address in full)	3. **Origin of game trophies** 3.1. Country: ..... 3.2. Code of territory: .....
	4. **Competent authority** 4.1. Responsible Ministry: ..... 4.2. Certifying department: ....
5. **Destination of the game trophies** 5.1. EU Member State: ..... 5.2. Name and address of the destination:	6. **Place of loading for exportation** ....
7. **Means of transport and consignment identification** (2) 7.1. (Lorry, rail wagon, ship, or aircraft) (3) 7.2. Number of seal (if applicable): ..... 7.3. Registration number(s), ship name or flight number:	7.4. Nature of packaging: .... 7.5. Number of parts or packages: .... 7.6. Reference number of Cites certificate: ....
8. **Identification of game trophies** 8.1. Game trophies of: ..... *(animal species)*	
9. **Health attestation** I, the undersigned official veterinarian, declare that I have read and understood Regulation (EC) No 1774/2002 (4) and certify that the game trophies described above: (3) *either* [9.1. with respect to game trophies of cloven-hoofed animals, excluding swine: (a) ..... (region) has been free from foot-and-mouth disease and rinderpest for the previous 12 months, and during the same period, no vaccination against any of those diseases has taken place; and (b) the game trophies described above:	

(b) (i) were obtained from animals which were killed in the territory of that region, which is authorised for export of fresh meat of the corresponding susceptible domestic species and where, during the last 60 days, there have been no animal health restrictions because of outbreaks of diseases to which the game animals are susceptible, and

(b) (ii) originated from animals that were killed at a distance of at least 20 km from the borders of another third country or part of a third country not authorised to export untreated game trophies of cloven-hoofed animals other than swine to the Community;]

(3) *or* [9.1. with respect to game trophies of wild swine:

(a) ..... (region) during the last 12 months was free from classical swine fever, African swine fever, swine vesicular disease, foot-and-mouth disease and porcine enteroviral encephalmiyelitis (Teschen disease) and no vaccinations have been carried out against any of those diseases during the last 12 months, and

(b) the game trophies described above:

(b) (i) were obtained from animals which were killed in that territory, which is authorised for export of fresh meat of the corresponding susceptible domestic species and where, during the last 60 days, there have been no animal health restrictions because of outbreaks of diseases to which the swine are susceptible, and

(b) (ii) originated from animals that were killed at a distance of at least 20 km from the borders of another third country or part of a third country not authorised to export untreated game trophies of wild swine to the Community;]

(3) *or* [9.1. with respect to game trophies of solipeds, the game trophies described above were obtained from wild solipeds that were killed in the territory of the exporting country mentioned above;]

(3) *or* [9.1. with respect to game trophies of game birds:

(a) ..... (region) is free from avian influenza and Newcastle disease, and

(b) the game trophies described above were obtained from wild game birds that were killed in that region and where during the last 30 days there have been no animal health restrictions because of outbreaks of disease to which the wild birds are susceptible;]

9.2. The game trophies described above have been packaged without being in contact with other products of animal origin likely to contaminate them, in individual, transparent and closed packages so as to avoid any subsequent contamination.

Official stamp and signature

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

(stamp) (5)

..... (signature of the official veterinarian) (5)

..... (name, qualifications and title, in capital letters)

Notes

(1) Issued by the competent authority.
(2) For goods vehicles the registration number should be given. For bulk containers, the container number and the seal number (if applicable) should be included).
(3) Delete as appropriate.
(4) OJ L 273, 10.10.2002, p. 1.
(5) The signature and the stamp must be in a different colour to that of the printing.

CHAPTER 7 (A)

Health certificate

For pig bristles from third countries or regions thereof that are free from African swine fever, intended for dispatch to the European Community

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

1. **Consignor** (name and address in full)	**VETERINARY CERTIFICATE** **For pig bristles from third countries or regions thereof that are free from African swine fever, intended for dispatch to the European Community** Reference number (1) ORIGINAL
2. **Consignee** (name and address in full)	3. **Origin of the pig bristles** 3.1. Country: ..... 3.2. Code of territory: .....
	4. **Competent authority** 4.1. Responsible Ministry: ..... 4.2. Certifying department: ....
5. **Destination of the pig bristles** 5.1. EU Member State: ..... 5.2. Name and address of the destination:	6. **Place of loading for exportation** ....
7. **Means of transport and consignment identification** (2) 7.1. (Lorry, rail wagon, ship, or aircraft) (3) 7.2. Number of seal (if applicable): ..... 7.3. Registration number(s), ship name or flight number:	7.4. Nature of packaging: .... 7.5. Number of parts or packages: .... 7.6. Net weight: .....
8. **Identification of the pig bristles** 8.1. Address and veterinary control number of the registered establishment:	
9. **Health attestation** I, the undersigned official veterinarian, declare that I have read and understood Regulation (EC) No 1774/2002 (4) and certify that: 9.1. the pig bristles described above have been obtained from pigs originating, and slaughtered in a slaughterhouse, in the country of origin; 9.2. the pigs from which the pig bristles have been obtained did not show during inspection, carried out at the time of slaughtering, signs of diseases communicable to humans or animals and were not killed to eradicate any epizootic disease; 9.3. the country of origin or, in case of regionalisation according to Community legislation, the region of origin, has been free from African swine fever for at least 12 months; 9.4. the pig bristles are dry and securely enclosed in packaging.	

Official stamp and signature

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

(stamp) (5)

..... (signature of the official veterinarian) (5)

..... (name, qualifications and title, in capital letters)

Notas

(1) Issued by the competent authority.
(2) For goods vehicles the registration number should be given. For bulk containers, the container number and the seal number (if applicable) should be included.
(3) Delete as appropriate.
(4) OJ L 273, 10.10.2002, p. 1.
(5) The signature and the stamp must be in a different colour to that of the printing.

CHAPTER 7 (B)

Health certificate

For pig bristles from third countries or regions thereof that are not free from African swine fever, intended for dispatch to the European Community

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

1. **Consignor** (name and address in full)	**VETERINARY CERTIFICATE** **For pig bristles from third countries or regions thereof that are not free from African swine fever, intended for dispatch to the European Community** Reference number (1) ORIGINAL
2. **Consignee** (name and address in full)	3. **Origin of the pig bristles** 3.1. Country: ..... 3.2. Code of territory: .....
	4. **Competent authority** 4.1. Responsible Ministry: ..... 4.2. Certifying department: ....
5. **Destination of the pig bristles** 5.1. EU Member State: ..... 5.2. Name and address of the destination:	6. **Place of loading for exportation** ....
7. **Means of transport and consignment identification** (2) 7.1. (Lorry, rail wagon, ship, or aircraft) (3) 7.2. Number of seal (if applicable): ..... 7.3. Registration number(s), ship name or flight number:	7.4. Nature of packaging: .... 7.5. Number of parts or packages: .... 7.6. Net weight: .....
8. **Identification of the pig bristles** 8.1. Address and veterinary control number of the registered establishment:	
9. **Health attestation** I, the undersigned official veterinarian, declare that I have read and understood Regulation (EC) No 1774/2002 (4) and certify that: 9.1. the pig bristles described above have been obtained from pigs originating, and slaughtered in a slaughterhouse, in the country of origin; 9.2. the pigs from which the pig bristles have been obtained did not show during inspection, carried out at the time of slaughtering, signs of diseases communicable to humans or animals and were not killed to eradicate any epizootic disease; 9.3. the pig bristles mentioned above have been: (3) *either* [boiled;] (3) *or* [dyed;]	

(3) *or* [bleached;]

9.4. the pig bristles are dry and securely enclosed in packaging.

Official stamp and signature

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

.....
(stamp) (5) (signature of the official veterinarian) (5)

.....
(name, qualifications and title, in capital letters)

Notes

(1) Issued by the competent authority.
(2) For goods vehicles the registration number should be given. For bulk containers, the container number and the seal number (if applicable) should be included.
(3) Delete as appropriate.
(4) OJ L 273, 10.10.2002, p. 1.
(5) The signature and the stamp must be in a different colour to that of the printing.

