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## ▶<u>C1</u> COMMISSION REGULATION (EU) No 206/2010

#### of 12 March 2010

laying down lists of third countries, territories or parts thereof authorised for the introduction into the European Union of certain animals and fresh meat and the veterinary certification requirements

(Text with EEA relevance) ◀

(OJ L 73, 20.3.2010, p. 1)

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► <u>M2</u>	Commission Regulation (EU) No 144/2011 of 17 February 2011	L 44	7	18.2.2011
► <u>M3</u>	Commission Implementing Regulation (EU) No 342/2011 of 8 April 2011	L 96	10	9.4.2011
► <u>M4</u>	Commission Implementing Regulation (EU) No 801/2011 of 9 August 2011	L 205	27	10.8.2011
► <u>M5</u>	Commission Implementing Regulation (EU) No 1112/2011 of 3 November 2011	L 287	32	4.11.2011
► <u>M6</u>	Commission Implementing Regulation (EU) No 497/2012 of 7 June 2012	L 152	1	13.6.2012
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- ▶<u>C1</u> Corrigendum, OJ L 146, 11.6.2010, p. 1 (206/2010)
- ► <u>C2</u> Corrigendum, OJ L 49, 24.2.2011, p. 53 (144/2011)
- ► <u>C3</u> Corrigendum, OJ L 63, 10.3.2011, p. 28 (144/2011)

#### **COMMISSION REGULATION (EU) No 206/2010**

#### of 12 March 2010

laying down lists of third countries, territories or parts thereof authorised for the introduction into the European Union of certain animals and fresh meat and the veterinary certification requirements

(Text with EEA relevance)

THE EUROPEAN COMMISSION,

Having regard to the Treaty on the Functioning of the European Union,

Having regard to 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 (1), and in particular Articles 17(2)(b) and 17(3)(a), the first subparagraph of Article 17(3)(c), the fourth indent of Article 18(1) and Article 19 thereof,

Having regard to Council Directive 2002/99/EC of 16 December 2002 laying down the animal health rules governing the production, processing, distribution and introduction of products of animal origin for human consumption (2), and in particular Article 8, Article 9(2)(b) and Article 9(4) thereof,

Having regard to Council Directive 2004/68/EC of 26 April 2004 laying down animal health rules for the importation into and transit through the Community of certain live ungulate animals, amending Directives 90/426/EEC and 92/65/EEC and repealing Directive 72/462/EEC (3), and in particular the first and second subparagraphs of Article 3(1), the first subparagraph of Article 6(1), Article 7(e), Article 8, the first paragraph of Article 10 and Article 13(1) thereof,

Having regard to Regulation (EC) No 852/2004 of the European Parliament and of the Council of 29 April 2004 on the hygiene of foodstuffs (4), and in particular Article 12 thereof,

Having regard to Regulation (EC) No 853/2004 of the European Parliament and of the Council of 29 April 2004 laying down specific hygiene rules for food of animal origin (5), and in particular Article 9 thereof,

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<sup>(1)</sup> OJ L 268, 14.9.1992, p. 54.

<sup>(&</sup>lt;sup>2</sup>) OJ L 18, 23.1.2003, p. 11.
(<sup>3</sup>) OJ L 139, 30.4.2004, p. 321.

<sup>(&</sup>lt;sup>4</sup>) OJ L 139, 30.4.2004, p. 1.

<sup>&</sup>lt;sup>(5)</sup> OJ L 139, 30.4.2004, p. 55.

Having regard to Regulation (EC) No 854/2004 of the European Parliament and of the Council of 29 April 2004 laying down specific rules for the organisation of official controls on products of animal origin intended for human consumption (<sup>1</sup>), and in particular Article 11(1) and Article 16 thereof,

Having regard to Regulation (EC) No 882/2004 of the European Parliament and of the Council of 29 April 2004 on official controls performed to ensure the verification of compliance with feed and food law, animal health and animal welfare rules (<sup>2</sup>), and in particular Article 48(1) thereof,

Whereas:

- (1) Council Directive 72/462/EEC of 12 December 1972 on health and veterinary inspection problems upon importation of bovine, ovine and caprine animals and swine, fresh meat or meat products from third countries (<sup>3</sup>) provided for a list to be drawn up of the countries or parts thereof from which Member States are to authorise the importation of certain live animals and fresh meat of certain animals.
- (2) Accordingly, Council Decision 79/542/EEC of 21 December 1976 drawing up a list of third countries or parts of third countries, and laying down animal and public health and veterinary certification conditions, for importation into the Community of certain live animals and their fresh meat (<sup>4</sup>) was adopted. That Decision establishes the sanitary conditions for the importation into the European Union of live animals excluding equidae, and for the importation of fresh meat of such animals, including equidae, but excluding meat preparations. Annexes I and II to that Decision also set out lists of third countries or parts thereof from which certain live animals and their fresh meat may be imported into the Union as well as models of veterinary certificates.
- (3) Since the date of adoption of that Decision, a number of new animal health and public health requirements have been laid down in other Union acts, constituting a new regulatory framework in this area. Also, Directive 72/462/EEC has been repealed by Directive 2004/68/EC.
- (4) Article 20 of Directive 2004/68/EC states that implementing rules on import established in accordance with Decisions adopted pursuant to Directive 72/462/EEC, inter alia Decision 79/542/EEC, shall remain in force until replaced by measures adopted under the new regulatory framework.

<sup>(&</sup>lt;sup>1</sup>) OJ L 139, 30.4.2004, p. 206.

<sup>&</sup>lt;sup>(2)</sup> OJ L 165, 30.4.2004, p. 1.

<sup>(&</sup>lt;sup>3</sup>) OJ L 302, 31.12.1972, p. 28.

<sup>(&</sup>lt;sup>4</sup>) OJ L 146, 14.6.1979, p. 15.

- (5) In accordance with Article 4(3) of Directive 2004/41/EC of the European Parliament and of the Council of 21 April 2004 repealing certain Directives concerning food hygiene and health conditions for the production and placing on the market of certain products of animal origin intended for human consumption and amending Council Directives 89/662/EEC and 92/118/EEC and Council Decision 95/408/EC (<sup>1</sup>), once the necessary provisions on the basis of Regulations (EC) No 852/2004, (EC) No 853/2004, (EC) No 854/2004 or Directive 2002/99/EC are adopted, the implementing rules adopted on the basis of Directive 72/462/EEC shall cease to apply.
- (6) Decision 79/542/EEC has been amended several times and import provisions based on the new regulatory framework have already been introduced in Decision 79/542/EEC. For the sake of clarity and transparency the measures that are laid down in Decision 79/542/EEC should be laid down in a new legal act. This Regulation includes all the provisions of Decision 79/542/EEC. Consequently, by the entry into force of the present Regulation Decision 79/542/EEC is lapsed and thus no longer applies, pending the explicit repeal of it.
- (7) Directive 92/65/EEC lays down the animal health requirements governing trade in and imports into the Union of live animals, semen, ova and embryos not subject to the animal health requirements laid down in the specific Union acts referred to in Annex F to that Directive. Pursuant to that Directive, those live animals, semen, ova and embryos may be imported into the Union only from a third country which is on a list drawn up in accordance with the procedure referred to in that Directive. In addition, such live animals are to be accompanied by a health certificate corresponding to a specimen drawn up in accordance with the procedure referred.
- (8) Council Directive 96/93/EC of 17 December 1996 on the certification of animals and animal products (<sup>2</sup>) lays down the rules to be observed in issuing the certificates required by veterinary legislation to prevent misleading or fraudulent certification. It is appropriate to ensure that rules and principles at least equivalent to those laid down in that Directive are applied by the official inspectors or veterinarians of third countries. Certain third countries, which are listed in Annex II to this Regulation, have provided sufficient guarantees as to the existence and implementation of such rules and principles. It is therefore appropriate to authorise the introduction of certain live animals into the Union from those third countries, provided that no further restrictions are required by their specific disease situation.
- (9) Directive 2002/99/EC lays down the animal health rules concerning the introduction into the Union of products of animal origin and products obtained therefrom intended for human consumption. Pursuant to that Directive, lists are to be drawn up of the third countries or regions of third countries from

<sup>(1)</sup> OJ L 157, 30.4.2004, p. 33.

<sup>&</sup>lt;sup>(2)</sup> OJ L 13, 16.1.1997, p. 28.

which imports of specified products of animal origin are permitted and those imports are to comply with certain veterinary certification requirements.

- (10) Directive 2004/68/EC lays down the animal health requirements for the importation into and transit through the Union of certain live ungulates. The importation of those live ungulates into and their transit through the Union is authorised only from third countries and territories that appear on a list or lists drawn up in accordance with the procedure referred to in that Directive and those imports are to comply with certain veterinary certification requirements.
- (11) Save the provisions of article 17(2) last subparagraph of Directive 92/65/EEC, live animals, and products of animal origin to which Directives 92/65/EEC, 2002/99/EC and 2004/68/EC apply are to be imported into or transit through the Union only if they are accompanied by a veterinary certificate and comply with the relevant requirements laid down in Union legislation.
- (12) Accordingly, for the implementation of Directives 92/65/EEC, 2002/99/EC and 2004/68/EC, it is appropriate to lay down in this Regulation lists of third countries, territories and parts thereof and the specific import conditions including model veterinary certificates for certain live animals and the fresh meat of certain animals.
- (13) In the interest of consistency of Union legislation, this Regulation should also take into account the public heath requirements laid down in other Union acts and in particular in Regulations (EC) Nos 852/2004, 853/2004 and 854/2004 which lay down rules concerning the hygiene of foodstuffs and food of animal origin and rules for the organisation of official controls on products of animal origin intended for human consumption, as well as the requirements of Council Directive 96/23/EC of 29 April 1996 on measures to monitor certain substances and residues thereof in live animals and animal products (<sup>1</sup>), and of Regulation (EC) No 999/2001 of the European Parliament and of the Council of 22 May 2001 laying down rules for the prevention, control and eradication of certain transmissible spongiform encephalopathies (<sup>2</sup>).
- (14) Regulation (EC) No 882/2004 lays down general rules governing the performance of official controls carried out in the areas of food and feed, animal health and animal welfare. Article 48 thereof empowers the Commission to adopt a list of third countries from which specific products may be imported into the Union. Regulation (EC) No 854/2004 provides specific rules for the organisation of official controls on products of animal origin intended for human consumption, including the establishment of lists of third countries from which imports of products of animal origin are permitted. Those rules provide that those lists may be combined with other lists drawn up for public and animal health purposes.

<sup>(&</sup>lt;sup>1</sup>) OJ L 125, 23.5.1996, p. 10.

<sup>&</sup>lt;sup>(2)</sup> OJ L 147, 31.5.2001, p. 1.

- (15) The model certificates set out in the Annexes to this Regulation should therefore include attestations certifying that the public health requirements laid down in Directive 96/23/EC and Regulations (EC) No 999/2001, 852/2004, 853/2004 and 854/2004, are fulfilled.
- (16) The model certificates set out in the Annexes to this Regulation should also include attestations certifying that animal welfare requirements laid down in Council Directive 93/119/EC of 22 December 1993 on the protection of animals at the time of slaughter and killing (<sup>1</sup>) and Council Regulation (EC) No 1/2005 of 22 December 2004 on the protection of animals during transport and related operations (<sup>2</sup>) are fulfilled.
- (17) In order to ensure that the health of live animals introduced into the Union is not jeopardised during their transport from the third country of origin to the Union, certain requirements relating to the transport of live animals should be laid down, including requirements on assembly centres.
- (18) In the interest of ensuring the protection of animal health in the Union, live animals should be conveyed directly to their place of destination in the Union.
- (19) Fresh meat introduced into the Union for transit to another third country poses a negligible risk to public health. Such meat should, however, comply with all the relevant animal health requirements. Accordingly, specific provisions on the transit, and storage before transit, of fresh meat should therefore be laid down.
- (20) Specific conditions for transit via the Union of consignments to and from Russia should be provided for owing to the geographical situation of Kaliningrad which affects only Latvia, Lithuania and Poland.
- (21) Consignments of fresh meat, excluding offal and minced meat, of farmed non-domesticated animals of the order Artiodactyla, originating from animals caught in the wild should be authorised for introduction into the Union. In order to rule out any possible animal health risks which could be posed by such introduction, it is appropriate that those animals be separated from wild animals for a period of three months prior to the introduction into the Union of such consignments. Accordingly, the model veterinary certificate for those consignments (RUF) should take that into account.
- (22) Commission Decision 2003/881/EC of 11 December 2003 concerning the animal health and certification conditions for imports of bees (*Apis mellifera* and *Bombus* spp.) from certain third countries (<sup>3</sup>) lays down the animal health and certification conditions for imports of bees from certain third countries. In the interest of simplification of Union legislation, the measures laid down in that Decision should be included in this Regulation. Consequently, Decision 2003/881/EC should be repealed.

<sup>(1)</sup> OJ L 340, 31.12.1993, p. 21.

<sup>(&</sup>lt;sup>2</sup>) OJ L 3, 5.1.2005, p. 1.

<sup>&</sup>lt;sup>(3)</sup> OJ L 328, 17.12.2003, p. 26.

- (23) It's appropriate to introduce a transitional period to allow Member States and industry to take the necessary measures to comply with the requirements laid down in this Regulation.
- (24) 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:

### CHAPTER I

#### SUBJECT MATTER, SCOPE AND DEFINITIONS

#### Article 1

#### Subject matter and scope

1. This Regulation lays down the veterinary certification requirements for the introduction into the Union of consignments containing the following live animals or fresh meat:

- (a) ungulates;
- (b) the animals listed in Part 2 of Annex IV;
- (c) fresh meat intended for human consumption, excluding meat preparations, of ungulates and equidae.

2. This Regulation lays down the lists of third countries, territories or parts thereof from which the consignments referred to in paragraph 1 may be introduced into the Union.

