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Council of 3 October 2002 laying down health rules concerning animal by-products not intended for human consumption (repealed)

ANNEX II

HYGIENE REQUIREMENTS FOR THE COLLECTION AND TRANSPORT OF ANIMAL BY-PRODUCTS AND PROCESSED PRODUCTS

CHAPTER I

Identification

1. All necessary measures must be taken to ensure that:

(a) Category 1, Category 2 and Category 3 materials are identifiable and kept separate and identifiable during collection and transportation; and

(b) processed products are identifiable and kept separate and identifiable during transportation.

2. During transport, a label attached to the vehicle, container, carton or other packaging material must clearly indicate:

(a) the category of the animal by-products or, in the case of processed products, the category of animal by-products from which the processed products were derived; and

(b) (i) **[**F1in the case of Category 3 material, the words ‘not for human consumption’;

(ii) in the case of Category 2 material (other than manure and digestive tract content) and processed products derived therefrom, the words ‘not for animal consumption’; however, when Category 2 material is intended for the feeding of animals referred to in point (c) of Article 23(2) under the conditions provided for in that Article, the label shall instead indicate ‘for feeding to …’ completed with the name of the specific species of those animal(s) for the feeding of which the material is intended;

(iii) in the case of Category 1 material and processed products derived therefrom, the words ‘for disposal only’;

(iv) in the case of manure and digestive tract content, the word ‘manure’.**]**

Textual Amendments

F1 Substituted by Commission Regulation (EC) No 808/2003 of 12 May 2003 amending Regulation (EC) No 1774/2002 of the European Parliament and of the Council laying down health rules concerning animal by-products not intended for human consumption (Text with EEA relevance).

CHAPTER II

Vehicles and containers

1. Animal by-products and processed products must be collected and transported in sealed new packaging or covered leak-proof containers or vehicles.

2. Vehicles and reusable containers, and all reusable items of equipment or appliances that come into contact with animal by-products or processed products, must be:

(a) cleaned, washed and disinfected after each use;

(b) maintained in a clean condition; and

(c) clean and dry before use.

3. Reusable containers must be dedicated to the carriage of a particular product to the extent necessary to avoid cross-contamination.

[[F2]4. Packaging material must be incinerated or disposed of by some other means in accordance with instructions from the competent authority.**]**

Textual Amendments

F2 Inserted by Commission Regulation (EC) No 808/2003 of 12 May 2003 amending Regulation (EC) No 1774/2002 of the European Parliament and of the Council laying down health rules concerning animal by-products not intended for human consumption (Text with EEA relevance).

CHAPTER III

Commercial documents and health certificates

[[F1]1. During transportation, a commercial document or, when required by this Regulation, a health certificate must accompany animal by-products and processed products except in the case of processed products originating from Category 3 material which are supplied within the same Member State by retailers to final users other than business operators.**]**

2. Commercial documents must specify:

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

(b) the description of the material, including the information referred to in Chapter I, the animal species for Category 3 material and processed products derived therefrom destined for use as feed material and, if applicable, the ear-tag number;

(c) the quantity of the material;

(d) the place of origin of the material;

(e) the name and the address of the carrier;

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

(g) if appropriate:

(i) the approval number of the plant of origin, and

(ii) the nature and the methods of the treatment.

3. The commercial document must be produced at least in triplicate (one original and two copies). The original must accompany the consignment to its final destination.

The receiver must retain it. The producer must retain one of the copies and the carrier the other.

4. A model for the commercial document may be laid down under the procedure referred to in Article 33(2).

5. Health certificates must be issued and signed by the competent authority.

CHAPTER IV

Records

The records referred to in Article 9 must contain the information referred to in Chapter III, paragraph 2, as follows. They must contain:

(a) the information referred to in subparagraphs (b) and (c); and

(b) in the case of records kept by any person consigning animal by-products, the information referred to in subparagraphs (a), (e) and, if known, (f); or

(c) in the case of records kept by any person transporting animal by-products, the information referred to in subparagraphs (a), (d) and (f); or

(d) in the case of records kept by any person receiving animal by-products, the date of reception and the information referred to in subparagraphs (d) and (e).