CHAPTER 8 (A)

Health certificate

For animal by-products (1) for the manufacture of petfood, intended for dispatch to the European Community

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

1. **Consignor** (name and address in full)	**Health certificate** **For animal by-products (1) for the manufacture of petfood, intended for dispatch to the European Community** Reference number (2) ORIGINAL
2. **Consignee** (name and address in full)	3. **Origin of animal by-products** 3.1. Country: ..... 3.2. Code of territory: .....
5. **Destination of animal by-products** 5.1. EU Member State: ..... 5.2. Name and address of the destination:	4. **Competent authority** 4.1. Responsible Ministry: ..... 4.2. Certifying department: .... 6. **Place of loading for exportation** ....
7. **Means of transport and consignment identification** (3) 7.1. (Lorry, rail wagon, ship, or aircraft) (4) 7.2. Number of seal (if applicable): ..... 7.3. Registration number(s), ship name or flight number:	7.4. Nature of packaging: .... 7.5. Number of packages: ..... 7.6. Net weight: ..... 7.7. Lot/batch production reference number: ....
8. **Identification of animal by-products** 8.1. Nature of animal by-products: ..... 8.2. Animal by-products of: ..... *(animal species)* 8.3. Address and veterinary control number of the approved establishment:	
9. **Health attestation** I, the undersigned official veterinarian, declare that I have read and understood Regulation (EC) No 1774/2002 (5) and certify that the animal by-products described above: 9.1. consist of animal by-products that satisfy the animal health requirements below;	

9.2. have been obtained in the territory of ..... (6) from animals:

(4) *either* [(a) that have remained in this territory since birth or for at least the last three months before slaughter;]

(4) *or* [(b) killed in the wild in this territory (7);]

9.3. have been obtained from animals:

(4) *either* [(a) coming from holdings:

(a) (i) where, for the following diseases for which the animals are susceptible, there has been neither case/outbreak of rinderpest, swine vesicular disease, Newcastle disease or avian influenza during the prior 30 days, nor of classical or African swine fever during the prior 40 days; nor in the holdings situated in their vicinity within 10 km, during the prior 30 days; and

(a) (ii) where there has been neither case/outbreak of foot and mouth disease during the prior 60 days, nor in the holdings situated in their vicinity within 25 km, during the prior 30 days; and

(b) which:

(b) (i) were not killed to eradicate any epizootic disease;

(b) (ii) have remained in their holdings of origin for at least forty days before departure and which have been transported directly to the slaughterhouse without contact with other animals which did not comply with the same health conditions;

(b) (iii) 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 to above for which the animals are susceptible; and

(b) (iv) 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 on animal welfare;]

(4) *or* [(a) captured and killed in the wild in an area:

(a) (i) in which within 25 km there has been no case/outbreak of any of the following diseases for which the animals are susceptible: foot and mouth disease, rinderpest, Newcastle disease or avian influenza during the prior 30 days , nor of classical or African swine fever during the prior 40 days; and

(a) (ii) that is situated at a distance that exceeds 20 km from the borders separating another territory of a country or part thereof, which is not authorised at these dates for exporting this material to the European Community; and

(b) which after killing were transported within 12 hours for chilling either to a collection centre and immediately afterwards to a game establishment, or directly to a game establishment;]

9.4. have been obtained in an establishment around which, within a radius of 10 km, there has been no case/outbreak of diseases referred to in point 9.3 for which the animals are susceptible during the prior 30 days or, in the event of a case of disease, the preparation of raw material for exportation to the European Community has been authorised only after removal of all meat, and the total cleaning and disinfection of the establishment under the control of an official veterinarian;

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

9.6. have been packed in new packaging preventing any leakage and in officially sealed containers bearing the label indicating 'RAW MATERIAL ONLY FOR THE MANUFACTURE OF PETFOOD' and the name and address of the EU establishment of destination;

9.7. consist only of the following animal by-products:

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

(4) *and/or* [- 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;]

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

(4) *and/or* [- former foodstuffs of animal origin, or former foodstuffs containing products of animal origin, other than catering waste (8) 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;]

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

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

(4) *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;]

(4) *and/or* [- raw material derived from animals which have been treated with certain substances prohibited in accordance with Directive 96/22/EC for the manufacture of petfood, as referred to in Article 28 of Regulation (EC) No 1774/2002;]

9.8. have been deep-frozen at the plant of origin or have been preserved in accordance with EU legislation in such a way that they will not spoil between dispatch and delivery to the plant of destination;

9.9. in the case of raw material derived from animals which have been treated with certain substances prohibited in accordance with Directive 96/22/EC for the manufacture of petfood , as referred to in Article 28 of Regulation (EC) No 1774/2002:

(a) it has been marked in the third country before entry into the territory of the Community by a cross of liquefied charcoal or activated carbon on each outer side of each frozen block in a way that the marking covers at least 70 % of the diagonal length of the frozen block and be of at least 10 cm width;

(b) in case of material which is not frozen, the raw material has been marked in the third country before entry into the territory of the Community by spraying it with liquefied charcoal or by applying charcoal powder in a way that the charcoal is clearly visible on the material; and

(c) in the case the animal by-products are made up of raw material which has been treated as referred to above and other non-treated raw material, all the raw materials have been marked as laid down in point (a) and (b) above.

(4) (9) [10. **Specific requirements**

(4) (10) 10.1. The by-products in this consignment come from animals that have been kept in the territory mentioned under (9.2), where vaccination programmes against foot-and-mouth disease are being regularly carried out and officially controlled in domestic bovine animals.

(4) (11) 10.2. The by-products in this consignment consists only of animal by-products derived from trimmed offal of domestic ruminants, which have maturated at an ambient temperature of more than +2 °C for at least three hours, or in the case of masseter muscles of bovine animals and de-boned meat of domestic animals, for at least 24 hours.]

Official stamp and signature

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

.....
(signature of the official veterinarian) (12)

(stamp) (12)

.....
(name, qualifications and title, in capital letters)

Notes

(1) Excluding raw blood, raw milk, hides and skins, hooves and horn, pig bristles and feathers (see relevant specific certificates for the import of these products).
(2) Issued by the competent authority.
(3) For goods vehicles the registration number should be given. For bulk containers, the container number and the seal number (if applicable) should be included).
(4) Delete as appropriate.
(5) OJ L 273, 10.10.2002, p. 1.
(6) The name and ISO code number of the exporting country as laid down in:
— part 1 of Annex II of Council Decision 79/542/EEC;
— the Annex to Commission Decision 94/984/EC; and
— the Annex to Commission Decision 2000/585/EC.
In addition the ISO code of regionalisation in this Annex (where applicable for the susceptible species concerned) should be included).
(7) Only for countries from where game meat intended for human consumption of the same animal species is authorised for importation into the European Community.
(8) Catering waste means all waste food, including used cooking oils, originating in restaurants, catering facilities and kitchens, including central kitchens and household kitchens.
(9) Supplementary guarantees to be provided when the material of domestic ruminants originated in the territory of a South American or South African country or part thereof from where only maturated and boned fresh meat of domestic ruminants for human consumption is permitted for exportation to the European Community. In the case of offal only trimmed offal of domestic ruminants which must be exclusively offal from which the bones, cartilage, trachea and main bronchi, lymphatic glands adhering connective tissue, fat and mucus have been completely removed is permitted. The whole masseter muscles of bovine animals, incised in accordance with paragraph 41(a) of Chapter VIII of Annex I to Council Directive 64/433/EEC, are also permitted.
(10) Only for certain South American countries.
(11) Only for certain South American and South African countries.
(12) The signature and the stamp must be in a different colour to that of the printing.