3. This Regulation shall not apply to the introduction into the Union of non-domesticated animals:

- (a) for shows or exhibitions where such live animals are not regularly kept or bred;
- (b) forming part of circuses;
- (c) intended for an approved body, institute or centre as defined in Article 2(1)(c) of Directive 92/65/EEC.

4. This Regulation shall apply without prejudice to any specific certification requirements laid down in other Union acts or in agreements concluded by the Union with third countries.

#### Article 2

#### Definitions

For the purposes of this Regulation, the following definitions shall apply:

(a) 'ungulates' means ungulates as defined in Article 2(d) of Directive 2004/68/EC;

- (b) 'fresh meat' means fresh meat as defined in point 1.10 of Annex I to Regulation (EC) No 853/2004;
- (c) 'equidae' means equidae as defined in Article 2(b) of Council Directive 90/426/EEC (<sup>1</sup>);
- (d) 'holding' means a farm or other officially supervised agricultural, industrial or commercial undertaking, including zoos, amusement parks and wildlife or hunting reserves where live animals are regularly kept or bred.

#### CHAPTER II

# CONDITIONS FOR THE INTRODUCTION OF LIVE ANIMALS INTO THE UNION

#### Article 3

#### General conditions for the introduction of ungulates into the Union

Consignments of ungulates shall only be introduced into the Union if they comply with the following conditions:

- (a) they come from the third countries, territories or parts thereof listed in columns 1, 2 and 3 of the table set out in Part 1 of Annex I for which there is a model veterinary certificate corresponding to the consignment concerned listed in column 4 of the table in Part 1 of Annex I;
- (b) they are accompanied by the appropriate veterinary certificate, drawn up in accordance with the relevant model veterinary certificate set out in Part 2 of Annex I, taking into account the specific conditions indicated in column 6 of the table in Part 1 of that Annex, and completed and signed by an official veterinarian of the exporting third country;
- (c) they comply with the requirements set out in the veterinary certificate referred to in point (b), including:
  - (i) the supplementary guarantees laid down in that certificate, where indicated in column 5 of the table in Part 1 of Annex I;
  - (ii) any additional veterinary certification requirements that the Member State of destination may impose in accordance with Union veterinary legislation and which are included in the certificate.

<sup>(&</sup>lt;sup>1</sup>) OJ L 224, 18.8.1990, p. 42.

#### Article 4

## Conditions for assembly centres for certain consignments of ungulates

Consignments of ungulates which contain live animals from more than one holding shall only be introduced into the Union if they are assembled in assembly centres approved by the competent authority of the third country of origin in accordance with the requirements set out in Part 5 of Annex I.

#### Article 5

## Protocols for the standardisation of materials and sampling and testing procedures for ungulates

Where sampling and testing is required by the veterinary certificates listed in column 4 of the table in Part 1 of Annex I for the diseases listed in Part 6 of that Annex, for the introduction into the Union of consignments of ungulates, such sampling and testing shall be carried out by or under the control of the competent authority of the third country of origin in accordance with the Protocols for the standard-isation of materials and testing procedures set out in Part 6 of that Annex.

#### Article 6

#### Special conditions for certain consignments of ungulates imported into St Pierre and Miquelon and introduced into the Union

Consignments of ungulates of the species listed in the table in Part 7 of Annex I which were introduced into St Pierre and Miquelon less than six months prior to the date of shipment from St Pierre and Miquelon to the Union shall only be introduced into the Union if:

- (a) they comply with the residence and quarantine requirements set out in Chapter 1 of that Part;
- (b) they have been tested in accordance with the animal health test requirements set out in Chapter 2 of that Part.

#### Article 7

# General conditions for the introduction into the Union of certain species of bees

1. Consignments of bees of the species listed in table 1 of Part 2 of Annex IV shall only be introduced into the Union from third countries or territories:

- (a) listed in Part 1 of Annex II;
- (b) where the presence of the American foulbrood, the small hive beetle (*Aethina tumida*) and the Tropilaelaps mite (*Tropilaelaps* spp.) is subject to compulsory notification throughout the whole territory of the third country or territory concerned.

2. By way of derogation from paragraph 1(a), consignments of bees may be introduced into the Union from a part of a third country or territory listed in Part 1 of Annex II which is:

- (a) a geographically and epidemiologically isolated part of the third country or territory
- (b) listed in the third column of the table in Section 1 of Part 1 of Annex IV.

When that derogation is applied, the introduction into the Union of consignments of bees shall be prohibited from all other parts of the third country or territory concerned not listed in the third column of the table in Section 1 of Part 1 of Annex IV.

3. Consignments of bees of the species listed in table 1 of Part 2 of Annex IV shall consist of either:

- (a) cages of queen bees (*Apis mellifera* and *Bombus* spp.), each containing one single queen bee with a maximum of 20 accompanying attendants; or
- (b) containers of bumble bees (*Bombus* spp.), each containing a colony of a maximum of 200 adult bumble bees.

4. Consignments of bees of the species listed in table 1 of Part 2 of Annex IV shall:

- (a) be accompanied by the appropriate veterinary certificate, drawn up in accordance with the relevant model veterinary certificate set out in Part 2 of Annex IV, and completed and signed by an official inspector of the exporting third country;
- (b) comply with the veterinary requirements set out in the veterinary certificate referred to in point (a).

#### Article 8

## General conditions concerning the transport of live animals to the Union

During the period after loading in the third country of origin and before arrival at the border inspection post of introduction into the Union, consignments of live animals shall not be:

- (a) transported together with live animals that:
  - (i) are not intended for introduction into the Union; or
  - (ii) are of a lower health status;
- (b) unloaded in, or when transported by air, moved to another aircraft, or transported by road, by rail or moved on foot through a third country, territory or a part thereof which is not listed in columns 1, 2 and 3 of the table set out in Part 1 of Annex I or for which there is no model veterinary certificate corresponding to the consignment concerned listed in column 4 of the table in Part 1 of Annex I.

#### Article 9

#### Time limit for the period of transport to the Union of live animals

Consignments of live animals shall only be introduced into the Union where the consignment arrives at the border inspection post of introduction into the Union within 10 days of the date of issue of the appropriate veterinary certificate.

In the case of transport by sea, that period of 10 days shall be extended by an additional period corresponding to the duration of the journey by sea, as certified by a signed declaration of the master of the ship, drawn up in accordance with Part 3 of Annex I and attached in its original form to the veterinary certificate.

#### Article 10

#### Special conditions regarding the spraying of consignments of live animals transported by air to the Union

Where consignments of live animals, excluding consignments of bees, are transported by air, the crate or container in which they are transported and the surrounding area shall be sprayed with an appropriate insecticide.

That spraying shall be carried out immediately prior to the closing of the aircraft doors after loading, and after any subsequent opening of the doors in a third country, until the aircraft reaches its final destination.

The captain of the aircraft shall certify that the spraying has been carried out by signing a declaration, drawn up in accordance with Part 4 of Annex I and attached in its original form to the veterinary certificate.

#### Article 11

#### Conditions to be applied following the introduction into the Union of certain consignments of ungulates

1. Following their introduction into the Union, consignments of ungulates intended for breeding and production, or intended for zoos, amusement parks and wildlife or hunting reserves, shall be conveyed without delay to the holding of destination.

The ungulates shall remain on that holding for a period of at least 30 days, unless they are dispatched directly to a slaughterhouse.

2. Following their introduction into the Union, consignments of ungulates intended for immediate slaughter shall be conveyed without delay to the slaughterhouse of destination where they shall be slaughtered within five working days from the date of arrival at the slaughterhouse.

#### Article 12

# Specific conditions concerning the transit through third countries of certain consignments of ungulates

Where specific condition I of Part 1 of Annex I applies, in order to allow consignments of the ungulates referred to in that condition originating in one Member State and destined for another Member State, to transit through a third country, territory or part thereof which is listed in the table in Part 1 of Annex I but for which there is no corresponding model veterinary certificate for consignments of the ungulates concerned indicated in column 4 of that table, the following conditions shall apply:

- (a) for bovine animals for fattening:
  - (i) the holdings of final destination must be designated in advance by the competent authority of the final destination;
  - (ii) the live animals comprised in the consignment must not be moved from the holding of final destination unless for immediate slaughter;
  - (iii) all movements of live animals into and out of the holding of final destination must be carried out under the control of the competent authority as long as the animals comprising the consignment are kept at the holding.
- (b) for ungulates for immediate slaughter, Article 11(2) shall apply.

#### Article 13

#### Conditions to be applied following the introduction into the Union of consignments of bees referred to in Article 7

1. Consignments of queen bees referred to in Article 7(3)(a) shall be conveyed without delay to the designated place of final destination where the hives shall be placed under the control of the competent authority and the queen bees transferred to new cages before being introduced to local colonies.

2. The cages, attendants, and other material that accompanied the queen bees from the third country of origin shall be sent to a laboratory designated by the competent authority for examination for the presence of:

(a) the small hive beetle (Aethina tumida), their eggs or larvae;

(b) signs of the Tropilaelaps mite (Tropilaelaps spp.).

After that laboratory examination, the cages, attendants and the material shall be destroyed.

3. Consignments of bumble bees (*Bombus spp.*) referred to in Article 7(3)(b) shall be conveyed without delay to the designated place of destination.

Those bumble bees may stay in the container in which they were introduced into the Union until the end of the lifespan of the colony.

That container and the material that accompanied the bumble bees from the third country of origin shall be destroyed at the end of the lifespan of the colony at the latest.

#### CHAPTER III

# CONDITIONS FOR THE INTRODUCTION OF FRESH MEAT INTO THE UNION

#### Article 14

### General conditions for the importation of fresh meat

Consignments of fresh meat intended for human consumption shall only be imported into the Union if they comply with the following conditions:

- (a) they come from the third countries, territories or parts thereof listed in columns 1, 2 and 3 of the table in Part 1 of Annex II for which there is a model veterinary certificate corresponding to the consignment concerned listed in column 4 of the table in Part 1 of Annex II;
- (b) they are presented at the border inspection post of introduction into the Union accompanied by the appropriate veterinary certificate, drawn up in accordance with the relevant model veterinary certificate set out in Part 2 of Annex II, taking into account the specific conditions indicated in column 6 of the table in Part 1 of that Annex, and completed and signed by an official veterinarian of the exporting third country;
- (c) they comply with the requirements set out in the veterinary certificate referred to in point (b), including:
  - (i) the supplementary guarantees laid down in that certificate, where indicated in column 5 of the table in Part 1 of Annex II;
  - (ii) any additional veterinary certification requirements that the Member State of destination may impose in accordance with Union veterinary legislation and which are included in the certificate.

#### Article 15

# Conditions to be applied following the importation of unskinned carcases of wild cloven-hoofed game

In accordance with Article 8(2) of Council Directive 97/78/EC (<sup>1</sup>), consignments of unskinned carcases of wild cloven-hoofed game for human consumption after further processing shall be conveyed without delay to the processing establishment of destination.

<sup>(&</sup>lt;sup>1</sup>) OJ L 24, 30.1.1998, p. 9.

#### Article 16

#### Transit and storage of fresh meat

The introduction into the Union of consignments of fresh meat not intended for importation into the Union but destined for a third country either by immediate transit or after storage in the Union in accordance with Article 12(4) and Article 13 of Directive 97/78/EC, shall only be authorised if the consignments comply with the following conditions:

- (a) they come from the third countries, territories or parts thereof listed in columns 1, 2 and 3 of the table in Part 1 of Annex II, for which there is a model veterinary certificate corresponding to the consignment concerned listed in column 4 of the table in Part 1 of Annex II;
- (b) they comply with the specific animal health requirements for the consignment concerned, as set out in the relevant model veterinary certificate referred to in point (a);
- (c) they are accompanied by a veterinary certificate, drawn up in accordance with the model veterinary certificate set out in Annex III, and completed and signed by an official veterinarian of the exporting third country;
- (d) they are certified as acceptable for transit, including for storage as appropriate, on the common veterinary entry document referred to in Article 2(1) of Commission Regulation (EC) No 136/2004 (<sup>1</sup>), signed by the official veterinarian of the border inspection post of introduction into the Union.

#### Article 17

#### Derogation for transit through Latvia, Lithuania and Poland

1. By way of derogation from Article 16 the transit by road or by rail through the Union, between the designated border inspection posts in Latvia, Lithuania and Poland listed in Commission Decision 2009/821/EC (<sup>2</sup>), of consignments coming from and destined to Russia directly or via another third country shall be authorised provided that the following conditions are complied with:

- (a) the consignment is sealed with a serially numbered seal at the border inspection post of introduction into the Union by the veterinary services of the competent authority;
- (b) the documents accompanying the consignment and referred to in Article 7 of Directive 97/78/EC are stamped 'ONLY FOR TRANSIT TO RUSSIA VIA THE EU' on each page by the official veterinarian of the competent authority responsible for the border inspection post of introduction into the Union;
- (c) the procedural requirements provided for in Article 11 of Directive 97/78/EC are complied with;
- (d) the consignment is certified as acceptable for transit on the common veterinary entry document signed by the official veterinarian of the border inspection post of introduction into the Union.

<sup>(&</sup>lt;sup>1</sup>) OJ L 21, 28.1.2004, p. 11.

<sup>&</sup>lt;sup>(2)</sup> OJ L 296, 12.11.2009, p. 1.

2. Unloading or storage, as defined in Article 12(4) or in Article 13 of Directive 97/78/EC, of such consignments on Union territory shall not be allowed.

3. Regular audits shall be made by the competent authority to ensure that the number of consignments and the quantities of products leaving the Union territory matches the number and quantities entering.

#### CHAPTER IV

#### GENERAL, TRANSITIONAL AND FINAL PROVISIONS

### Article 18

### Certification

The veterinary certificates required by this Regulation shall be completed in accordance with the explanatory notes set out in Annex V.

However, that requirement shall not preclude the use of electronic certification or of other agreed systems, harmonised at Union level.