CHAPTER V

Retention of documents

The commercial document and the health certificate referred to in Chapter III, and the records referred to in Chapter IV, must be kept for a period of at least two years for presentation to the competent authority.

CHAPTER VI

Temperature conditions

1. The transport of animal by-products must take place at an appropriate temperature, to avoid any risk to animal or public health.

2. Unprocessed Category 3 material destined for the production of feed material or pet food must be transported chilled or frozen, unless processed within 24 hours of departure.

3. The design of vehicles used for refrigerated transport must ensure the maintenance of an appropriate temperature throughout transport.

CHAPTER VII

Specific rules for transit

The carriage of animal by-products and processed products in transit must meet the requirements of Chapters I, II, III and VI.

CHAPTER VIII

Control measures

The competent authority must take the necessary measures to control the collection, transport, use and disposal of animal by-products and processed products, including by checking the keeping of required records and documents and, when this Regulation requires it or the competent authority considers it necessary, by sealing.

When the competent authority applies a seal to a consignment of animal by-products or processed products, it must inform the competent authority of the place of destination.

[F2CHAPTER IX

Collection of animal material when treating waste water

1. Category 1 processing plants and other premises where specified risk material is removed, slaughterhouses and Category 2 processing plants shall have a pre-treatment process for the retention and collection of animal material as an initial step in the treatment of waste water. The equipment used in the pre-treatment process shall consist of drain traps or screen with apertures or a mesh size of no more than 6 mm in the downstream end of the process or equivalent systems that ensures that the solid particles in the waste water passing through them are no more than 6 mm.

2. Waste water from the premises as referred to in paragraph 1 must enter a pre-treatment process which shall ensure that all waste water has been filtered through the process before being drained off the premises. No grinding or maceration shall take place which could facilitate the passage of animal material through the pre-treatment process.

3. All animal material retained in the pre-treatment process in premises as referred to in paragraph 1 shall be collected and transported as Category 1 or Category 2 material, as appropriate, and disposed of in accordance with this Regulation.

4. Waste water having passed the pre-treatment process in premises referred to in paragraph 1 and waste water from premises only receiving Category 3 material shall be treated in accordance with other relevant Community legislation.]

[F3CHAPTER X

Commercial document

1. The following commercial document shall accompany animal by-products and processed products during transportation. However, Member States may decide to

use a different commercial document for animal by-products and processed products transported within the same Member State.

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

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

Notes:

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

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

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

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

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

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

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

— the date on which the material was taken from the premises,

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

— the quantity of the material,

— the place of origin of the material,

— the name and the address of the carrier,

— the name and the address of the receiver and, if applicable, its approval number, and

— if appropriate, the approval number of the plant of origin, and the nature and the methods of the treatment.

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

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

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)
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2. **Consignee** (name and address in full)
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5. **Destination of the blood products**
5.1. EU Member State: .....
5.2. Name and address of the destination: .....
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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: .
.....
.....

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

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: .....
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6. **Place of loading for exportation**
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7.4. Nature of packaging: .....
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7.5. Number of packages: .....
7.6. Net weight: .....
7.7. Lot/batch production reference number:
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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;

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

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6. **Place of loading for exportation**
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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: .
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7.5. Number of packages: .....
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VETERINARY CERTIFICATE
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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;

Textual Amendments

F3 Inserted by Commission Regulation (EC) No 93/2005 of 19 January 2005 amending Regulation (EC) No 1774/2002 of the European Parliament and of the Council as regards processing of animal by-products of fish origin and commercial documents for the transportation of animal by-products (Text with EEA relevance).

Status:
Point in time view as at 01/01/2005.

Changes to legislation:
There are currently no known outstanding effects for the Regulation (EC) No 1774/2002 of the European Parliament and of the Council (repealed), ANNEX II.