CHAPTER 8 (B)

Health certificate

For animal by-products for the manufacture of technical products (including pharmaceutical products) (1) , intended for dispatch to the European Community

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

1. **Consignor** (name and address in full)	**Health certificate** **For animal by-products for the manufacture of technical products (including pharmaceutical products) (1) , intended for dispatch to the European Community** Reference number (2) ORIGINAL
2. **Consignee** (name and address in full)	3. **Origin of animal by-products** 3.1. Country: ..... 3.2. Code of territory: .....
	4. **Competent authority** 4.1. Responsible Ministry: ..... 4.2. Certifying department: ....
5. **Destination of animal by-products** 5.1. EU Member State: ..... 5.2. Name and address of the destination:	6. **Place of loading for exportation** ....
7. **Means of transport and consignment identification** (3) 7.1. (Lorry, rail wagon, ship, or aircraft) (4) 7.2. Number of seal (if applicable): ..... 7.3. Registration number(s), ship name or flight number:	7.4. Nature of packaging: .... 7.5. Number of packages: ..... 7.6. Net weight: ..... 7.7. Lot/batch production reference number: ....
8. **Identification of animal by-products** 8.1. Nature of animal by-products: ..... 8.2. Animal by-products of: ..... *(animal species)* 8.2. Address and veterinary control number of the approved establishment:	
9. **Health attestation** I, the undersigned official veterinarian, declare that I have read and understood Regulation (EC) No 1774/2002 (5) and certify that the animal by-products described above: 9.1. consist of animal by-products that satisfy the animal health requirements below;	

9.2. have been obtained in the territory of: ..... (6) from animals:

(4) *either* [(a) that have remained in this territory since birth or for at least the last three months before slaughter;]

(4) *or* [(b) killed in the wild in this territory (7);]

9.3. have been obtained from animals:

(4) *either* [(a) coming from holdings:

(i) where, for the following diseases for which the animals are susceptible, there has been neither case/outbreak of rinderpest, swine vesicular disease, Newcastle disease or avian influenza during the prior 30 days; nor of classical or African swine fever during the prior 40 days; nor in the holdings situated in their vicinity within 10 km, during the prior 30 days; and

(ii) where there has been neither case/outbreak of foot and mouth disease during the prior 60 days, nor in the holdings situated in their vicinity within 25 km, during the prior 30 days; and

(b) which:

(i) were not killed to eradicate any epizootic disease;

(ii) have remained in their holdings of origin for at least forty days before departure and which have been transported directly to the slaughterhouse without contact with other animals which did not comply with the same health conditions;

(iii)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 to above for which the animals are susceptible; and

(iv) 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 on animal welfare;]

(4) *or* [(a) captured and killed in the wild in an area:

(i) in which within 25 km there has been no case/outbreak of any of the following diseases for which the animals are susceptible: foot and mouth disease, rinderpest, Newcastle disease or avian influenza during the prior 30 days nor of classical or African swine fever during the prior 40 days and

(ii) that is situated at a distance that exceeds 20 km from the borders separating another territory of a country or part thereof, which is not authorised at these dates for exporting this material to the European Community; and

(b) which after killing were transported within 12 hours for chilling either to a collection centre and immediately afterwards to a game establishment, or directly to a game establishment;]

9.4. have been obtained in an establishment around which, within a radius of 10 km, there has been no case/outbreak of diseases referred to in point 9.3 for which the animals are susceptible during the prior 30 days or, in the event of a case of disease, the preparation of raw material for exportation to the European Community has been authorised only after removal of all meat, and the total cleaning and disinfection of the establishment under the control of an official veterinarian;

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

9.6. have been packed in new packaging preventing any leakage and in officially sealed containers bearing the label indicating 'RAW MATERIAL ONLY FOR THE MANUFACTURE OF TECHNICAL PRODUCTS INCLUDING PHARMACEUTICAL PRODUCTS' and the name and address of the EU establishment of destination;

9.7. consist only of the following animal by-products:

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

(4) *and/or* [- 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;]

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

(4) *and/or* [- former foodstuffs of animal origin, or former foodstuffs containing products of animal origin, other than catering waste (8) 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;]

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

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

(4) *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;]

(4) *and/or* [- fur originating from animals that did not show clinical signs of any disease communicable through that product to humans or animals;]

9.8. have been deep-frozen at the plant of origin or have been preserved in accordance with EU legislation in such a way that they will not spoil between dispatch and delivery to the plant of destination.

(4) (9) [10. **Specific requirements**

(4) (10) 10.1. The by-products in this consignment come from animals that have been obtained in the territory mentioned under (9.2), where vaccination programmes against foot-and-mouth disease are being regularly carried out and officially controlled in domestic bovine animals.

(4) (11) 10.2. The by-products in this consignment consists of animal by-products derived from offal or boned meat.]

Official stamp and signature

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

.....
(signature of the official veterinarian) (12)

(stamp) (12)

.....
(name, qualifications and title, in capital letters)

Notes

(1) Excluding raw blood, raw milk, hides and skins, pig bristles and feathers (see relevant specific certificates for the import of these products).
(2) Issued by the competent authority.
(3) For goods vehicles the registration number should be given. For bulk containers, the container number and the seal number (if applicable) should be included.
(4) Delete as appropriate.
(5) OJ L 273, 10.10.2002, p. 1.
(6) The name and ISO code number of the exporting country as laid down in:
— part 1 of Annex II of Council Decision 79/542/EEC;
— the Annex to Commission Decision 94/984/EC; and
— the Annex to Commission Decision 2000/585/EC.
In addition the ISO code of regionalisation in this Annex (where applicable for the susceptible species concerned) should be included.
(7) Only for countries from where game meat intended for human consumption of the same animal species is authorised for importation into the European Community.
(8) Catering waste means all waste food, including used cooking oils, originating in restaurants, catering facilities and kitchens, including central kitchens and household kitchens.
(9) Supplementary guarantees to be provided when the material of domestic ruminants originated in the territory of a South American or South African country or part thereof from where only maturated and boned fresh meat of domestic ruminants for human consumption is permitted for exportation to the European Community. In the case of offal only trimmed offal of domestic ruminants which must be exclusively offal from which the bones, cartilage, trachea and main bronchi, lymphatic glands adhering connective tissue, fat and mucus have been completely removed is permitted. The whole masseter muscles of bovine animals, incised in accordance with paragraph 41(a) of Chapter VIII of Annex I to Council Directive 64/433/EEC, are also permitted.
(10) Only for certain South American countries.
(11) Only for certain South American and South African countries.
(12) The signature and the stamp must be in a different colour to that of the printing.

CHAPTER 9

Health certificate

For fish oil not intended for human consumption to be used as feed material or for technical purposes, intended for dispatch to the European Community

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

1. **Consignor** (name and address in full)	**VETERINARY CERTIFICATE** **For fish oil not intended for human consumption to be used as feed material or for technical purposes, intended for dispatch to the European Community** Reference number (1) ORIGINAL
2. **Consignee** (name and address in full)	3. **Origin of the fish oil** 3.1. Country: ..... 3.2. Code of territory: .....
	4. **Competent authority** 4.1. Responsible Ministry: ..... 4.2. Certifying department: ....
5. **Intended destination of the fish oil** 5.1. EU Member State: ..... 5.2. Name and address of the destination:	6. **Place of loading for exportation** ....
7. **Means of transport and consignment identification** (2) 7.1. (Lorry, rail wagon, ship, or aircraft) (3) 7.2. Number of seal (if applicable): ..... 7.3. Registration number(s), ship name or flight number:	7.4. Nature of packaging: .... 7.5. Number of packages: ..... 7.6. Net weight: ..... 7.7. Lot/batch production reference number: ....
8. **Identification of the fish oil** 8.1. Description of the fish oil: ..... 8.2. Address and registration number of treatment/processing establishment (3):	
9. **Health attestation** I, the undersigned official veterinarian, declare that I have read and understood Regulation (EC) No 1774/2002 (4) and certify that the fish oil described above: 9.1. consists of fish oil that satisfy the health requirements below; 9.2. contains exclusively fish oil not intended for human consumption; 9.3. has been prepared and stored in a dedicated fish 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;	

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

(3) *either* [- former foodstuffs of fish origin, other than catering waste (5), 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;]

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

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

9.5. the fish oil:

(a) has been subjected to processing in accordance with Annex VII, Chapter IV of Regulation 1774/2002/EC, in order to kill pathogenic agents;

(b) has not been in contact with other types of oils including rendered fats from other animal species, and

(3) *either* [(c) is packaged in new containers or in containers that have been cleaned and all precautions taken to prevent their contamination;]

(3) *or* [(c) where bulk transport is intended, the pipe, pumps and bulk tanks and any other bulk container or bulk road tanker used in the transportation of the product from the manufacturing plant either directly on to the ship or into shore tanks or direct to plants have been inspected and found to be clean before use;]

and which bear labels indicating 'NOT FOR HUMAN CONSUMPTION'.