#### Article 19

#### **Transitional provisions**

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For a transitional period those consignments of live animals, except bees coming from the State of Hawaii, and fresh meat intended for human consumption certified before 30 November 2010 in accordance with Decisions 79/542/EEC and 2003/881/EC may continue to be introduced into the Union until 31 May 2011.

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#### Article 20

#### Repeal

Decision 2003/881/EC is repealed.

#### Article 21

#### Entry into force

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

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

## ANNEX I

### UNGULATES

### PART 1

## List of third countries, territories or parts thereof (\*)

	ISO code and name Code of		Description of third country, territory or part	Veterinary certificate	Specific condi-	
	of third country	Territory	thereof	Model(s)	SG	tions
	1	2	3	4	5	6
▼ <u>M7</u>		CA-0	Whole country	POR-X		
	CA — Canada	CA-1	<ul> <li>Whole country, except the Okanagan Valley region of British Columbia described as follows:</li> <li>From a point on the Canada/United States border 120°15' longitude, 49° latitude</li> <li>Northerly to a point 119°35' longitude, 50°30' latitude</li> <li>North-easterly to a point 119° longitude, 50°45' latitude</li> <li>Southerly to a point on the Canada/United States border 118°15' longitude, 49° latitude</li> </ul>	BOV-X, OVI-X, OVI-Y RUM (*)	A	IVb IX V
▼ <u>C1</u>	CH – Swit- zerland	- Swit- CH 0 Whole country		(**)		
	CL – Chile	CL-0	Whole country	BOV-X, OVI-X, RUM		
		CL-0	Whole country	POR-X, SUI	В	
	GL – Greenland GL		Whole country	OVI-X, RUM		V
	HR – Croatia HR-0		Whole country	BOV-X, BOV-Y, RUM, OVI-X, OVI-Y		
	IS – Iceland IS-0		Whole country	BOV-X, BOV-Y RUM, OVI-X, OVI-Y		
				POR-X, POR-Y	В	
	ME – Montenegro	ME-0	Whole country			I
	MK – The former Yugoslav Republic of Macedonia (***)	MK-0	Whole country			I

<sup>(\*)</sup> Without prejudice to specific certification requirements provided for by any relevant agreement between the Union and third countries.

ISO code and name	Code of	Description of third country, territory or part	Veterinary certificat	Specific	
of third country	Territory	thereof	Model(s)	SG	condi- tions
1	2	3	4	5	6
NZ – New Zealand	NZ-0	Whole country	BOV-X, BOV-Y, RUM, POR-X, POR-Y OVI-X, OVI-Y		ш v
PM – St Pierre and Miquelon	PM-0	Whole country	BOV-X, BOV-Y, RUM, OVI-X, OVI-Y CAM		
RS – Serbia (****)	RS-0	Whole country			I

(\*) Exclusively for live animals other than animals belonging to the cervidae species.

(\*\*) Certificates in accordance with the Agreement between the European Community and the Swiss Confederation on trade in agricultural products (OJ L 114, 30.4.2002, p. 132).

(\*\*\*) The former Yugoslav Republic of Macedonia: the definitive nomenclature for this country will be agreed following current negotiations at UN level.

(\*\*\*\*) Not including Kosovo which is at present under international administration pursuant to United Nations Security Council Resolution 1244 of 10 June 1999.

Specific Conditions (see footnotes in each certificate):

**'I':** for transit through the territory of a third country of live animals for immediate slaughter or live bovine animals for fattening which are consigned from a Member State and destined to another Member State in lorries which have been sealed with a serially numbered seal.

The seal number must be entered on the health certificate issued in accordance with the model laid down in Annex F to Directive 64/432/EEC (<sup>1</sup>) for live bovine animals for slaughter and fattening and in accordance with Model I of Annex E to Directive 91/68/EEC (<sup>2</sup>) for ovine and caprine animals for slaughter.

In addition, the seal must be intact on arrival at the designated border inspection post of entry into the Union and the seal number recorded in the integrated computerised veterinary system of the Union (TRACES).

The certificate must be stamped at the exit point of the Union by the competent veterinary authority prior to transiting one or more third countries, with the following wording 'ONLY FOR TRANSIT BETWEEN DIFFERENT PARTS OF THE EUROPEAN UNION VIA THE FORMER YUGOSLAV REPUBLIC OF MACEDONIA/ MONTENEGRO/SERBIA (\*) (\*\*)'.

Bovine animals for fattening must be transported directly to the holding of destination designated by the competent veterinary authority of destination. Those animals must not be moved from that holding unless for immediate slaughter.

<sup>(\*)</sup> Delete country as applicable.

<sup>(\*\*)</sup> Serbia does not include Kosovo which is at present under international administration pursuant to United Nations Security Council Resolution 1244 of 10 June 1999.

**<sup>&#</sup>x27;II':** territory recognised as having an official tuberculosis-free status for the purposes of exports to the Union of live animals certified according to the model of certificate BOV-X

<sup>(1)</sup> OJ 121, 29.7.1964, p. 1977/64.

<sup>(&</sup>lt;sup>2</sup>) OJ L 46, 19.2.1991, p. 19.

v <u>er</u>	'III':	territory recognised as having an official brucellosis-free status for the purposes of exports to the Union of live animals certified according to the model of certificate BOV-X.
	'IVa':	territory recognised as having an official enzootic-bovine-leukosis (EBL) free status for the purposes of exports to the Union of live animals certified according to the model of certificate BOV $-X$ .
▼ <u>M2</u>	'IVb':	recognised as having officially enzootic-bovine-leukosis (EBL)-free herds equivalent to the requirements set out in Annex D to Directive 64/432/EEC for the purposes of exports to the Union of live animals certified according to the model of certificate BOV–X.
▼ <u>C1</u>	'V':	territory recognised as having an official brucellosis-free status for the purposes of exports to the Union of live animals certified according to the model of certificate OVI-X.
	'VI':	Geographical constraints:
	'VII':	territory recognised as having an official tuberculosis-free status for the purposes of exports to the Union of live animals certified according to the model of certificate RUM.
	'VIII':	territory recognised as having an official brucellosis-free status for the purposes of exports to the Union of live animals certified according to the model of certificate RUM.
	'IX':	territory recognised as having an official Aujeszky's disease -free status for the purposes of exports to the Union of live animals certified according to the model of certificate POR-X.
		PART 2
		Models of Veterinary Certificates
	Models:	
	'BOV-X	<sup>2</sup> : Model of veterinary certificate for domestic bovine animals (including <i>Bubalus</i> and <i>Bison</i> species and their cross-breeds) intended for breeding and/or production after importation.

- 'BOV-Y': Model of veterinary certificate for domestic bovine animals (including *Bubalus* and *Bison* species and their cross-breeds) intended for immediate slaughter after importation.
- 'OVI-X': Model of veterinary certificate for domestic ovine animals (*Ovis aries*) and domestic caprine animals (*Capra hircus*) intended for breeding and/or production after importation.
- 'OVI-Y': Model of veterinary certificate for domestic ovine animals (*Ovis aries*) and domestic caprine animals (*Capra hircus*) intended for immediate slaughter after importation.
- 'POR-X': Model of veterinary certificate for domestic porcine animals (*Sus scrofa*) intended for breeding and/or production after importation;
- "POR-Y": Model of veterinary certificate for domestic porcine animals (*Sus scrofa*) intended for immediate slaughter after importation.

'RUM':	Model of veterinary certificate for animals of the order Artiodactyla
	(excluding bovine animals (including Bubalus and Bison species
	and their cross-breeds), Ovis aries, Capra hircus, Suidae and
	Tayassuidae), and of the families Rhinocerotidae and Elephantidae.

- 'SUI': Model of veterinary certificate for non-domestic *Suidae*, *Tayas-suidae* and *Tapiridae*.
- 'CAM': Model of specific attestation for animals imported from St Pierre and Miquelon under the conditions provided for in Part 7 of Annex I.
- SG (Supplementary guarantees):
- <sup>•</sup>A<sup>•</sup>: guarantees regarding Bluetongue and Epizootic-haemorrhagicdisease tests on animals certified according to the model of certificate BOV-X (point II.2.8 B), OVI-X (point II.2.6 D) and RUM (point II.2.6).
- 'B': guarantees regarding Swine-vesicular-disease and Classical-swinefever tests on animals certified according to the model of certificate POR-X (point II.2.4 B) and SUI (point II.2.4 B).
- 'C': guarantees regarding Brucellosis test on animals certified according to the model of certificate POR-X (point II.2.4 C) and SUI (point II.2.4 C).

Model BOV-X

col	INTR	(				Veterinary cert	tificate to EU
	l.1.	Consignor Name		I.2. Certificate referen		l.2.a.	
		Address		I.3. Central competer	nt authority		
ŧ		Tel.		I.4. Local competent	authority		
dispatched consignment	1.5.	Consignee		I.6.			
sigr		Name					
S		Address					
hed		Postal code					
patc		Tel.					
s of dis	1.7.	Country of origin ISO code	I.8. Region of origin Code	I.9. Country of destination	ISO code	I.10. Region of destination	Code
Part I: Details of	l.11.	Place of origin		l.12.	I		
μ		Name	Approval number				
Pa		Address					
	1.10	Disco of localizer					
	1.13.	Place of loading		I.14. Date of departure			
		Address	Approval number				
	l.15.	Means of transport		I.16. Entry BIP in EU			
		Aeroplane Ship [ Road vehicle Other					
		Identification Documentary references		1.17.			
	l.18.	Description of commodity		I.19. Co	ommodity code 01.02	e (HS code)	
					1.20.	. Quantity	
	1.21.				1.22.	. Number of package	s
	1.23.	Seal/Container No			1.24.		
	1.25.	Commodities certified for:					
		Breeding		Fattening			
				-			
	1.26.			I.27. For import or adr	nission into El	U 🗌	
	1.28.	Identification of the commoditie	S	•			
		Species (scientific name)	Breed Identificat system			Age	Sex

COL	INTRY						Model BOV-X				
	П.	Health	information			II.a. Certificate reference number	II.b.				
	II.1.	Public	Health Attesta	tion							
		I, the	undersigned offic	cial v	eterinarian, hereby certify, that th	ne animals described in this certificate	ə:				
Part II: Certification		II.1.1. come from holdings which have been free from any official prohibition on health grounds, for the past 42 days in brucellosis, for the past 30 days in the case of anthrax and for the past six months in the case of rabies, and, have contact with animals from holdings which did not satisfy these conditions;									
t II:		II.1.2.	have not receiv	ed:	vd:						
Par			— any stilbene	or t	hyrostatic substances,						
					ogenic, gestagenic or β- agonist rective 96/22/EC);	substances for purposes other than t	herapeutic or zootechnic treatment				
		ll.1.3.	with regard to b	ovin	e spongiform encephalopathy (B	SE):					
			( <sup>1</sup> ) ( <sup>2</sup> ) <i>either</i>	[(a)		a permanent identification system en nd are not exposed bovine animals a Regulation (EC) No 999/2001;					
				(b)	from which the ban on the fee	bus cases in the country concerned, the ding of ruminants with meat-and-bor enforced or after the date of birth o ban.]	ne meal and greaves derived from				
			( <sup>1</sup> ) ( <sup>3</sup> ) or	[(a)		a permanent identification system en id are not exposed bovine animals a Regulation (EC) No 999/2001;					
				(b)	meal and greaves derived from	date from which the ban on the feedir ruminants had been effectively enform n after the date of the feed ban.]					
			( <sup>1</sup> ) ( <sup>4</sup> ) or	[(a)		a permanent identification system en id are not exposed bovine animals a Regulation (EC) No 999/2001;					
				(b)	with meat-and-bone meal and g	two years after the date from which the reaves derived from ruminants had be ligenous case if born after the date o	een effectively enforced or after the				
	II.2.	Anima	al Health attesta	ation	:						
		I, the	undersigned offic	cial v	eterinarian, hereby certify, that th	ne animals described above meet the	following requirements:				
		II.2.1.	they come from	the	territory with code:	( <sup>5</sup> ) which, at the date of	of issuing this certificate:				
			( <sup>1</sup> ) either	[(a)	has been free for 24 months fro	om foot-and-mouth disease]					
			( <sup>1</sup> ) or	[(a)	having had cases/outbreaks af	foot-and-mouth disease since ter that date, and authorised to exp Io/, of	oort these animals by Commission				
				(b)		n rinderpest, Rift valley fever, contagic norrhagic disease, and for six monthe					
				(c)		, no vaccination against the diseases domestic cloven-hoofed animals vac					
			( <sup>1</sup> ) either	[(d)	has been free for 24 months fro	om bluetongue;]					
			( <sup>1</sup> ) ( <sup>9</sup> ) or	[(d)	test for the detection of antibody occasions on samples of blood	orn bluetongue, and the animals have y for bluetongue and epizootic haemo taken at the beginning of the isolation (dd/mm/yyyy) and on thin 10 days before export;]	orrhagic disease, carried out on two n/quarantine period and at least 28				