Official stamp and signature

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

(stamp) (6)

..... (signature of the official veterinarian) (6)

..... (name, qualifications and title, in capital letters)

Notes

(1) Issued by the competent authority.
(2) For goods vehicles the registration number should be given. For bulk containers, the container number and the seal number (if applicable) should be included.
(3) Delete as appropriate.
(4) OJ L 273, 10.10.2002, p. 1.
(5) Catering waste means all waste food, including used cooking oils, originating in restaurants, catering facilities and kitchens, including central kitchens and household kitchens.
(6) The signature and the stamp must be in a different colour to that of the printing.

CHAPTER 10 (A)

Health certificate

For rendered fats not intended for human consumption to be used as feed material or for technical purposes, intended for dispatch to the European Community

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

1. **Consignor** (name and address in full)	**VETERINARY CERTIFICATE** **For rendered fats not intended for human consumption to be used as feed material or for technical purposes, intended for dispatch to the European Community** Reference number (1) ORIGINAL
2. **Consignee** (name and address in full)	3. **Origin of the rendered fat** 3.1. Country: ..... 3.2. Code of territory: .....
	4. **Competent Authority** 4.1. Responsible Ministry: ..... 4.2. Certifying department: ....
5. **Intended destination of the rendered fat** 5.1. EU Member State: ..... 5.2. Name and address of the destination:	6. **Place of loading for exportation** ....
7. **Means of transport and consignment identification** (2) 7.1. (Lorry, rail wagon, ship, or aircraft) (3) 7.2. Number of seal (if applicable): ..... 7.3. Registration number(s), ship name or flight number:	7.4. Nature of packaging: .... 7.5. Number of packages: ..... 7.6. Net weight: ..... 7.7. Lot/batch production reference number: ....
8. **Identification of the rendered fat** 8.1. Description of the rendered fat: ..... 8.2. Rendered fat of: ..... *(animal species)* 8.3. Address and registration number of treatment/processing establishment (3):	
9. **Health attestation** I, the undersigned official veterinarian, declare that I have read and understood Regulation (EC) No 1774/2002 (4) and certify that the rendered fats described above: 9.1. consist of rendered fats that satisfy the health requirements below; 9.2. consist of rendered fats not intended for human consumption;	

9.3. have 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 or in accordance with Chapter II of Annex C to Council Directive 77/99/EEC ([5]) or Chapter IX of Annex 1 to Council Directive 92/118/EEC ([6]), in order to kill pathogenic agents;

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

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

([3]) *and/or* [- 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;]

([3]) *and/or* [- hides and skins, hooves and horns, pig bristles and feathers originating from animals that 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;]

([3]) *and/or* [- blood obtained from animals other than ruminants that are 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;]

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

([3]) *and/or* [- former foodstuffs of animal origin, or former foodstuffs containing products of animal origin, other than catering waste ([7]), 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;]

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

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

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

([3]) *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;]

9.5. if derived from ruminant animals were purified in such way that the maximum levels of remaining total insoluble impurities does not exceed 0,15 % in weight;

9.6. the rendered fats:

(a) have been subjected to processing in accordance with Annex VII, Chapter IV of Regulation 1774/2002/EC, or treatment in accordance with Council Directives 77/99/EEC or 92/118/EEC, in order to kill pathogenic agents, and

([3]) *either* [(b)are packaged in new containers or in containers that have been cleaned and all precautions taken to prevent their contamination;]

([3]) or [(b)where bulk transport is intended, the pipe, pumps and bulk tanks and any other bulk container or bulk road tanker used in the transportation of the product from the manufacturing plant either directly on to the ship or into shore tanks or direct to plants have been inspected and found to be clean before use;]

and which bear labels indicating 'NOT FOR HUMAN CONSUMPTION'.

Official stamp and signature

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

.....
(stamp) ([8]) (signature of the official veterinarian) ([8])

.....
(name, qualifications and title, in capital letters)

(1) Issued by the competent authority.
(2) For goods vehicles the registration number should be given. For bulk containers, the container number and the seal number (if applicable) should be included.
(3) Delete as appropriate.
(4) OJ L 273, 10.10.2002, p. 1.
(5) OJ L 26, 31.1.1977, p. 85.
(6) DO L 62 de 15.3.1993, p. 49.
(7) Catering waste means all waste food, including used cooking oils, originating in restaurants, catering facilities and kitchens, including central kitchens and household kitchens.
(8) The signature and the stamp must be in a different colour to that of the printing.

CHAPTER 10 (B)

Health certificate

For rendered fats not intended for human consumption to be used for technical purposes, intended for dispatch to the European Community

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

1. **Consignor** (name and address in full)	**VETERINARY CERTIFICATE** **For rendered fats not intended for human consumption to be used for technical purposes, intended for dispatch to the European Community** Reference number (1) ORIGINAL
2. **Consignee** (name and address in full)	3. **Origin of the rendered fat** 3.1. Country: ..... 3.2. Code of territory: ..... 4. **Competent authority** 4.1. Responsible Ministry: ..... 4.2. Certifying department: ....
5. **Intended destination of the rendered fat** 5.1. EU Member State: ..... 5.2. Name and address of the destination:	6. **Place of loading for exportation** ....
7. **Means of transport and consignment identification** (2) 7.1. (Lorry, rail wagon, ship, or aircraft) (3) 7.2. Number of seal (if applicable): ..... 7.3. Registration number(s), ship name or flight number:	7.4. Nature of packaging: .... 7.5. Number of packages: ..... 7.6. Net weight: ..... 7.7. Lot/batch production reference number: ....
8. **Identification of the rendered fat** 8.1. Description of the rendered fat: ..... 8.2. Rendered fat of: ..... *(animal species)* 8.3. Address and registration number of treatment/processing establishment (3):	
9. **Health attestation** I, the undersigned official veterinarian, declare that I have read and understood Regulation (EC) No 1774/2002 (4) and certify that the rendered fats described above: 9.1. consist of rendered fats that satisfy the health requirements below; 9.2. consist of rendered fats not intended for human or animal consumption;	

9.3. have been prepared and stored in a plant approved, validated and supervised by the competent authority in accordance with Article 13 and where appropriate Article 11 of Regulation (EC) No 1774/2002, in order to kill pathogenic agents;

9.4. have been prepared with the following animal by-products:

(3) *either* [category 2 materials (5);]

(3) *or* [a mixture of category 2 materials with category 3 materials (6);]

9.5. if derived from ruminant animals were purified in such way that the maximum levels of remaining total insoluble impurities does not exceed 0,15 % in weight;

9.6. the rendered fats:

(a) have been subjected to processing in accordance with Annex VII, Chapter XII of Regulation (EC) No 1774/2002/EC , in order to kill pathogenic agents; and

(3) *either* [(b)are packaged in new containers or in containers that have been cleaned and all precautions taken to prevent their contamination;]

(3) *or* [(b)where bulk transport is intended, the pipe, pumps and bulk tanks and any other bulk container or bulk road tanker used in the transportation of the product from the manufacturing plant either directly on to the ship or into shore tanks or direct to plants have been inspected and found to be clean before use;]

and which bear labels indicating "NOT FOR HUMAN OR ANIMAL CONSUMPTION".

Official stamp and signature

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

(stamp) (7)

..... (signature of the official veterinarian) (7)

..... (name, qualifications and title, in capital letters)

Notes

(1) Issued by the competent authority.
(2) For goods vehicles the registration number should be given. For bulk containers, the container number and the seal number (if applicable) should be included.
(3) Delete as appropriate.
(4) OJ L 273, 10.10.2002, p. 1.
(5) List of category 2 materials:
(a) all animal materials collected when treating waste water from slaughterhouses other than slaughterhouses covered by Article 4(1)(d) or from category 2 processing plants, including screenings, materials from desanding, grease and oil mixtures, sludge and materials removed from drains from those premises;
(b) products of animal origin containing residues of veterinary drugs and contaminants listed in group B(1) and (2) of Annex I to Directive 96/23/EC, if such residues exceed the permitted level laid down by Community legislation;
(c) products of animal origin, other than category 1 material, that are imported from third countries and, in the course of the inspections provided for in Community legislation, fail to comply with the veterinary requirements for their importation into the Community, unless they are returned or their importation is accepted under restrictions laid down under Community legislation;
(d) animals and parts of animals, other than those referred to in Article 4, that die other than by being slaughtered for human consumption, including animals killed to eradicate an epizootic disease;
(e) mixtures of category 2 material with category 3 material, including any material destined for processing in a category 2 processing plant; and
(f) animal by-products other than category 1 material or category 3 material.
(6) List of category 3 materials:
(a) parts of slaughtered animals, which are fit for human consumption in accordance with Community legislation, but are not intended for human consumption for commercial reasons;
(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;
(c) hides and skins, hooves and horns, pig bristles and feathers originating from animals that 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;
d) blood obtained from animals other than ruminants that are 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;
(e) animal by-products derived from the production of products intended for human consumption, including degreased bones and greaves;
(f) former foodstuffs of animal origin, or former foodstuffs containing products of animal origin, other than catering waste (1), 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;
(g) milk originating from animals which do not show any clinical signs of any disease communicable through that product to humans or animals;
(h) fish or other sea animals, except sea mammals, caught in the open sea for the purposes of fishmeal production;
(i) by-products from fish from plants manufacturing fish products for human consumption;
(j) 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.
(7) The signature and the stamp must be in a different colour to that of the printing.

CHAPTER 11

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 the European Community

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

1. **Consignor** (name and address in full)	**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 the European Community** Reference number (1) ORIGINAL
2. **Consignee** (name and address in full)	3. **Origin of the gelatine/collagen** (2) 3.1. Country: ..... 3.2. Code of territory: .....
	4. **Competent authority** 4.1. Responsible Ministry: ..... 4.2. Certifying department: ....
5. **Intended destination of the gelatine/collagen** (2) 5.1. EU Member State: ..... 5.2. Name and address of the destination:	6. **Place of loading for exportation** ....
7. **Means of transport and consignment identification** (3) 7.1. (Lorry, rail wagon, ship, or aircraft) (2) 7.2. Number of seal (if applicable): ..... 7.3. Registration number(s), ship name or flight number:	7.4. Nature of packaging: .... 7.5. Number of packages: ..... 7.6. Net weight: ..... 7.7. Lot/batch production reference number: ....
8. **Identification of the gelatine/collagen** (2) 8.1. Description of the gelatine/collagen (2): ..... 8.2. Gelatine/collagen (2) of: ..... *(animal species)* 8.3. Address and registration number of treatment/processing establishment (2):	
9. **Health attestation** I, the undersigned official veterinarian, declare that I have read and understood Regulation (EC) No 1774/2002 (4) and certify that the gelatine/collagen (2) described above: 9.1. consists of gelatine/collagen (2) that satisfy the health requirements below; 9.2. consist exclusively of gelatine/collagen (2) not intended for human consumption;	

9.3. 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 , in order to kill pathogenic agents;

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

(2) *either* [- parts of slaughtered animals, which are 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 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;]

(2) *and/or* [- hides and skins, hooves and horns, pig bristles and feathers originating from animals that 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;]

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

(2) *and/or* [- former foodstuffs of animal origin, or former foodstuffs containing products of animal origin, other than catering waste (5), 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;]

9.5. the gelatine/collagen (2):

(a) was wrapped, packaged, stored and transported under satisfactory hygiene conditions, and in particular wrapping and packaging took place in a dedicated room, and only preservatives permitted under Community legislation were used;

Wrappings and packages containing gelatine/collagen (2) must carry the words 'GELATINE/COLLAGEN (2) SUITABLE FOR ANIMAL CONSUMPTION', and

(2) *either* [(b)in the case of gelatine, has been produced by a process that ensuring that unprocessed Category 3 material is subjected to a treatment with acid or alkali, followed by one or more rinses, involving pH adjustment, extraction by heating one or several times in succession, followed by purification by means of filtration and sterilisation, in order to kill pathogenic agents;]

(2) *or* [(b)in the case of collagen, has been produced by a process that ensuring that unprocessed category 3 material is subjected to a treatment involving washing, pH adjustment using acid or alkali followed by one or more rinses, filtration and extrusion, in order to kill pathogenic agents.]

Official stamp and signature

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

(stamp) (6)

..... (signature of the official veterinarian) (6)

..... (name, qualifications and title, in capital letters)

Notes

(1) Issued by the competent authority.
(2) Delete as appropriate.
(3) For goods vehicles the registration number should be given. For bulk containers, the container number and the seal number (if applicable) should be included.
(4) OJ L 273, 10.10.2002, p. 1.
(5) Catering waste means all waste food, including used cooking oils, originating in restaurants, catering facilities and kitchens, including central kitchens and household kitchens.
(6) The signature and the stamp must be in a different colour to that of the printing.

CHAPTER 12

Health certificate

For hydrolysed protein, dicalcium phosphate and tricalcium phosphate not intended for human consumption to be used as feed material or for technical purposes, intended for dispatch to the European Community

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

1. **Consignor** (name and address in full)	**VETERINARY CERTIFICATE** **For hydrolysed protein, dicalcium phosphate and tricalcium phosphate not intended for human consumption to be used as feed material or for technical purposes, intended for dispatch to the European Community** Reference number (1) ORIGINAL
2. **Consignee** (name and address in full)	3. **Origin of the hydrolysed protein/dicalcium phosphate/tricalcium phosphate** (2) 3.1. Country: ..... 3.2. Code of territory: ..... 4. **Competent authority** 4.1. Responsible Ministry: ..... 4.2. Certifying department: ....
5. **Intended destination of the hydrolysed protein/dicalcium phosphate/tricalcium phosphate** (2) 5.1. EU Member State: ..... 5.2. Name and address of the destination:	6. **Place of loading for exportation** ....
7. **Means of transport and consignment identification** (3) 7.1. (Lorry, rail wagon, ship, or aircraft) (2) 7.2. Number of seal (if applicable): ..... 7.3. Registration number(s), ship name or flight number:	7.4. Nature of packaging: .... 7.5. Number of packages: ..... 7.6. Net weight: ..... 7.7. Lot/batch production reference number: ....
8. **Identification of the hydrolysed protein/dicalcium phosphate/tricalcium phosphate** (2) 8.1. Description of the [hydrolysed protein]/[dicalcium phosphate]/[tricalcium phosphate] (2): .... 8.2. [hydrolysed protein]/[dicalcium phosphate]/[tricalcium phosphate] (2) of: .... *(animal species)* 8.3. Address and registration number of treatment/processing establishment (2):	
9. **Health attestation** I, the undersigned official veterinarian, declare that I have read and understood Regulation (EC) No 1774/2002 (4) and certify that the hydrolysed protein/dicalcium phosphate/tricalcium phosphate (2) described above:	

9.1. consists of hydrolysed protein/dicalcium phosphate/tricalcium phosphate (2) that satisfy the health requirements below;

9.2. consists exclusively of hydrolysed protein/dicalcium phosphate/tricalcium phosphate (2) not intended for human consumption;

9.3. 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 , in order to kill pathogenic agents;

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

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

(3) *and/or* [- 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 carcases that are fit for human consumption in accordance with Community legislation;]

(3) *and/or* [- hides and skins, hooves and horns, pig bristles and feathers originating from animals that 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;]

(3) *and/or* [- blood obtained from animals other than ruminants that are 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;]

(3) *and/or* [- animal by-products derived from the production of products intended for human consumption;]

(3) *and/or* [- former foodstuffs of animal origin, or former foodstuffs containing products of animal origin, other than catering waste (5), 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;]

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

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

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

(3) *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;]

9.5. the hydrolysed protein/dicalcium phosphate/tricalcium phosphate (2):

(a) was wrapped and packaged in packaging which bear labels indicating 'NOT FOR HUMAN CONSUMPTION' and stored and transported under satisfactory hygiene conditions, and in particular wrapping and packaging took place in a dedicated room, and only preservatives permitted under Community legislation were used, and

(2) *bien* [(b)in the case of hydrolysed protein, has been produced by a process involving appropriate measures to minimise contamination of raw category 3 material. In the case of hydrolysed proteins entirely or partly derived from ruminants hides and skins, has been produced in a processing plant dedicated only to hydrolysed proteins production, using a process involving the preparation of the raw category 3 material by brining, liming and intensive washing followed by:

(b) (i) exposure of the material to a pH of more than 11 for more than three hours at temperature of more than 80 C and subsequently by heat treatment at more than 140 °C for 30 minutes at more than 3,6 bar; and

(b) (ii) exposure of the material to a pH of 1 to 2, followed by a pH of more than 11, followed by heat treatment at 140 °C for 30 minutes at 3 bar;]

(2) *or* [(b)in the case of dicalcium phosphate, has been produced by a process that :

(b) (i) ensures that all category 3 bone-material is finely crushed and degreased with hot water and treated with dilute hydrochloric acid (at a minimum concentration of 4 % and a pH of less than 1,5) over a period of at least two days;

(b) (ii) followed by treatment of the obtained phosphoric liquor with lime, resulting in a precipitate of dicalcium phosphate at pH 4 to 7; and

(b) (iii) finally air-dries this precipitate for 15 minutes, with inlet temperature of 270 to 325 °C and end temperature between 60 and 65 °C;]

(2) *or* [(b)in the case of tricalcium phosphate, has been produced by a process ensuring:

(i) that all category 3 bone-material is finely crushed and degreased in counter-flow with hot water (bone chips less than 14 mm);

(ii) continuous cooking with steam at 145 °C during 30 minutes at 4 bars;

(iii) separation of the protein broth from the hydroxyapatite (tricalcium phosphate) by centrifugation; and

(iv) granulation of the tricalcium phosphate after drying in a fluid bed with air at 200 °C.]