COUNTRY					Model BOV-X	
II.	Health	information		II.a. Certificate reference number	II.b.	
		( <sup>1</sup> ) or	inactivated vaccine, at least 60 serotype/s (inse demonstrated through a surve holding(s) of origin described	but the animation of the animation of the animation of the animation of the date of dispatch to the are those present and the are those present of the animatic and the are the animatic and the are the animatic and the animatic	the Union, against all bluetongue sent in the source population as vith a 150 km radius around the	
	II.2.2.			point II.2.1 since birth, or for at least the last six months before dispatch to n-hoofed animals for the last 30 days;		
	II.2.3.	they have rem reference I.11.:	ained since birth or at least 40 c	lays before dispatch in the holding(	s) of origin described under box	
			nd which, in an area with a 150 km ra previous 60 days,	adius, there has been no case/outbreal	<ul> <li>of epizootic haemorrhagic disease</li> </ul>	
		rinderpest, I		n radius, there has been no case/ou bus bovine pleuropneumonia, lumpy sk		
	II.2.4.		imals to be killed under a national preases referred to under point II.2.1,(a	rogramme for the eradication of diseas a) and (b);	ses, nor have they been vaccinated	
	II.2.5.		n herds that are not restricted und enzootic bovine leukosis;	er the national legislation pertaining	to the eradication of tuberculosis,	
	II.2.6.	they come from	herds recognised as officially tuber	culosis-free ( <sup>6</sup> );		
	and	( <sup>1</sup> ) ( <sup>7</sup> ) <i>either</i>	[come from a region which is recog	nised as officially tuberculosis-free (6)	]	
		( <sup>1</sup> ) or	[have been subjected to an intrade 30 days before dispatch to the Univ	ermal tuberculin test ( <sup>8</sup> ) carried out w on;]	ith negative results within the past	
		( <sup>1</sup> ) <i>or</i>	[are less than six weeks old;]			
	II.2.7.	they have not b	een vaccinated against brucellosis a	and come from herds recognised as o	fficially brucellosis-free ( <sup>6</sup> );	
	and	( <sup>1</sup> ) ( <sup>7</sup> ) <i>either</i>	[come from a region which is recog	nised as officially brucellosis-free (6);]		
		( <sup>1</sup> ) or	[have been subjected to at least one 30 days before dispatch to the Univ	e test for bovine brucellosis ( <sup>8</sup> ) carried c on,]	out on samples taken within the past	
		( <sup>1</sup> ) <i>or</i>	[are less than 12 months old,]			
		( <sup>1</sup> ) <i>or</i>	[are castrated males of any age,]			
( <sup>1</sup> ) either	[11.2.8.			for the control of enzoctic bovine leuko y test of this disease during the past f		
( <sup>1</sup> ) or	[II.2.8.	they come from	herds recognised as officially enzo	otic-bovine-leukosis-free ( <sup>6</sup> ) ( <sup>6a</sup> ),]		
	and	( <sup>1</sup> ) ( <sup>7</sup> ) <i>either</i>	[come from a region which is recog	nised as officially enzootic-bovine-leul	kosis-free ( <sup>6</sup> );]	
		( <sup>1</sup> ) or	[have been subjected to an individu samples taken within the past 30 d	ial test for enzootic bovine leukosis ( <sup>8</sup> ) ays before dispatch to the Union;]	carried out with negative result on	
		( <sup>1</sup> ) or	[are less than 12 months old;]			
	II.2.9.	they are/were ( <sup>1</sup>	) dispatched from their holding(s) of	origin, without passing through any m	narket:	
		( <sup>1</sup> ) either	[directly to the Union,]			
		( <sup>1</sup> ) or	[to the officially authorised assemb described under point II.2.1,]	ly centre described under box referer	nce I.13 situated within the territory	

II.	Health	information	II.a. Certificate reference number	II.b.
		and, until dispatched to the Union:		
		(a) they did not come in contact with other	cloven-boofed animals not complying with th	e health requirements as described i
		this certificate,		
		(b) they were not at any place where, or are case/outbreak of any of the diseases re		e previous 30 days there has been
	II.2.10.	any transport vehicles or containers in whic authorised disinfectant;	h they were loaded were cleaned and disinf	ected before loading with an official
	II.2.11.	they were examined by an official veterinar	ian within 24 hours of loading and showed	no clinical sign of disease;
	II.2.12.		Union on(dd/mm/yyyy) ( <sup>10</sup> leaned and disinfected before loading with a odder could not flow or fall out of the vehic	n officially authorised disinfectant an
II.3.	Anima	I transport attestation		
	loading	indersigned official veterinarian, hereby certif i in accordance with the relevant provisions of e fit for the intended transport.		
( <sup>1</sup> ) ( <sup>11</sup> ) [II.4.	Specif	ic requirements		
	II.4.1.	According to official information, no clinica recorded in the holding(s) of origin referred	al or pathological evidence of infectious b to in box reference I.11, for the last 12 mc	
	II.4.2.	the animals referred to in box reference I.2	8.:	
		(a) have been isolated in accommodation dispatch for export,	approved by the competent authority for th	ne last 30 days immediately prior t
		(b) have been subjected to a serological te results, and all animals in isolation have	st for IBR on sera taken at least 21 days a also given negative results to this test,	fter entry into isolation, with negativ
		(c) have not been vaccinated against IBR.]		
Notes				
This certifica production.	ate is m	eant for domestic bovine animals (including	Bubalus and Bison species and their cross-	breeds) intended for breeding and/o
		animals must be conveyed without delay to t ment outside the holding, except in the case		main for a minimum period of 30 day
Part I:				
— Box refe	rence I.8	3.: Provide the code of territory as appearing	in Part 1 of Annex I to Regulation (EU) No	o 206/2010.
— Box refe No 206/2		3.: The assembly centre, if any, must fulfil th	e conditions for its approval, as laid down ir	Part 5 of Annex I to Regulation (EU
		15.: Registration number (railway wagons or ding and reloading, the consignor must information in the second s		aft) or name (ship) is to be provided
— Box refe	rence I.2	23.: For containers or boxes, the container n	umber and the seal number (if applicable) :	should be included.
— Box refe	rence I.2	28.: Identification system: The animals must	bear:	
	dividual ponder).	number which permits tracing of their premi	ses of origin. Specify the identification syste	em (such as tag, tattoos, brand, chip
		nat includes the ISO code of the exporting	The individual constants and the second s	

COUNTRY			Model BOV-X							
П. Н	ealth information	II.a. Certificate reference number	II.b.							
Species:	Species: Select amongst "Bos", "Bison" and "Bubalus" as appropriate.									
Age: Dat	Age: Date of birth (dd/mm/yy).									
Sex (M =	= male, F = female, C = castrated).									
Breed: s	elect purebred, crossbreed.									
Part II:	Part II:									
( <sup>1</sup> ) Keep as	s appropriate.									
	he animals were born and continuously reared in a country 2001 as a country or region posing a negligible BSE risk ar									
	he country or region of origin is categorised in accordance a controlled BSE risk and is listed as such in Decision 2007		lo 999/2001 as a country or region							
	he country or region of origin has not been categorised in a sed as a country or region with undetermined BSE risk and									
( <sup>5</sup> ) Code of	the territory as it appears in Part 1 of Annex I to Regulation	on (EU) No 206/2010.								
	v tuberculosis/brucellosis-free regions and herds as laid dow and herds as laid down in Chapter I of Annex D to Directiv		; and enzootic-bovine-leukosis-free							
Directive	officially enzootic-bovine-leukosis-free herds recognised as e 64/432/EEC for the purpose of exports to the EU of live ar 6 of Part 1 of Annex I to Regulation (EU) No 206/2010, app	nimals according to the model certifica	ate BOV-X from the territory that, in							
	a territory that, in column 6 of Part 1 of Annex I to Regulation regards brucellosis, and/or "IVa" as regards enzootic bovine		e entry "II", as regards tuberculosis,							
( <sup>8</sup> ) Tests ca No 206/	arried out in accordance with the protocols that, for the dis 2010.	ease concerned, are described in Pa	rt 6 of Annex I to Regulation (EU)							
( <sup>9</sup> ) Supplen entry " <b>A</b>	nentary guarantees to be provided when required in columr ".	n 5 "SG" of Part 1 of Annex I to Regi	ulation (EU) No 206/2010, with the							
Tests fo	or bluetongue and for epizootic haemorrhagic disease in acc	cordance with Part 6 of Annex I to Re	gulation (EU) No 206/2010.							
exportat	loading. Imports of these animals shall not be allowed wh ion to the Union of the third country, territory or part there as have been adopted by the Union against imports of these	of referred to in Boxes I.7 and I.8, o	r during a period where restrictive							
	equired by the EU Member State of destination or Switzerlar ent between the Community and the Swiss Confederation o									
( <sup>12</sup> ) Surveilla	ance programme as laid down in Annex I to Commission reg	gulation (EC) No 1266/2007 (OJ L 28	3, 27.10.2007, p. 37.).							
Official veter	inarian									
Name (i	n capital letters):	Qualification and title:								
Date:		Signature:								
Stamp:										

Model BOV-Y

col	NTR	(	Veterinary certificate to EU				
	l.1.	Consignor Name Address	I.2. Certificate reference No     I.2.a.				
		Tel.	I.3. Central competent authority				
-		10.	I.4. Local competent authority				
ment	1.5.	Consignee	1.6.				
nsign		Name Address					
d co		Postal code					
itche		Tel.					
s of dispatched consignment	1.7.	Country of origin ISO code I.8. Region of origin Code	I.9. Country of ISO code I.10. Region of Code destination				
Part I: Details	l.11.	Place of origin	1.12.				
art I		Name Approval number Address					
-							
	l.13.	Place of loading	I.14. Date of departure				
		Address Approval number					
	l.15.	Means of transport	I.16. Entry BIP in EU				
		Aeroplane 🗌 Ship 🗌 Railway wagon 🗌					
		Road vehicle Other I	1.17.				
		Documentary references					
	l.18.	Description of commodity	I.19. Commodity code (HS code) 01.02				
			I.20. Quantity				
	I.21.		I.22. Number of packages				
	1.23.	Seal/Container No	1.24.				
	1.25.	Commodities certified for:					
		Slaughter 🗌					
	1.26.		I.27. For import or admission into EU				
	1.28.	Identification of the commodities	1				
		Species Breed Identification system (scientific name)	Identification number Age Sex				
		······,					

col	NTRY						Model BOV-Y			
	II.	Health	information			II.a. Certificate reference number	II.b.			
	II.1.	Public	Health Attestation	I						
		I, the u	ndersigned official v	/eteri	narian, hereby certify, that the an	imals described in this certificate:				
Part II: Certification		II.1.1. come from holdings which have been free from any official prohibition on health grounds, for the last 42 days in the brucellosis, for the last 30 days in the case of anthrax, for the last six months in the case of rabies, and, have not been with animals from holdings which did not satisfy these conditions;								
s S I		II.1.2.	have not received	:						
Part I			— any stilbene or	thyr	ostatic substances,					
			<ul> <li>oestrogenic, ar defined in Dire</li> </ul>			ostances for purposes other than ther	apeutic or zootechnic treatment (as			
		II.1.3.	with regard to bov	ine s	pongiform encephalopathy (BSE)	:				
			( <sup>1</sup> ) ( <sup>2</sup> ) <i>either</i>	[(a)		permanent identification system enab ot exposed bovine animals as descrit C) No 999/2001;				
				(b)	from which the ban on the feed	us cases in the country concerned, to ding of ruminants with meat-and-bor enforced or after the date of birth o an.]	ne meal and greaves derived from			
			( <sup>1</sup> ) ( <sup>3</sup> ) or	[(a)		permanent identification system enab not exposed bovine animals as desc (EC) No 999/2001;				
				(b)	and-bone meal and greaves der	e date from which the ban on the ived from ruminants had been effec case if born after the date of the fee	tively enforced or after the date of			
			( <sup>1</sup> ) ( <sup>4</sup> ) or	[(a)		permanent identification system enab not exposed bovine animals as desc (EC) No 999/2001;				
				(b)	with meat-and-bone meal and gr	wo years after the date from which t eaves derived from ruminants had b genous case if born after the date o	een effectively enforced or after the			
	II.2.	Animal	Health Attestation	n						
		I, the u	ndersigned official v	/eteri	narian, hereby certify, that the an	imals described above meet the follo	owing requirements:			
		II.2.1.	they come from th	ie tei	ritory with code:	( <sup>5</sup> ) which, a	t the date of issuing this certificate:			
			( <sup>1</sup> ) either	[(a)	has been free for 24 months from	n foot-and-mouth disease]				
			( <sup>1</sup> ) or	[(a)	had cases/outbreaks after that	oot-and-mouth disease since t date, and authorised to export o/, of	t these animals by Commission			
				(b)		n rinderpest, Rift valley fever, contagio orrhagic disease, and for six months				
				(c)		no vaccination against the diseases domestic cloven-hoofed animals vac				
			( <sup>1</sup> ) either	[(d)	has been free for 24 months from	n bluetongue;]				

Health	nformation		II.a. Certificate reference number	II.b.				
	( <sup>1</sup> ) or [	inactivated vaccine, at least 6 serotype/s demonstrated through a surveil	onths from bluetongue, and the anim 0 days before the date of dispatch to ( <i>insert serotype</i> /s) which are those lance programme ( <sup>9</sup> ) in an area with a reference I.11, and the animals are sti s of the vaccine;]	o the Union, against present in the source 150 km radius around	all bluetongu e population a the holding(s			
II.2.2.			int II.2.1 since birth, or for at least the bofed animals for the last 30 days;	last three months bef	ore dispatch t			
II.2.3.	they have remained	I since birth or at least 40 days be	fore dispatch in the holding(s) describ	ed under box referer	ice I.11:			
		which, in an area with a 150 km ra ious 60 days, and	dius, there has been no case/outbreal	k of epizootic haemo	rhagic diseas			
		r, bluetongue, contagious bovine p	is, there has been no case/outbreak of leuropneumonia, lumpy skin disease					
II.2.4.		ls to be killed under a national pro s referred to in point II.2.1(a) and i	ogramme for the eradication of diseas (b);	es, nor have they b	een vaccinate			
II.2.5.	they come from her	rds:						
	(a) included in an c	official system for the control of en:	zootic bovine leukosis, and					
	(b) that are not rest	tricted under the national legislation	n regarding eradication of tuberculosis	and brucellosis, and				
	(c) recognised as o	fficially tuberculosis free; ( <sup>6</sup> )						
II.2.6.	they have not been	vaccinated against brucellosis and	t they:					
	( <sup>1</sup> ) <i>either</i> [com	ne from herds which are recognise	d as officially brucellosis free;] ( <sup>6</sup> )					
	( <sup>1</sup> ) <i>or</i> [are	castrated males of any age;]						
II.2.7.	<ol> <li>they are individually marked on at least two places on their hindquarters as to show that they are exclusively immediate slaughter; (<sup>7</sup>)</li> </ol>							
II.2.8.	they are/were (1) dis	spatched from their holding(s) of o	rigin, without passing through any mar	ket:				
	( <sup>1</sup> ) <i>either</i> [dire	ctly to the Union,]						
		he officially authorised assembly ribed under point II.2.1]	centre described under box reference	e I.13 situated with	in the territo			
	and, until dispatche	d to the Union:						
	(a) they did not com certificate, and	ne in contact with other cloven-hoof	ed animals not complying with the hea	lth requirements as d	escribed in th			
		at any place where, or around whi f any of the diseases referred to ir	ch within a 10 km radius, during the j n point II.2.1;	previous 30 days the	re has been			
II.2.9.	any transport vehicl authorised disinfecta		ere loaded were cleaned and disinfec	ted before loading w	ith an official			
II.2.10.	they were examined	d by an official veterinarian within 2	24 hours of loading and showed no cli	nical sign of disease	;			
II.2.11.	under box reference	e I.15 above that were cleaned and	(dd/mm/yyyy) ( <sup>8</sup> ) i I disinfected before loading with an offi I not flow or fall out of the vehicle	cially authorised disir	nfectant and s			