Official stamp and signature

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

.....
(stamp) (6) (signature of the official veterinarian) (6)

.....
(name, qualifications and title, in capital letters)

Notes

(1) Issued by the competent authority.
(2) Delete as appropriate.
(3) For goods vehicles the registration number should be given. For bulk containers, the container number and the seal number (if applicable) should be included.
(4) OJ L 273, 10.10.2002, p. 1.
(5) Catering waste means all waste food, including used cooking oils, originating in restaurants, catering facilities and kitchens, including central kitchens and household kitchens.
(6) The signature and the stamp must be in a different colour to that of the printing.

CHAPTER 13

Health certificate

For apiculture products, intended for dispatch to the European Community

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

1. **Consignor** (name and address in full)	**VETERINARY CERTIFICATE** **For apiculture products, intended for dispatch to the European Community** Reference number (1) ORIGINAL
2. **Consignee** (name and address in full)	3. **Origin of the apiculture products** 3.1. Country: ..... 3.2. Code of territory: .....
	4. **Competent Authority** 4.1. Responsible Ministry: ..... 4.2. Certifying department: ....
5. **Destination of the apiculture products** 5.1. EU Member State: ..... 5.2. Name and address of the destination:	6. **Place of loading for exportation** ....
7. **Means of transport and consignment identification** (2) 7.1. (Lorry, rail wagon, ship, or aircraft) (3) 7.2. Number of seal (if applicable): ..... 7.3. Registration number(s), ship name or flight number:	7.4. Nature of packaging: .... 7.5. Number of packages: ..... 7.6. Net weight: ..... 7.7. Lot/batch production reference number: ....
8. **Identification of the apiculture products** 8.1. Description of the apiculture products: .... 8.2. Address and registration number of the establishment of production:	
9. **Health attestation** I, the undersigned official veterinarian, declare that I have read and understood Regulation (EC) No 1774/2002 (4) and certify that the apiculture products described above: 9.1. consist of apiculture products that satisfy the health requirements below; 9.2. (3) *either* [are new and have not been in use before and have not come into contact with bees or used apiculture products;]	

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

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

9.3. 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.

Official stamp and signature

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

.....
(stamp) (5) (signature of the official veterinarian) (5)

.....
(name, qualifications and title, in capital letters)

Notes

(1) Issued by the competent authority.
(2) For goods vehicles the registration number should be given. For bulk containers, the container number and the seal number (if applicable) should be included
(3) Delete as appropriate.
(4) OJ L 273, 10.10.2002, p. 1.
(5) The signature and the stamp must be in a different colour to that of the printing.

CHAPTER 14 (A)

Health certificate

For fat derivatives not intended for human consumption to be used for technical purposes, intended for dispatch to the European Community

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

1. **Consignor** (name and address in full)	**VETERINARY CERTIFICATE** **For fat derivatives not intended for human consumption to be used for technical purposes, intended for dispatch to the European Community** Reference number (1) ORIGINAL
2. **Consignee** (name and address in full)	3. **Origin of the fat derivatives** 3.1. Country: ..... 3.2. Code of territory: .....
	4. **Competent Authority** 4.1. Responsible Ministry: ..... 4.2. Certifying department: ....
5. **Intended destination of the fat derivatives** 5.1. EU Member State: ..... 5.2. Name and address of the destination:	6. **Place of loading for exportation** ....
7. **Means of transport and consignment identification** (2) 7.1. (Lorry, rail wagon, ship, or aircraft) (3) 7.2. Number of seal (if applicable): ..... 7.3. Registration number(s), ship name or flight number:	7.4. Nature of packaging: .... 7.5. Number of packages: ..... 7.6. Net weight: ..... 7.7. Lot/batch production reference number: ....
8. **Identification of the fat derivatives** 8.1. Description of the fat derivatives: ..... 8.2. Fat derivatives of: ..... *(animal species)* 8.3. Address and registration number of treatment/processing establishment (3):	
9. **Health attestation** I, the undersigned official veterinarian, declare that I have read and understood Regulation (EC) No 1774/2002 (4) and certify that the fat derivatives described above: 9.1. consist of fat derivatives that satisfy the health requirements below; 9.2. consist of fat derivatives containing exclusively fat derivatives not intended for human nor animal consumption;	

9.3. have been prepared and stored in a plant approved, validated and supervised by the competent authority in accordance with Article 14 and where appropriate Article 11 of Regulation (EC) No 1774/2002, in order to kill pathogenic agents;

9.4. have been prepared from rendered fats exclusively produced from category 2 and/or category 3 materials (5);

9.5. the fat derivatives produced from category 2 materials:

(a) have been produced using the following methods:

(3) *either* [transesterification or hydrolysis at least 200 °C, under corresponding appropriate pressure, for 20 minutes (glycerol, fatty acids and esters); and]

(3) *or* [saponification with NaOH 12M (glycerol and soap):

(3) *either* [in a batch process at 95 °C for three hours; and]

(3) *or* [in a continuous process at 140 °C, 2 bars (2 000 hPa) for eight minutes; and]]

(b) are packaged in new containers or in containers that have been cleaned, and all precautions are taken to prevent its contamination which bear labels indicating 'NOT FOR HUMAN OR ANIMAL COSUMPTION'.

Official stamp and signature

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

(stamp) (6)

..... (signature of the official veterinarian) (6)

..... (name, qualifications and title, in capital letters)

Notes

(1) Issued by the competent authority.
(2) For goods vehicles the registration number should be given. For bulk containers, the container number and the seal number (if applicable) should be included.
(3) Delete as appropriate.
(4) OJ L 273, 10.10.2002, p. 1.
(5) List of category 2 materials:
a) all animal materials collected when treating waste water from slaughterhouses other than slaughterhouses covered by Article 4(1)(d) or from category 2 processing plants, including screenings, materials from desanding, grease and oil mixtures, sludge and materials removed from drains from those premises;
b) products of animal origin containing residues of veterinary drugs and contaminants listed in group B(1) and (2) of Annex I to Directive 96/23/EC, if such residues exceed the permitted level laid down by Community legislation;
c) products of animal origin, other than category 1 material, that are imported from third countries and, in the course of the inspections provided for in Community legislation, fail to comply with the veterinary requirements for their importation into the Community, unless they are returned or their importation is accepted under restrictions laid down under Community legislation;
d) animals and parts of animals, other than those referred to in Article 4, that die other than by being slaughtered for human consumption, including animals killed to eradicate an epizootic disease;
e) mixtures of category 2 material with category 3 material, including any material destined for processing in a category 2 processing plant; and
f) animal by-products other than category 1 material or category 3 material.
(6) The signature and the stamp must be in a different colour to that of the printing.

CHAPTER 14 (B)

Health certificate

For fat derivatives not intended for human consumption to be used as feed or for technical purposes, intended for dispatch to the European Community

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

1. **Consignor** (name and address in full)	**VETERINARY CERTIFICATE** **For fat derivatives not intended for human consumption to be used as feed, intended for dispatch to the European Community** Reference number (1) ORIGINAL
2. **Consignee** (name and address in full)	3. **Origin of the fat derivatives** 3.1. Country: ..... 3.2. Code of territory: .....
	4. **Competent Authority** 4.1. Responsible Ministry: ..... 4.2. Certifying department: ....
5. **Intended destination of the fat derivatives** 5.1. EU Member State: ..... 5.2. Name and address of the destination:	6. **Place of loading for exportation** ....
7. **Means of transport and consignment identification** (2) 7.1. (Lorry, rail wagon, ship, or aircraft) (3) 7.2. Number of seal (if applicable): ..... 7.3. Registration number(s), ship name or flight number:	7.4. Nature of packaging: .... 7.5. Number of packages: ..... 7.6. Net weight: ..... 7.7. Lot/batch production reference number: ....
8. **Identification of the fat derivatives** 8.1. Description of the fat derivatives: ..... 8.2. Fat derivatives of: ..... *(animal species)* 8.3. Address and registration number of treatment/processing establishment (3):	
9. **Health attestation** I, the undersigned official veterinarian, declare that I have read and understood Regulation (EC) No 1774/2002 (4) and certify that the fat derivatives described above: 9.1. consist of fat derivatives that satisfy the health requirements below; 9.2. consist of fat derivatives containing exclusively fat derivatives not intended for human consumption;	

9.3. have been prepared and stored in a plant approved, validated and supervised by the competent authority in accordance with Article 14 and where appropriate Article 11 of Regulation (EC) No 1774/2002, in order to kill pathogenic agents;

9.4. have been prepared from rendered fats exclusively produced from the following category 3 materials:

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

(3) *and/or* [- 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;]

(3) *and/or* [- hides and skins, hooves and horns, pig bristles and feathers originating from animals that 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;]

(3) *and/or* [- blood obtained from animals other than ruminants that are 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;]

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

(3) *and/or* [- former foodstuffs of animal origin, or former foodstuffs containing products of animal origin, other than catering waste (5), 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;]

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

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

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

(3) *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;]

9.5. are packaged in new containers or in containers which bear labels indicating 'NOT FOR HUMAN CONSUMPTION', that have been cleaned, and all precautions are taken to prevent its contamination.

Official stamp and signature

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

(stamp) (6)

..... (signature of the official veterinarian) (6)

..... (name, qualifications and title, in capital letters)

Notes

(1) Issued by the competent authority.
(2) For goods vehicles the registration number should be given. For bulk containers, the container number and the seal number (if applicable) should be included.
(3) Delete as appropriate.
(4) OJ L 273, 10.10.2002, p. 1.
(5) Catering waste means all waste food, including used cooking oils, originating in restaurants, catering facilities and kitchens, including central kitchens and household kitchens.
(6) The signature and the stamp must be in a different colour to that of the printing.

CHAPTER 15

Health certificate

For egg products not intended for human consumption that could be used as feed material, intended for dispatch to the European Community

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

1. **Consignor** (name and address in full)	**VETERINARY CERTIFICATE** **For egg products not intended for human consumption that could be used as feed material, intended for dispatch to the European Community** Reference number (1) ORIGINAL
2. **Consignee** (name and address in full)	3. **Origin of the egg products** 3.1. Country: ..... 3.2. Code of territory: .....
	4. **Competent Authority** 4.1. Responsible Ministry: ..... 4.2. Certifying department: ....
5. **Destination of the egg products** 5.1. EU Member State: ..... 5.2. Name and address of the destination:	6. **Place of loading for exportation** ....
7. **Means of transport and consignment identification** (2) 7.1. (Lorry, railwagon, ship, or aircraft) (3) 7.2. Number of seal (if applicable): ..... 7.3. Registration number(s), ship name or flight number:	7.4. Nature of packaging: .... 7.5. Number of packages: ..... 7.6. Net weight: ..... 7.7. Lot/batch production reference number: ....
8. **Identification of the egg products** 8.1. Nature of the egg products: ..... 8.2. Species of animals from which the egg products derive: .... 8.3. Address and registration number of the approved establishment:	
9. **Health attestation** I, the undersigned official veterinarian, declare that I have read and understood Regulation (EC) No 1774/2002 (4) and certify that the egg products described above: 9.1. consist of egg products that satisfy the health requirements below; 9.2. consist exclusively of egg products not intended for human consumption;	

9.3. have 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 or Council Directive 89/437/EEC (5), in order to kill pathogenic agents;

9.4. se have been prepared (derived) exclusively with the following animal by-product:

— eggs originating from animals which did not show clinical signs of any disease communicable through that product to humans or animals;

9.5. have been subjected to processing:

(3) *either* [in accordance with processing method ..... (6) as set out in Annex V, Chapter III of Regulation (EC) No 1774/2002/EC;]

(3) *or* [in accordance to a method and parameters which ensure that the products comply with the microbiological standards set in Chapter I, paragraph 10 of Annex VII to Regulation (EC) No 1774/2002/EC;]

(3) *or* [treated in accordance with Chapter V of the Annex to Council Directive 89/437/EC;]

9.6. have been examined by the competent authority taking a random sample immediately prior to dispatch and found it to comply with the following standards (7):

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

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

9.7. meet Community standards on residues of substances that are harmful or might alter the organoleptic characteristics of the product or make its use as feed dangerous or harmful to animal health;

9.8. the end product was:

(3) *either* [packed in new or sterilised bags;]

(3) *either* [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';

9.9. the end product was stored in enclosed storage;

9.10. the product has undergone all precautions to avoid contamination with pathogenic agents after treatment:

Official stamp and signature

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

(stamp) (8)

..... (signature of the official veterinarian) (8)

..... (name, qualifications and title, in capital letters)

Notes

(1) Issued by the competent authority.
(2) For goods vehicles the registration number should be given. For bulk containers, the container number and the seal number (if applicable) should be included).
(3) Delete as appropriate.
(4) OJ L 273, 10.10.2002, p. 1.
(5) OJ L 212, 22.07.1989, p. 89.
(6) Insert method 1 to 5 or 7 as applicable:.
(7) 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.
(8) The signature and the stamp must be in a different colour to that of the printing.

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 European 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 (1):

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

(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 fertilisers or soil improvers and will be conveyed directly to the following processing establishment:

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

The importer

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

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

Signature .....

Reference number as indicated on the certificate provided for in Annex B to Commission Decision 93/13/EEC: .....

.....

Official stamp of the border inspection post of entry into the EC (2)

Signature .....
(Signature of the official veterinarian of the border inspection post) (2)

Name: .....
(Name in capital letters)

(1) Delete as appropriate.
(2) The signature and the stamp must be in a different colour to that of the printing.

5. Annex XI is replaced by the following:

ANNEX XI

Lists of third countries from which Member States may authorise imports of animal by-products not intended for human consumption

The inclusion of a country on one of the following lists is a necessary, but not sufficient, condition for the importation of relevant products from that country. Imports must also fulfil the relevant animal health and public health requirements.

PART I

List of third countries from which Member States may authorise imports of milk and milk-based products (health certificates Chapters 2(A), 2(B) and 2(C))

Third countries listed in column B or column C of the Annex to Commission Decision 95/340/EC[(6)].

PART II

List of third countries from which Member States may authorise imports of processed animal proteins (excluding fishmeal) (health certificate Chapter 1)

Third countries listed in part 1 of Annex II to Council Decision 79/542/EEC[(7)].

PART III

List of third countries from which Member States may authorise imports of fishmeal and fish oil (health certificate Chapters 1 and 9)

Third countries listed in the Annex to Commission Decision 97/296/EC[(8)].

PART IV

List of third countries from which Member States may authorise imports of rendered fats (excluding fish oil) (health certificate Chapters 10(A) and 10(B))

Third countries listed in part 1 of Annex II to Council Decision 79/542/EEC.

PART V

List of third countries from which Member States may authorise imports of blood products for feed material (health certificate Chapter 4(B))

A. Blood products from ungulates

Third countries or parts of countries listed in part 1 of Annex II to Decision 79/542/EEC, from which imports of all categories of fresh meat of the respective species are authorised.

B. Blood products from other species

Third countries listed in part 1 of Annex II to Council Decision 79/542/EEC.

PART VI

List of third countries from which Member States may authorise imports of raw material including blood products (with the exception of equidae) intended for technical purposes including pharmaceutical products (health certificate Chapters 4(C) and 8(B))

A. Blood products

1. Blood products from ungulates:

third countries or parts of third countries listed in part 1 of Annex II of Decision 79/542/EEC, from which imports of all categories of fresh meat of the respective species are authorised.