II.	Health information	II.a. Certificate reference number	er II.b.
1.3.	Animal transport attestation		
	I, the undersigned official veterinarian, hereby certify, t in accordance with the relevant provisions of Regulati the intended transport.		
Notes	5		
This o	certificate is meant for live bovine animals (including Bu	<i>Ibalus</i> and <i>Bison</i> species and their cross-br	eeds) intended for immediate slaughte
After	importation the animals must be conveyed without dela	ay to the slaughterhouse of destination to b	e slaughtered within five working days
Part I	Ŀ		
— Вс	ox reference I.8: Provide the code of territory as appear	ing in Part 1 of Annex I to Regulation (EU)	No 206/2010.
	ox reference I.13: The assembly centre, if any, must fulfil o 206/2010.	I the conditions for its approval, as laid dowr	n in Part 5 of Annex I to Regulation (EU
	ox reference I.15: Registration number (railway wagons on second se		aft) or name (ship) is to be provided. I
— Во	ox reference I.23: For containers or boxes, the container	r number and the seal number (if applicable	) should be included.
— Во	ox reference I.28: Identification system: the animals mus	at bear:	
_	An individual number which permits tracing of their pre transponder).	emises of origin. Specify the identification sy	rstem (such as tag, tattoos, brand, chi
_	An ear tag that includes the ISO code of the exporti	ng country. The individual number must pe	rmit tracing of their premises of origi
Sp	pecies: Select amongst "Bos", "Bison" and "Bubalus" as	appropriate.	
Ag	ye: Date of birth (dd/mm/yy).		
Se	ex (M = male, F = female, C = castrated).		
Part I	11:		
( <sup>1</sup> ) Ke	eep as appropriate.		
	nly if the animals were born and continuously reared in o 999/2001 as a country or region posing a negligible E		
	nly if the country or region of origin is categorised in a osing a controlled BSE risk and is listed as such in Dec		C) No 999/2001 as a country or regic
	nly if the country or region of origin has not been categ ttegorised as a country or region with undetermined BSI		
<sup>(5</sup> ) Co	ode of the territory as it appears in Part 1 of Annex I to	Regulation (EU) No 206/2010.	
( <sup>6</sup> ) Of	fficially tuberculosis/brucellosis free regions and herds a	is laid down in Annex A to Directive 64/432/	ÆEC.
( <sup>7</sup> ) <b>T</b> ł	nis mark shall take the form of "L" having 13 cm in the l oplied using the technique known as "freeze-branding".	left side and 7 cm in the bottom side with 1	cm of strength in both lines. It shall b

COUNTRY	Model BO\								
II. Health information	II.a. Certificate reference number II.b.								
( <sup>8</sup> ) Date of loading. Imports of these animals shall not be allowed when the animals were loaded either prior to the date of authorisation for exportation to the Union of the third country, territory or part thereof referred to in boxes 1.7 and 1.8, or during a period where restrictive measures have been adopted by the Union against imports of these animals from this third country, territory or part thereof.									
( <sup>9</sup> ) Surveillance programme as laid down in Annex I to Commission regr Official veterinarian	ulauon (EC) No 1266/2007 (CJ L 263, 27.10.2007, p. 37.).								
Name (in capital letters):	Qualification and title:								
Date: Signature:									
Stamp:									

Model OVI-X

cou	NTR	,				Veterinary ce	ertificate to EU		
	l.1.	Consignor Name		I.2. Certificat	e reference No	l.2.a.			
		Address		I.3. Central competent authority					
-		Tel.		I.4. Local co	mpetent authority				
Part I: Details of dispatched consignment	I.5.	Consignee Name Address			1.6.				
atched c		Postal code Tel.							
ls of disp	I.7. Country of origin ISO code I.8. Region of origin Code		Code	I.9. Country destinatio	of ISO code on	e I.10. Region of destination	Code		
Detai	l.11.	Place of origin		l.12.					
Part I:		Name Approval number Address		I.14. Date of departure					
	l.13.	Place of loading							
		Address Approval number							
	l.15.	Means of transport		I.16. Entry BIP in EU					
		Aeroplane Ship Railway wagon Road vehicle Other							
		Identification Documentary references		l.17.					
	l.18.	Description of commodity		I.19. Commodity code (HS code)					
						I.20. Quantity			
	I.21.				ies				
	1.23.	Seal/Container No		1.24.					
	l.25.	Commodities certified for:			L.	~			
		Breeding		Fattening					
	I.26.			I.27. For import or admission into EU					
	1.28.	Identification of the commodities							
		Species Breed (scientific name)	Identificatic system	on Identification number		Age	Sex		

οι	JNTRY					1			Model OVI-		
	11.	Health i	h information II.a. Certificate reference number II.b.								
	11.1.	Public Health Attestation									
		I, the ur	ndersigned	offici	al veterinarian, hereby certify, that the	anima	ls described in this certificat	e:			
Part II: Certification		II.1.1. come from holdings which have been free from any official prohibition on health grounds, for the last 42 days in the orbit brucellosis, for the last 30 days in the case of anthrax, for the last six months in the case of rables, and, have not b contact with animals from holdings which did not satisfy these conditions;									
а) П.Се		II.1.2.	have no	t rece	ived:						
L D			— any	stilbe	ne or thyrostatic substances,						
					ic, androgenic, gestagenic or β- agoni d in Directive 96/22/EC).	st subs	tances for purposes other the	an therapeu	utic or zootechnic treatmer		
	11.2.	Animal	Health at	testa	ion						
		I, the ur	ndersigned	l offici	al veterinarian, hereby certify, that the	e anima	ls described above meet the	e following	requirements:		
		II.2.1.	they cor	ne fro	m the territory with code:		$(^1)$ which, at the date of is	suing this o	certificate:		
			(²) either	<sup>-</sup> [(a)	has been free for 24 months from fo	ot-and-	mouth disease ]				
			(²) or	[(a)	has been considered free from foot- having had cases/outbreaks after th menting Regulation (EU) No/, c	at date	, and authorised to export	hese anim			
				(b)	has been free for 12 months from rin pox, contagious caprine pleuropner vesicular stomatitis,						
				(c)	where during the last 12 months, no carried out and imports of domes permitted;						
			(²) either	· [(d)	has been free for 24 months from b	uetong	ue;]				
			( <sup>2</sup> ) ( <sup>9</sup> ) or	[(d)	has been free for 24 months from blu the detection of antibody for bluetons samples of blood taken at the be on(dd/mm/yyyy have been taken within 10 days before	gue and ginning ) and d	l epizootic haemorrhagic dise of the isolation/quarantine on	ease, carrie period an	ed out on two occasions of nd at least 28 days later		
			( <sup>2</sup> ) or	[(d)	has not been free for 24 months fro vaccine, at least 60 days before the serotype/s) which are those prese programme ( <sup>11</sup> ) in an area with a reference I.11, and the animals are the vaccine;]	date of nt in 1 150 k	dispatch to the Union, agair he source population as o m radius around the holdir	st all bluet lemonstrate ng(s) of or	ongue serotype/s (inse ed through a surveillanc rigin described under bo		
		II.2.2.			ained in the territory described under I without contact with imported clover				x months before dispatch t		
		II.2.3.	they hav	/e rer	nained since birth or at least 40 day	s in th	ə holding(s) described unde	r box refer	rence I.11 before dispatch		
					round which, in an area with a 150 during the previous 60 days, and	km rad	ius, there has been no cas	e/outbreak	of epizootic haemorrhagi		
			rind	erpes	round which, in an area with a 10 k t, Rift valley fever, bluetongue, pesi eumonia and vesicular stomatitis durir	e des	petits ruminants, sheep po				

COUNTRY						Model OVI-X				
II.	Health in	formation			II.a. Certificate reference number	II.b.				
	II.2.4.	according to	o my	knowledge and to the written decla	ration made by the owner, the animal	ls:				
				from holdings, and have not been i ly detected:	n contact with animals of a holding, in	which the following diseases have				
				us agalactia of sheep or goats ( <i>Myc</i> s large colony), within the last six n	oplasma agalactiae, Mycoplasma capr nonths,	icolum, Mycoplasma mycoides var.				
		(ii) paratuberculosis and caseous lymphadenitis, within the last 12 months,								
		(iii) puln	nona	ry adenomatosis, within the last thre	ee years, and					
		(iv) Mae	edi/Vi	sna or caprine viral arthritis/enceph	alitis:					
		(²) eithei	r [wi	thin the last three years,]						
		(²) or			the infected animals were slaught o tests carried out at least six months					
		(b) are inclu	Ided	in an official system for notification	of these diseases, and					
		(c) have be	en f	ree from clinical or other evidence	of tuberculosis and brucellosis durin	ng the three years prior to export;				
	II.2.5.			nals to be killed under a national pr ases referred to in point II.2.1(a) and	ogramme for the eradication of diseas d (b);	es, nor have they been vaccinated				
	II.2.6.	they originat	te:							
		( <sup>2</sup> ) ( <sup>3</sup> ) either	[fror	n the territory described under box	reference I.8, which has been recog	nised as officially brucellosis-free;]				
		(²) or	[fror	n the holding(s) described under b	ox reference I.11, where, in respect o	of brucellosis ( <i>Brucella melitensis</i> ):				
			(a)	all susceptible animals have been	free from clinical or any signs of thi	is disease for the last 12 months,				
			(b)	a representative number of the domeach year to a serological test, $(^{4})$	nestic ovine and caprine animals over a	an age of six months are submitted				
		( <sup>2</sup> ) ( <sup>5</sup> ) either	[(c)	all domestic ovine or caprine anima with Rev. 1 vaccine more than two	als have not been vaccinated against t years ago;	his disease, save those vaccinated				
			(d)	the last two tests ( <sup>6</sup> ), separated by (dd/mm/yyyy) and on months of age gave negative resul	an interval of at least six months, ca 	rried out on ovine and caprine animals over six				
		( <sup>2</sup> ) or	[(c)	domestic ovine or caprine animals Rev. 1 vaccine;	under the age of seven months are v	accinated against this disease with				
			(d)	the last two tests (6), separated by	an interval of at least six months, ca	rried out:				
					v/yyyy) and on(d mals over six months of age , and	ld/mm/yyyy) on all non-vaccinated				
				— on (dd/mm domestic ovine and caprine ani	v/yyyy) and on(d mals over 18 months of age	ld/mm/yyyy) on all vaccinated				
				gave negative results, and]						
			(e)	there are only domestic ovine and o	caprine animals that fulfil at least the a	bove conditions and requirements;]				

COUNTRY					Model OVI-X
II. He	ealth inf	ormation	II.a. Certificate	reference number	II.b.
( <sup>2</sup> ) [II		the uncastrated rams have been kept continuously epididymitis ( <i>Brucella ovis</i> ) has been diagnosed in th days a complement fixation test to detect contagiou	e last 12 months	and, these rams have	e undergone during the previous 30
Ш.	2.8.	In respect of scrapie			
( <sup>2</sup> ) ( <sup>7</sup> ) [II		if they are destined for a Member State which benefi or (c) of Chapter A(I) of Annex VIII to Regulation (EC) programmes referred to in those points and the anir destination regarding scrapie, and]	No 999/2001, th	ne animals comply with	h the guarantees provided for in the
( <sup>1</sup> ) <i>either</i> [II.		are animals intended for production born in and con diagnosed;]	tinuously reared	on holdings in which	a case of scrapie has never been
( <sup>2</sup> ) ( <sup>8</sup> ) or [II.2		they shall have been kept continuously since birth or following requirements for at least three years:	for the last three	e years on a holding c	or holdings which have satisfied the
		- they are subject to regular official veterinary che	cks,		
		- the animals are identified in conformity with Unic	n legislation,		
		<ul> <li>no case of scrapie has been confirmed;</li> </ul>			
		<ul> <li>all animals over the age of 18 months which ha framework of a disease eradication campaign or accordance with the laboratory methods laid No 999/2001;</li> </ul>	slaughtered for	human consumption)	have been examined for scrapie in
		<ul> <li>domestic ovine and caprine animals, with the exc have been introduced into the holding only if t</li> </ul>			
( <sup>2</sup> ) or [11.:	2.8.2.	they are domestic ovine animals of the ARR/ARR p	rion protein geno	otype, as defined in A	nnex I to Decision 2002/1003/EC;]
11.2	2.9.	they are/were ( <sup>1</sup> ) dispatched from their holding(s) of	origin, without p	assing through any m	narket,
		( <sup>2</sup> ) <i>either</i> [directly to the Union,]			
		( <sup>2</sup> ) or [to the officially authorised assembly ca described under point II.2.1.]	entre described	under box reference	e I.13 situated within the territory
		and, until dispatched to the Union:			
		<ul> <li>(a) they did not come in contact with other cloven-he this certificate, and</li> </ul>	ofed animals no	ot complying with the h	nealth requirements as described in
		(b) they were not at any place where, or around wh case/outbreak of any of the diseases referred to		m radius, during the	previous 30 days there has been a
11.2		any transport vehicles or containers in which they w authorised disinfectant;	ere loaded were	cleaned and disinfec	ted before loading with an officially
.2	2.11.	they were examined by an official veterinarian withir	1 24 hours of loa	ading and showed no	clinical sign of disease;
.2		they have been loaded for dispatch to the Union on described under box reference I.15 above that we disinfectant and so constructed that faeces, urine, during transportation.	re cleaned and	disinfected before lo	ading with an officially authorised