2. Blood products of other species:

third countries listed in part 1 of Annex II of Decision 79/542/EEC.

B. Raw material (except blood products) for pharmaceutical use

Third countries listed in part 1 of Annex II of Decision 79/542/EEC, in the Annex to Commission Decision 94/85/EEC[9] or in Annex I to Commission Decision 2000/585/EC[10] and the following countries:

— (JP) Japan,
— (PH) Philippines and
— (TW) Taiwan.

C. Raw material for technical purposes other than pharmaceutical uses

Third countries listed in part 1 of Annex II of Decision 79/542/EEC from which imports of that category of fresh meat of the respective species is authorised, in the Annex to Decision 94/85/EEC, or in the Annex to Decision 2000/585/EC.

PART VII(A)

List of third countries from which Member States may authorise imports of animal by-products for the manufacture of processed petfood (health certificate Chapter 3(B) and 8(A))

A. Animal by-products from bovine, ovine, caprine, porcine and equine animals, including farmed and wild animals

third countries or parts of third countries listed in part 1 of Annex II to Decision 79/542/EEC, from which imports of that category of fresh meat of the respective species is authorised and the following countries for the by-products specified:

— animal by-products from Bulgaria (BG), Latvia (LV), Romania (RO), [(Slovenia (SI)], concerning material from pigs;
— southern American and southern African countries or parts thereof where matured and boned meat of the corresponding species

is authorised, concerning matured and boned meat (including diaphragm) and/or matured trimmed offal of bovine, caprine, ovine animals and game (wild or farmed).

B. Raw material from poultry including ratites

Third countries or parts of third countries from which Member States authorise imports of fresh poultrymeat, which are listed in Annex I to Commission Decision 94/984/EC[(11)] and/or in Annex I to Commission Decision 2000/609/EC[(12)].

C. Raw material from fish

Third countries listed in the Annex to Decision 97/296/EC.

D. Raw material from other species, including feathered game, other wild land mammals and leparopidae

Third countries listed in Part 1 of Annex II to Decision 79/542/EEC or in the Annex I to Decision 2000/585/EC, from which Member States authorise imports of fresh meat from the same species.

PART VII(B)

List of third countries from which Member States may authorise imports of raw petfood intended for dispatch to the European Community for direct sale or animal by-products to be fed to farmed fur animals (health certificate Chapter 3(D))

Third countries listed in part 1 of Annex II to Decision 79/542/EEC, in Annex I to Decision 94/984/EC, or in Annex I to Decision 2000/609/EC, from which Member States authorise imports of fresh meat from the same species and where only bone in meat is authorised.

In the case of fish materials, third countries listed in the Annex to Decision 97/296/EC.

PART VII(C)

List of third countries from which Member States may authorise imports of flavouring innards for use in the manufacture of petfood, intended for dispatch to the European Community (health certificate Chapter 3(E))

Third countries listed in part 1 of Annex II to Decision 79/542/EEC, in Annex I to Decision 94/984/EC, or in Annex I to Decision 2000/609/EC, from which Member States authorise imports of fresh meat from the same species and where only bone in meat is authorised.

In the case of flavouring innards fish materials, third countries listed in the Annex to Commission Decision 97/296/EC.

PART VIII

List of third countries from which Member States may authorise imports of pig bristles (health certificate Chapter 7(A) and 7(B))

A. For untreated pig bristles, third countries listed in part 1 of Annex II to Decision 79/542/EEC, which are free of African swine fever for the last 12 months.

B. For treated pig bristles, third countries listed in part 1 of Annex II to Decision 79/542/EEC, which may not be free of African swine fever for the last 12 months.

PART IX

List of third countries from which Member States may authorise imports of manure for treatment of the soil

A. Processed manure products
Third countries listed in part 1 of Annex II to Decision 79/542/EEC.

B. Processed manure from equidae
Third countries listed in Part 1 of Annex II to Decision 79/542/EEC for live equidae.

C. Unprocessed manure from poultry
Third countries listed in Annex I to Decision 94/984/EC.

PART X

List of third countries from which Member States may authorise imports of petfood and dogchews (health certificate Chapters 3(A), 3(B) and 3(C))

Third countries listed in part 1 of Annex II to Decision 79/542/EEC, and the following countries:

(LK) Sri Lanka[(13)]

(JP) Japan[(14)]

(TW) Taiwan[(14)].

PART XI

List of third countries from which Member States may authorise imports of gelatine, hydrolysed protein, collagen, dicalcium phosphate and tricalcium phosphate (health certificate Chapters 11 and 12).

Third countries listed in part 1 of Annex II to Decision 79/542/EEC, and the following countries:

(KR) The Republic of Korea[(15)]

(MY) Malaysia[15]
(PK) Pakistan[15]
(TW) Taiwan[15].

PART XII

List of third countries from which Member States may authorise imports of apiculture products (health certificate Chapter 13)

Third countries listed in part 1 of Annex II to Decision 79/542/EEC.

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[16], from which the importation of horses for slaughter is allowed.

PART XIV

List of third countries from which Member States may authorise imports of hides and skins of ungulates (health certificate Chapters 5(A), 5(B) and 5(C))

A. For fresh or chilled hides and skins of ungulates, third countries listed in part 1 of Annex II to Decision 79/542/EEC, from which Member States authorise imports of fresh meat from the same species.

B. For treated hides and skins of ungulates, third countries or parts of third countries listed in Part 1 of Annex II to Decision 79/542/EEC.

C. For treated hides and skins of ruminants that are intended for dispatch to the European Community and which have been kept separate for 21 days or will undergo transport for 21 uninterrupted days before importation, any third country.

PART XV

List of third countries from which Member States may authorise imports of game trophies (health certificate Chapters 6(A) and 6(B))

A. For treated game trophies of birds and ungulates, being solely bones, horns, hooves, claws, antlers, teeth, hides or skins, any third country.

B. For game trophies of birds consisting of entire parts not having been treated, third countries listed in the Annex to Commission Decision 94/85/EC, from which Member States authorise imports of fresh poultrymeat, and the following countries:
(GL) Greenland
(TN) Tunisia.

C. For game trophies of ungulates consisting of entire parts not having been treated, third countries listed in the appropriate columns for fresh meat of ungulates in part 1 of Annex II to Decision 79/542/EEC, including any restrictions laid down in the column for special remarks for fresh meat.

PART XVI

List of third countries from which Member States may authorise imports of egg products not intended for human consumption that could be used as feed material (health certificate Chapter 15)

Third countries listed in part 1 of Annex II to Decision 79/542/EEC, and third countries or parts of third countries from which Member States authorise imports of fresh poultrymeat, which are listed in Annex I to Decision 94/984/EC and/or in Annex I to Decision 2000/609/EC.

PART XVII

List of third countries from which Member States may authorise imports of bones and bone products (excluding bonemeal), 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 (declaration Chapter 16)

Any third country.

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

(**2**) OJ L 117, 13.5.2003, p. 1.

(**3**) OJ L 212, 22.7.1989, p. 87.

(**4**) This includes countries with sero-positive ruminant animals.'

(**5**) Council Directive 92/65/EEC of 13 July 1992 laying down animal health requirements governing trade in and imports into the Community of animals, semen, ova and embryos not subject to animal health requirements laid down in specific Community rules referred to in Annex A(I) to Directive 90/425/EEC (OJ L 268, 14.9.1992, p. 54).

(**6**) OJ L 200, 24.8.1995, p. 38.

(**7**) OJ L 146, 14.6.1979, p. 15.

(**8**) OJ L 196, 24.7.1997, p. 82.

(**9**) OJ L 44, 17.2.1994, p. 31.

(**10**) OJ L 251, 6.10.2000, p. 1.

(**11**) OJ L 378, 31.12.1994, p. 11.

(**12**) OJ L 258, 12.10.2000, p. 49.

(**13**) Dogchews made from hides and skins of ungulates only.

(**14**) Processed petfood for ornamental fish only.

(**15**) Gelatine only.

(**16**) OJ L 73, 11.3.2004, p. 1.

Status:
Point in time view as at 22/04/2004.

Changes to legislation:
There are currently no known outstanding effects for the Commission Regulation (EC) No 668/2004 (repealed).