II.	Health information	II.a. Certificate reference number	II.b.							
II.3.	Animal transport attestation									
	I, the undersigned official veterinarian, hereby certify, that the animals described above have been treated before and at the time of loading in accordance with the relevant provisions of Regulation (EC) No 1/2005, in particular as regards watering and feeding, and they are fit for the intended transport.									
Notes										
This ce product	ertificate is meant for live domestic ovine animals ( <i>Ovis</i> ion.	aries) and domestic caprine animals (Cap	ora hircus) intended for breeding or							
	portation the animals must be conveyed without delay to further movement outside the holding, except in the case		nain for a minimum period of 30 days							
Part I:										
— Вох	reference I.8: Provide the code of territory as appearing	in Part 1 of Annex I to Regulation (EU) No	206/2010.							
	reference I.13: The assembly centre, if any, must fulfil th 206/2010.	e conditions for its approval, as laid down in	Part 5 of Annex I to Regulation (EU)							
	reference I.15: Registration number (railway wagons or a of unloading and reloading, the consignor must inform		or name (ship) is to be provided. In							
— Вох	reference I.19: Use the appropriate HS code: 01.04.10	or 01.04.20.								
— Вох	reference I.23: For containers or boxes, the container n	umber and the seal number (if applicable) sh	nould be included.							
— Вох	reference I.28: Identification system: The animals must	bear:								
	An individual number which permits tracing of their premi transponder) and the anatomic place used in the animal.		m (such as tag, tattoos, brand, chip,							
— /	An ear tag that includes the ISO code of the exporting	country. The individual number must permit	t tracing of their premises of origin.							
Spe	cies: Select amongst "Ovis aries" and "Capra hircus" as	appropriate.								
Age	: (months).									
Sex	M = male, F = female, C = castrated).									
Part II:										
( <sup>1</sup> ) Co	de of the territory as it appears in Part 1 of Annex I to F	Regulation (EU) No 206/2010.								
(²) Ke	ep as appropriate.									
( <sup>3</sup> ) On	ly for a territory appearing with the entry "V" in column 6	of Part 1 of Annex I to Regulation (EU) No	206/2010.							
( <sup>4</sup> ) The	e representative number of animals to be tested for bruc	ellosis must, for each holding, consist of:								
_	all non-castrated male animals, which have not been va	ccinated against brucellosis, over six months	old,							
_	all non-castrated male animals, which have been vaccin	ated against brucellosis, over 18 months old	,							
_	all animals brought onto the holding since the previous	tests, and								
_	25% of females which are sexually mature, within a min	imum of 50 females.								
	is must be completed when the destination is a Membe /52/EEC.	r State or part of a Member State laid down	n in one of the Annexes of Decision							

cou	NTRY		Model OVI-X					
11.	Health information	II.a. Certificate reference number	II.b.					
(6)	In accordance with Part 6 of Annex I to Regulation (EU) No 206/2	2010.						
	Where more than one holding of origin is involved the date of the most recent test on each holding must be clearly indicated.							
(7)	Guarantees in relation to a programme of control of scrapie, as req and Chapter E of Annex IX to Regulation (EC) No 999/2001.	uested by the EU Member State of de	stination, in application of Article 15					
(8)	In the case of animals intended, exclusively, for breeding purpose	s.						
( <sup>9</sup> )	Supplementary guarantees to be provided when required in column "A". Tests for Bluetongue and for Epizootic-haemorrhagic-disease							
(10)	Date of loading. Imports of these animals shall not be allowed we exportation to the Union of the third country, territory or part them measures have been adopted by the Union against imports of the	eof referred to in boxes I.7 and I.8, c	or during a period where restrictive					
(11)	Surveillance programme as laid down in Annex I to Commission R	Regulation (EC) No 1266/2007 (OJ L 2	283, 27.10.2007, p. 37.).					
Offic	ial veterinarian							
	Name (in capital letters):	Qualification and title:						
	Date: Signature:							
	Stamp:							

Model OVI-Y

col	INTR	(							Veterinary ce	ertificate to EU	
	l.1.	Consignor Name				I.2. Certificate reference No I.2.a.					
		Address				1.3.	Central competence	tent authority			
ent		Tel.				1.4.	Local compete	nt authority			
gnme	1.5.	Consignee				1.6.					
consi		Name Address									
hed		Postal code									
patc		Tel.						-			
Part I: Details of dispatched consignment	1.7.	Country of origin	ISO code	I.8. Region of origin	Code	1.9.	Country of destination	ISO code	I.10. Region of destination	Code	
Deta	1.11.	Place of origin				I.12.					
Part I:		Name Approval number Address									
	I.13.	13. Place of loading Address Approval number					Date of depart	ure			
	l.15.	Means of transport	t			I.16.	Entry BIP in El	U			
		Aeroplane 🗖	Ship 🗌	Railway wagor							
		Road vehicle 🗌	Other [			l.17.					
		Identification									
	118	Documentary refer					119	Commodity co	de (HS code)		
	1.10.	Description of con	mounty								
									0. Quantity		
	1.21.							1.2	2. Number of packag	ges	
	1.23.	Seal/Container No						1.2	4.		
	1.05	Commodition contif	ind for								
	1.20.	Commodities certif									
	1.26.					I.27. For import or admission into EU					
	1.28.	Identification of the	e commodities			I					
		Species (scientific name)	Bree	d Identificat system		lo	lentification num	nber	Age	Sex	

col	INTRY				Model OVI-Y
	II.	Health informatic	วท	II.a. Certificate reference number	II.b.
	II.1.	Public Health A	Ittestation		
		I, the undersigne	ed official veterinarian, hereby certify, that	the animals described in this certificate	
Part II: Certification		brucellosis	m holdings which have been free from an s, for the last 30 days in the case of anthra als from holdings which did not satisfy the	x, for the last six months in the case of r	
II: Cer		II.1.2. have not	received:		
Part		— any sti	ilbene or thyrostatic substances,		
			genic, androgenic, gestagenic or β- agonisi d in Directive 96/22/EC).	t substances for purposes other than the	arapeutic or zootechnic treatment (as
	II.2.	Animal Health a	attestation		
		I, the undersigne	ed official veterinarian, hereby certify, that	the animals described above meet the	following requirements:
		II.2.1. they come this certifi	e from the territory with code:		( <sup>1</sup> ) which, at the date of issuing
		(²) either	[(a) has been free for 24 months from	foot-and-mouth disease ]	
		( <sup>2</sup> ) or		t-and-mouth disease since s after that date, and authorised to ex 	port these animals by Commission
			<ul> <li>(b) has been free for 12 months from r pox, contagious caprine pleuropneu stomatitis,</li> </ul>	inderpest, Rift valley fever, peste des p monia, and epizootic haemorrhagic dise	
			(c) where during the last 12 months, no carried out and imports of domestic	o vaccination against the diseases ment cloven-hoofed animals vaccinated agair	
		( <sup>2</sup> ) either	[(d) has been free for 24 months from	bluetongue;]	
		( <sup>2</sup> ) or	( <i>insert serotype/s</i> ) which are those programme ( <sup>5</sup> ) in an area with a 15	om bluetongue, and the animals have e date of dispatch to the Union, agains present in the source population as c 0 km radius around the holding(s) of o the immunity period of time guaranteed	t all bluetongue serotype/s lemonstrated through a surveillance rigin described under box reference
			remained in the territory described under p and without contact with imported cloven		last three months before dispatch to
		II.2.3. they have	e remained since birth or at least 40 day	ys before dispatch in the holding(s) d	escribed under box reference I.11:
			d around which in an area with a 150 km g the previous 60 days, and	radius there has been no case/outbreal	< of epizootic haemorrhagic disease
		rinder	d around which, in an area with a 10 k rpest, Rift valley fever, bluetongue, peste o ionia and vesicular stomatitis during the pr	des petits ruminants, sheep pox and go	
			not animals to be killed under a national p e diseases referred to in point II.2.1(a) an		ses, nor have they been vaccinated
		II.2.5. they are/w	vere ( <sup>2</sup> ) dispatched from their holding(s) of	<sup>;</sup> origin, without passing through any ma	arket,
		( <sup>2</sup> ) either	[directly to the Union]		

COUNTRY				Model OVI-Y
П.	Health in	nformation	II.a. Certificate reference number	II.b.
		( <sup>2</sup> ) <i>or</i> [to the officially authorised assembly centre under point II.2.1,]	described under box reference I.13 s	ituated within the territory described
		and, until dispatched to the Union:		
		<ul> <li>(a) they did not come in contact with other cloven-ho this certificate, and</li> </ul>	oofed animals not complying with the	health requirements as described in
		(b) they were not at any place where, or around wh case/outbreak of any of the diseases referred to		previous 30 days there has been a
	II.2.6.	in respect of scrapie:		
(2) (3)	[II.2.6.1.	if they are destined for a Member State which benefi or (c) of Chapter A(I) of Annex VIII to Regulation (Ed the programmes referred to in those points, as laid	C) No 999/2001, the animals comply	with the guarantees provided for in
( <sup>2</sup> ) either	[11.2.6.2.	were born in and continuously reared on holdings ir	n which a case of scrapie has never	been diagnosed;]
( <sup>2</sup> ) or	[II.2.6.2.	are domestic ovine animals of the ARR/ARR prion pr from a holding where no case of scrapie has been		to Decision 2002/1003/EC, coming
	II.2.7.	any transport vehicles or containers in which they we authorised disinfectant;	ere loaded were cleaned and disinfed	ted before loading with an officially
	II.2.8.	they were examined by an official veterinarian within	n 24 hours of loading and showed no	o clinical sign of disease;
	II.2.9.	they have been loaded for dispatch to the Union on described under box reference I.15 above that we disinfectant and so constructed that faeces, urine, during transportation.	re cleaned and disinfected before lo	pading with an officially authorised
II.3.	Animal	welfare attestation		
	loading i	ndersigned official veterinarian, hereby certify, that the in accordance with the relevant provisions of Regulatio or the intended transport.		
Notes				
This certific after impor		eant for live domestic ovine animals (Ovis aries) and d	omestic caprine animals (Capra hircu	s) intended for immediate slaughter
After impo	tation the	e animals must be conveyed without delay to the sla	aughterhouse of destination to be sla	aughtered within five working days.
Part I:				
- Box ref	erence I.8	8: Provide the code of territory as appearing in Part 1	I of Annex I to Regulation (EU) No 2	206/2010.
— Box ref No 206		13: The assembly centre, if any, must fulfil the condition	ons for its approval, as laid down in F	Part 5 of Annex I to Regulation (EU)
		15: Registration number (railway wagons or container ig and reloading, the consignor must inform the BIP o		or name (ship) is to be provided. In
- Box ref	erence I. <sup>-</sup>	19: Use the appropriate HS code: 01.04.10 or 01.04.2	20.	
- Box ref	erence I.2	23: For containers or boxes, the container number an	d the seal number (if applicable) sho	ould be included.

со	UNTRY		Model OVI-Y				
11.	Health information	II.a. Certificate reference number	II.b.				
-	- Box reference I.28: Identification system: The animals must bear:						
	<ul> <li>An individual number which permits tracing of their premises of o transponder) and the anatomic place used in the animal.</li> </ul>	origin. Specify the identification system	(such as tag, tattoos, brand, chip,				
	- An ear tag that includes the ISO code of the exporting country	. The individual number must permit	tracing of their premises of origin.				
	Species: Select amongst "Ovis aries" and "Capra hircus" as appropr	iate.					
	Age: months.						
	Sex (M = male, F = female, C = castrated).						
Pa	rt II:						
(1)	Code of the territory as it appears in Part 1 of Annex I to Regulation	n (EU) No 206/2010.					
(2)	Keep as appropriate.						
(3)	Guarantees in relation to a programme of control of scrapie, as requ and Chapter E of Annex IX to Regulation (EC) No 999/2001.	ested by the EU Member State of des	tination, in application of Article 15				
(4)	Date of loading. Imports of these animals shall not be allowed wh exportation to the Union of the third country, territory or part there measures have been adopted by the Union against imports of these	of referred to in boxes I.7 and I.8, or	r during a period where restrictive				
(5)	Surveillance programme as laid down in Annex I to Commission Re	gulation (EC) No 1266/2007 (OJ L 28	3, 27.10.2007, p. 37.).				
Of	ficial veterinarian						
	Name (in capital letters):	Qualification and title:					
	Date:	Signature:					
	Stamp:						
	Date:						

	l.1.	UNTRY Consignor Name Address				1.2. 0	Cortifics	to referen		Veterinary ce	rtificate to EL
		Name				1.2. (	Cortifics	to roforon			
ignment	1.5.			ů					ice numbe	er I.2.a.	
gnment	1.5.	Address	.					Competer	nt Authorit	y	
gnment	1.5.							ompetent	Authority	-	
gnment	1.5.	Tel. No				1.4. L		Jinpeterit	Authonity		
gnme		Consignee				I.6.					
0		Name									
nsi		Address									
200		Postal code									
che		Tel. No									
of dispat	I.7. Country ISO I.8. Region Code of origin code of origin					Country destinat		ISO code	I.10. Region of destination	Code	
ils o	I.11.	Place of origin				I.12.					
I: Deta		Name Address		Approval number							
Part		Name Address		Approval number							
	Name Approval number Address										
	I.13. Place of loading Address Approval number					I.14. Date of departure time of departure					
	I.15. Means of transport Aeroplane Ship Railway wagon					I.16. Entry BIP in EU					
		Road vehicle	Oth	er 🗌		I.17.					
		Identification: Documentary ref	erences:			1.17.					
	I.18.	Description of co	mmodity					I.19. Cor	nmodity c	ode (HS code)	01.03
							-		1.20.	Quantity	
	1.21								l.22.	Number of package	es
	I.23. Identification of container/seal number								1.24.		
	I.25. Commodities certified for: Breeding						F	attening			
	1.26					I.27. F	For impo	ort or adm	ission into	EU	
-	1.28	. Identification of t	he commo	dities		1					
		Species (Scientific name)		Identification system			ification mber	I	۵	ge	Sex

	COUNTR	łY				Model POR-X		
	П.	Health	information		II.a. Certificate reference number	II.b.		
	II.1.	Public	ic Health Attestation					
		I, the u	ndersigned offi	cial veterina	arian, hereby certify, that the animals described	I in this certificate:		
tion		II.1.1	case of bruce	llosis, for the	h have been free from any official prohibition o e last 30 days in the case of anthrax and for the n in contact with animals from holdings which o	e past six months in the case of rabies and,		
Part II: Certification		II.1.2	have not rece	ived:				
ll: Ce			<ul> <li>any stilbe</li> </ul>	ne or thyros	tatic substances,			
Part			•	-	nic, gestagenic or β- agonist substances for pu l in Directive 96/22/EC).	rposes other than therapeutic or zootechnic		
	II.2.	Anima	l Health attest	ation				
		I, the u	ndersigned offi	cial veterina	arian, hereby certify, that the animals described	I above meet the following requirements:		
		II.2.1	they come fro	m the territo	pry with code: (1) which,	at the date of issuing this certificate:		
			(²) either	swine	been free for 24 months from foot-and-mouth dis e fever, classical swine fever, swine vesicular nths from vesicular stomatitis, and]			
			(²) or	A	has been free [for 24 months from foot-and-mout African swine fever, vesicular exanthema, [cla lisease] ( <sup>2</sup> ), and for 6 months from vesicular sto	ssical swine fever] (2) and [swine vesicular		
				[: h	has been considered free from [foot-and-mouth swine vesicular disease] (²), since had cases/outbreaks from that date, and authori Regulation (EU) No/, of	(dd/mm/yyyy), without having sed to export these animals by Commission		
				and	e during the last 12 months, no vaccination ag imports of domestic cloven-hoofed animals v iitted;			
		II.2.2			e territory described under point II.2.1 since bin I without contact with imported cloven-hoofed a			
		II.2.3	dispatch, and	, during this	e holding(s) described under box reference I.1 period, in the holding(s) and in an area with a 1 outbreak of the diseases referred to in point II.2	0 km radius around the holding(s) of origin,		
		II.2.4 A			e killed under a national programme for the e seases referred to in point II.2.1;	radication of diseases, nor have they been		
	(2) (3)	[II.2.4 B	-		within the past 30 days to a test for swine vesicu n negative results in both cases];	lar disease antibodies and a test for classical		
	(²) (4) [	II.2.4 C	they have been negative result		d within the past 30 days to a buffered Bruce	lla antigen test for porcine brucellosis with		
		II.2.5	they come fro	m herds wh	ich are not restricted under the national brucel	losis eradication programme;		
		II.2.6	they are/were	(²) dispatch	ed from their holding(s) of origin, without pass	ing through any market,		
			(²) either	[directly t	o the Union,]			
			(²) or		officially authorised assembly centre described under box reference I.13 situated within the y described under point II.2.1,]			

COUNT	COUNTRY Model POR-X						
II.	Health	information	II.a. Certificate reference number	II.b.			
	II.2.7 II.2.8	<ul> <li>described in this cert</li> <li>(b) they were not at any been a case/outbrea</li> <li>any transport vehicles or officially authorised disin</li> </ul>	contact with other cloven-hoofed animals no lificate, and place where, or around which within a 10 km r k of any of the diseases referred to in point II.2 containers in which they were loaded were cle	adius, during the previous 40 days there has 2.1; eaned and disinfected before loading with an			
	II.2.9	transport described und	for dispatch to the Union on ler box reference I.15 above that were clear fectant and so constructed that faeces, urine, ng transportation.	ned and disinfected before loading with an			
II.3.	l, the u at the	time of loading in accorda	inarian, hereby certify, that the animals desc ance with the relevant provisions of Regulation are fit for the intended transport.				
(²) ( <sup>6</sup> ) [II.4	. Specif	ic requirements					
	[11.4.1	Aujeszky's disease is not	tifiable in the country referred to in box referen	ce I.7;			
	II.4.2	•	rmation, no clinical, pathological or serologica months in the holding(s) of origin referred to nin 5 km;				
	II.4.3	the animals referred to in	box reference I.28:				
			r exportation, have remained since birth in y have remained in this(ese) holdings(s) for th				
			n accommodation approved by the competer export, without direct or indirect contact with of				
			d to an ELISA test for the presence of gl antil vith negative results; and, all animals in isolation				
	(d) have not been vaccinated against Aujeszky's disease and have not been in contact with vaccinated animals and the herd of origin has not been vaccinated during the previous 12 months.]						
(2) (8	<sup>3</sup> ) [II.4.4		]	(further requirements and/or tests)			
Notes	Notes						
This cert	ificate is	meant for live domestic po	prcine animals (Sus scrofa) intended for breed	ing or production.			
Aftor imr	ortation	the enimele must be een	waved without delay to the helding of destine	tion where they shall remain for a minimum			

After importation the animals must be conveyed without delay to the holding of destination where they shall remain for a minimum period of 30 days before further movement outside the holding, except in the case of a dispatch to a slaughterhouse.

со	DUNTRY		Model POR-X
II.	Health information	II.a. Certificate reference number	II.b.
Pa	rt I:		
_	Box reference 1.8. Provide the code of t	erritory as appearing in Part 1 of Annex I	to Begulation (EU) No 206/2010
_			its approval, as laid down in Part 5 of Annex I to
—	ð	er (railway wagons or container and lorrie ading, the consignor must inform the BIF	es), flight number (aircraft) or name (ship) is to be P of entry into the Union.
—	Box reference I.23: For containers or bo	oxes, the container number and the seal	number (if applicable) should be included.
—	Box reference I.28: Identification system	n: the animals must bear:	
	<ul> <li>An individual number which permit brand, chip, transponder).</li> </ul>	s tracing of their premises of origin. Spe	cify the identification system (such as tag, tattoos,
	origin.	de of the exporting country. The individu	al number must permit tracing of their premises of
—	Box reference I.28: Age: months.		
_	Box reference I.28: <i>Sex</i> (M = male, F =	female, $C = castrated$ ).	
Pa	rt II:		
( <sup>1</sup> )	Code of the territory as it appears in Pa	rt 1 of Annex I to Regulation (EU) No 206	6/2010.
(²)	Keep as appropriate.		
(3)	Supplementary guarantees to be provi with the entry 'B'.	ded when required in column 5 'SG' of F	Part 1 of Annex I to Regulation (EU) No 206/2010,
(4)	Supplementary guarantees to be provi with the entry 'C'.	ded when required in column 5 'SG' of F	Part 1 of Annex I to Regulation (EU) No 206/2010,
(5)	for exportation to the Union of the third	l country, territory or part thereof referre	were loaded either prior to the date of authorisation d to in boxes I.7 and I.8, or during a period where animals from this third country, territory or part
( <sup>6</sup> )	between the Community and the Swiss		nce with Decision 2008/185/EC and the Agreement products (OJ L 114, 30.4.2002, p. 132) except for Regulation (EU) No 206/2010.
(7)	To be carried out according to the stand the test used shall be the whole virus E		08/185/EC. In the case of pigs aged over 4 months,
(8)	Further requirements requested by Finl	and in respect of transmissible gastro-er	nteritis.
Off	icial veterinarian		
	Name (in capital letters):	Qualif	ication and title:
	Date:	Signa	ture:
	Stamp:		

					Mode	I POR-Y					
		UNTRY								Veterinary ce	rtificate to EU
	1.1.	Consignor				1.2. C	Sertifica	te referend	ce numbe	er I.2.a.	
		Name				I.3. C	Central (	Competen	t Authorit	y	
		Address				14.1	ocal Co	mpetent A	Authority		
		Tel. No						, inpotonit,			
ent	1.5.	Consignee				I.6.					
nm		Name									
nsiç		Address									
d co		Postal code									
che		Tel. No									
Part I: Details of dispatched consignment	I.7.	Country of origin	ISO code	I.8. Region of origin	Code		Country Iestinati		ISO code	I.10. Region of destination	Code
ils o	l.11.	Place of origin				I.12.					
l: Deta		Name Address		Approval number							
Part		Name Address		Approval number							
	Name Approval number Address										
	I.13. Place of loading Address Approval number				I.14. Date of departure time of departure						
	I.15. Means of transport Aeroplane Ship Railway wagon				I.16. Entry BIP in EU						
		Road vehicle	Oth	er 🗌		l.17.					
		Identification: Documentary ref	erences:								
	I.18	. Description of co	ommodity					I.19. Com	nmodity c	ode (HS code)	01.03
									1.20.	Quantity	
	I.21	-							1.22.	Number of package	es
	I.23. Identification of container/seal number								1.24.		
	I.25. Commodities certified for: Slaughter										
	1.26.					I.27. F	For impo	ort or admi	ssion into	EU	
	1.28	. Identification of t	he commo	dities		I					
		Species (Scientific name)		Identification system			ication nber		A	de	Sex

	COUNTRY Model POR					
	П.	Health	information		II.a. Certificate reference number	II.b.
	II.1.	Public	Health Attest	ation		~
		I, the u	indersigned off	icial veterina	arian, hereby certify, that the animals described	I in this certificate:
		II.1.1	case of bruce	ellosis, for th	h have been free from any official prohibition o e last 30 days in the case of anthrax and for th n in contact with animals from holdings which	e past six months in the case of rabies and,
5		II.1.2	have not rece	eived:		
			— any stilbe	ene or thyros	static substances,	
			-	-	enic, gestagenic or β- agonist substances for pu l in Directive 96/22/EC).	rposes other than therapeutic or zootechnic
	II.2.	Anima	I Health attes	tation		
		I, the u	indersigned off	icial veterina	arian, hereby certify, that the animals described	above meet the following requirements:
		II.2.1	they come fro	om the territo	pry with code: (1) which	, at the date of issuing this certificate:
			(²) either	swin	been free for 24 months from foot-and-mouth dis e fever, classical swine fever, swine vesicular onths from vesicular stomatitis, and]	
( <sup>2</sup> ) or [(a) (i) has been free [for 24 months from foot-and-mouth disease] ( <sup>2</sup> ), for 12 months from African swine fever, vesicular exanthema, [classical swine fever] ( <sup>2</sup> ) and [swine disease] ( <sup>2</sup> ), and for 6 months from vesicular stomatitis, and				assical swine fever] (2) and [swine vesicular		
				[	has been considered free from [foot-and-mout swine vesicular disease] ( <sup>2</sup> ), since cases/outbreaks from that date, and authorise Regulation (EU) No/, of	
				and	e during the last 12 months, no vaccination an imports of domestic cloven-hoofed animals v nitted.	
		II.2.2			e territory described under point II.2.1 since birt d without contact with imported cloven-hoofed	
		II.2.3	dispatch, and	I, during this	e holding(s) described under box reference I.1 period, in the holding(s) and in an area with a putbreak of the diseases referred to in point II.2	10 km radius around the holding(s) of origin,
		II.2.4			e killed under a national programme for the e seases referred to in point II.2.1;	radication of diseases, nor have they been
		II.2.5	they are/were	e (²) dispatch	ned from their holding(s) of origin, without pass	ing through any market,
			(²) either	[directly 1	to the Union,]	
			(²) or	-	ficially authorised assembly centre described described under point II.2.1,]	under box reference I.13 situated within the
			and, until disp	patched to th	he Union:	
			., .	not come in d in this cert	contact with other cloven-hoofed animals not ificate, and	complying with the health requirements as
					place where, or around which within a 10 km ra k of any of the diseases referred to in point II.2.	

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Ι.	Health	information	II.a. Certificate reference number	II.b.
	II.2.6	any transport vehicles o officially authorised disir	I r containers in which they were loaded were cl nfectant;	eaned and disinfected before loading with ar
	II.2.7	they were examined by a	an official veterinarian within 24 hours of loadii	ng and showed no clinical sign of disease;
	II.2.8	transport described und	I for dispatch to the Union on der box reference I.15 that were cleaned and and so constructed that faeces, urine, litter or sportation.	disinfected before loading with an officially
1.3.	Anima	I transport attestation		
	time of		narian, hereby certify, that the animals describe rith the relevant provisions of Regulation (EC) the intended transport.	
(²) (⁴) [II.	.4. Specif	ic requirements		
	II.4.1	Aujeszky's disease is no	tifiable in the country referred to in box referer	nce I.7;
	II.4.2		prmation, no clinical, pathological or serologic s) of origin referred to in box reference I.11, for	
	II.4.3	the animals referred to i	n box reference I.28:	
		(a) have remained in the to dispatch for expo	e holding(s) of origin referred to in box referen rtation, and	ce I.11 since birth or for the last 60 days prio
		(b) have not been vacci	inated against Aujeszky's disease.]	
Notes				
	tificate is	meant for live domestic p	orcine animals ( <i>Sus scrofa</i> ) intended for imme	diate slaughter after importation.
After imj days.	portation	the animals must be conve	eyed without delay to the slaughterhouse of de	stination to be slaughtered within five working
Part I:				
— Box	reference	e I.8: Provide the code of t	territory as appearing in Part 1 of Annex I to Re	egulation (EU) No 206/2010.
		e I.13: The assembly cen EU) No 206/2010.	tre, if any, must fulfil the conditions for its ap	oproval, as laid down in Part 5 of Annex I to
			er (railway wagons or container and lorries), fl ading, the consignor must inform the BIP of er	
— Вох	reference	e I.23: For containers or b	oxes, the container number and the seal numb	per (if applicable) should be included.
		-	<i>m</i> : The animals must bear:	
			ts tracing of their premises of origin. Specify the anatomic place used in the animal.	ne identification system (such as tag, tattoos
	An ear ta origin.	ig that includes the ISO co	ode of the exporting country. The individual nu	mber must permit tracing of their premises o
— Box	reference	e I.28: Age: months.		
	reference	e I.28: <i>Sex</i> (M = male, F =	female C = castrated)	

II. a. Certificate reference number       ILb.         Part II: <ul> <li>Code of the territory as it appears in Part 1 of Annex I to Regulation (EU) No 208/2010.</li> <li>Reep as appropriate.</li> <li>Code of load loading, imposts of these animals shall not be allowed when the animals were loaded ellifer prior to the date of authorisation for expondant in to the third country, territory or part thereor restrictive measures have been adopted by the Union against imports of these animals shall not be allowed when the animals were loaded ellifer prior to the date of authorisation for expondant in to the third country, territory or part thereor restrictive measures have been adopted by the Union against imports of these animals troin this third country, territory or part thereor</li> <li>When required by the EU Member State of destination, in accordance with Decision 2008/185/EC.</li> </ul> <li>Official voterinarian         <ul> <li>Mane (in capital letters):</li> <li>Date:</li> <li>Stamp:</li> <li>Coulification and title:</li> <li>Stamp:</li> </ul> </li>	co	COUNTRY Model POR-Y					
<ul> <li>C) Code of the territory as it appears in Part 1 of Annex 1 to Regulation (EU) No 206/2010.</li> <li>(*) Keep as appropriate.</li> <li>(*) Date of Loading. Imports of these animals shall not be allowed when the animals were loaded either prior to the date of authorisation for exportation to the Union of the Hird county; territory or part thereof reference to in boxes 17 and 1.8, or during a period where restrictive measures have been adopted by the Union against imports of these animals from this third county; territory or part thereof.</li> <li>(*) When required by the EU Member State of destination, in accordance with Decision 2008/185/EC.</li> </ul>	II.	Health information	II.a. Certificate reference number	II.b.			
(*)       Keep as appropriate.         (*)       Lete of loading, imports of these animals shall not be allowed when the animals were loaded either prior to the date of authorisation for exportation to the Union of the third country, territory or part thereof referred to in boxes I.7 and I.8, or during a period when the animals mere loaded either prior to the date of authorisation for exportation to the Union of the third country, territory or part thereof referred to in boxes I.7 and I.8, or during a period when the animals mere loaded either prior to the date of authorisation for exportation to the Union of the third country, territory or part thereof referred to in boxes I.7 and I.8, or during a period when the animals mere loaded either prior to the date of authorisation or part thereof referred to in boxes I.7 and I.8, or during a period when the animals mere loaded either prior to the date of authorisation or part thereof.         (*)       When required by the EU Member State of destination, in accordance with Decision 2008/185/EC.         (*)       When required by the EU Member State of destination, in accordance with Decision 2008/185/EC.         (*)       When required by the EU Member State of destination, in accordance with Decision 2008/185/EC.         (*)       When required by the EU Member State of destination, in accordance with Decision 2008/185/EC.         Official veterinarian       E         Name (in capital letters):       Cualification and title:         Date:       Signature:	Pa	rt II:					
(*)       Date of loading. Imports of these animals shall not be allowed when the animals were loaded either prior to the date of authorisation for exportation to the Union of the third country, territory or part therefor telered to in boxes I.7 and I.8, or during a period where restricte weessures have been adopted by the Union against imports of these animals from this third country, territory or part therefort.         (*)       When required by the EU Member State of destination, in accordance with Decision 2008/185/EC.         (*)       When required by the EU Member State of destination, in accordance with Decision 2008/185/EC.         Official veterinarian       Name (in capital letters):         Name (in capital letters):       Cualification and title:         Date:       Signature:	( <sup>1</sup> )	Code of the territory as it appears in P	art 1 of Annex I to Regulation (EU) No 206/2010	0.			
Official veterinarian         Official veterinarian         Name (in capital letters):       Qualification and title:         Date:       Signature:	(²)	Keep as appropriate.					
Official veterinarian Name (in capital letters): Date: Qualification and title: Signature: Qualification and title: Signature: Qualification and title: Signature: Qualification and title: Qualification and title: Signature: Qualification and title: Qualification and titl	(3)	for exportation to the Union of the thir restrictive measures have been adop	d country, territory or part thereof referred to ir	h boxes I.7 and I.8, or during a period where			
Name (in capital letters):Qualification and title:Date:Signature:	(4)	When required by the EU Member Sta	te of destination, in accordance with Decision 2	2008/185/EC.			
Name (in capital letters):Qualification and title:Date:Signature:							
Name (in capital letters):Qualification and title:Date:Signature:							
Name (in capital letters):Qualification and title:Date:Signature:							
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Name (in capital letters):Qualification and title:Date:Signature:							
Name (in capital letters):Qualification and title:Date:Signature:							
Date: Signature:	Off	icial veterinarian					
		Name (in capital letters):	Qualification	n and title:			
Stamp:		Date:	Signature:				
		Stamp:					

Model RUM

cou	NTR	(	Veterinary certificate to EL
	l.1.	Consignor Name Address	I.2. Certificate reference No I.2.a.
			I.3. Central competent authority
¥		Tel.	I.4. Local competent authority
nmer	l.5.	Consignee	1.6.
nsig		Name Address	
о р		Postal code	
atche		Tel.	
Part I: Details of dispatched consignment	l.7.	Country of origin ISO code I.8. Region of origin Code	I.9. Country of ISO code I.10. Region of Code destination
etail	l.11.	Place of origin	1.12.
Part I: D		Name Approval number Address	
	l.13.	Place of loading	I.14. Date of departure
		Address Approval number	
	l.15.	Means of transport	I.16. Entry BIP in EU
		Aeroplane 🗌 Ship 📃 Railway wagon 🗌	
		Road vehicle Other I Identification Documentary references	I.17. No(s) of CITES
	l.18.	Description of commodity	I.19. Commodity code (HS code)
			I.20. Quantity
	l.21.		I.22. Number of packages
	1.23.	Seal/Container No	1.24.
	1.25.	Commodities certified for:	
		Breeding  Fattening	Slaughter
	1.26.		I.27. For import or admission into EU
	1.28.	Identification of the commodities	
		Species Identification system Identif (scientific name)	ication number Age Sex

co	UNTRY					Model RUM		
	П.	Health	information		II.a. Certificate reference number	II.b.		
	II.1.	Public	Health Attest	tation				
		l, the ı	undersigned of	ficial veterinarian, hereby certify, that the	e animals described in this certificate:			
tion		II.1.1.	come from a holding which has been free from any official prohibition on health grounds, for the last 42 days in the case of brucellosis and tuberculosis, for the last 30 days in the case of anthrax, for the last six months in the case of rabies, and, have not been in contact with animals from holdings which did not satisfy these conditions;					
Part II: Certification		II.1.2.	have not rece	eived:				
Ŭ ≓			— any stilbe	ne or thyrostatic substances,				
Part				ic, androgenic, gestagenic or β- agonist d in Directive 96/22/EC).	t substances for purposes other than	therapeutic or zootechnic treatment		
	11.2.	Anima	l Health Attes	station				
		l, the ι	undersigned of	ficial veterinarian, hereby certify, that the	e animals described above meet the t	following requirements:		
		II.2.1.	they come fro	om the territory with code:	( <sup>1</sup> ) which, at the d	ate of issuing this certificate:		
	-		contagiou	free for 24 months from foot-and-mouth is bovine pleuropneumonia, lumpy skin leuropneumonia and epizootic haemorri	disease, peste des petits ruminants, s	heep pox and goat pox, contagious		
<ul> <li>(b) where during the last 12 months, no vaccination against foot-and-mouth disease, rinderpe bovine pleuropneumonia, lumpy skin disease, peste des petits ruminants, sheep pox and pleuropneumonia and epizootic haemorrhagic disease and during the last 24 months no vac been carried out and imports of cloven-hoofed animals vaccinated against these diseases</li> <li>II.2.2. they have remained</li> </ul>					and goat pox, contagious caprine vaccination against bluetongue has			
			( <sup>2</sup> ) either	[in the territory described under point I Union and without contact with clove				
			(²) or	[in the country of dispatch for at least Part 7 of Annex I to Regulation (EU) N for each species in Part 7 of Annex I to than six months prior to embarkation to not of the same health status after Union ( <sup>3</sup> )]	o 206/2010 and they were imported di Regulation (EU) No 206/2010 from a the Union and in any case they have	rectly under the conditions specified third country during a period of less been separated from other animals		
		II.2.3.	they have real reference I.1	mained since birth or at least 40 days 1 and I.13:	before dispatch in the holding/establ	ishment ( <sup>2</sup> ) described under boxes		
				round which in an area of radius of 1 agic disease during the previous 60 da		break of bluetongue and epizootic		
				ound which in an area of 10 km radius, t ing the previous 40 days;	here has been no case/outbreak of the	e other diseases referred to in point		
		II.2.4.		animals to be killed under a national pro of the diseases referred to in point II.2.1		es, nor have they been vaccinated		
			( <sup>2</sup> ) ( <sup>4</sup> ) <i>either</i>	[come from a herd which is recognise	ed as officially tuberculosis free, and]			
			( <sup>2</sup> ) ( <sup>5</sup> ) or	[have been subjected to an intraderr	mal tuberculin test within the past 3	0 days with negative results, and]		
			they have no	t been vaccinated against brucellosis a	nd they:			
			( <sup>2</sup> ) ( <sup>4</sup> ) <i>either</i>	[come from a herd which is recognise	ed as officially brucellosis free;]			
			( <sup>2</sup> ) ( <sup>5</sup> ) or	[have been subjected to a serum at agglutination per ml, within the past 3		cella count of less than 30 IU of		
			( <sup>2</sup> ) or	[are castrated males of any age;]				

COUN	TRY			Model RUM				
11.	Health	n information	II.a. Certificate reference number	II.b.				
	II.2.5.	according to my knowledge and to the written declara	ation made by the owner, the animals					
		(a) do not come from holdings/establishments $(^{2}),{\rm and}$ which the following diseases have been clinically		nals of a holding/establishment, in				
		<ul> <li>(i) contagious agalactia of sheep or goats (Myco mycoides 'large colony'), within the last six me</li> </ul>		icolum, Mycoplasma mycoides var.				
		(ii) paratuberculosis and caseous lymphadenitis, v	within the last 12 months,					
		(iii) pulmonary adenomatosis, within the last three	years, and					
		(iv) Maedi/Visna or caprine viral arthritis/encephalitis,						
		( <sup>2</sup> ) <i>either</i> [within the last three years,]						
			he infected animals were slaughtered asts carried out at least six months ap					
		(b) are included in an official system for notification of	f these diseases, and					
		(c) have been free from clinical or other evidence of	tuberculosis and brucellosis during th	e three years prior to export;				
	( <sup>2</sup> ) ( <sup>6</sup> ) [II.2.6.	the animals have reacted negatively to a serological rhagic-disease, carried out on two occasions on samp at least 28 days later on	les of blood taken at the beginning of	the isolation/quarantine period and				
	II.2.7.	they are dispatched from the holding/establishment des dispatched to the Union:	scribed under boxes reference I.11 and	d I.13 directly to the Union and, until				
		<ul> <li>(a) they did not come in contact with other cloven-hoo this certificate, and</li> </ul>	ofed animals not complying with the h	ealth requirements as described in				
		(b) they were not at any place where, or around whice case/outbreak of any of the diseases referred to in		previous 30 days there has been a				
	II.2.8.	any transport vehicles or containers in which they we authorised disinfectant;	re loaded were cleaned and disinfect	ed before loading with an officially				
	II.2.9.	they were examined by an official veterinarian within 2	24 hours of loading and showed no c	linical sign of disease;				
	II.2.10.	they have been loaded for dispatch to the Union on . under box reference I.15. above that were cleaned and constructed that faeces, urine, litter or fodder could no	disinfected before loading with an offi	cially authorised disinfectant and so				
11.3.	Anima	I transport attestation						
	loading	undersigned official veterinarian, hereby certify, that the g in accordance with the relevant provisions of Regulatio for the intended transport.						
( <sup>2</sup> ) ( <sup>8</sup> )	[II.4. Specif	fic requirements						
	II.4.1.	According to official information, no clinical or patholog in the holding/establishment ( <sup>2</sup> ) of origin referred to in						
	II.4.2.	the animals referred to in box reference I.28 .:						
		<ul> <li>(a) have been isolated in accommodation approved by for export, and</li> </ul>	the competent authority for the last 30	) days immediately prior to dispatch				
		(b) have been subjected to a serological test for IBR results, and all animals in isolation have also give		r entry into isolation, with negative				

COUNTRY	/			Model RUM			
II.	Health ir	formation	II.a. Certificate reference number	II.b.			
	(c)	) have not been vaccinated against IBR.;					
(2)	[II.4.3	(further requirement	s and/or tests)	]]			
Notes							
	This certificate is meant for live animals of the order Artiodactyla (excluding bovine animals (including <i>Bubalus</i> and <i>Bison</i> species and their cross- breeds), <i>Ovis aries, Capra hircus</i> , Suidae and Tayassuidae), and of the families Rhinocerotidae and Elephantidae. Use one certificate per species.						
		animals must be conveyed without delay to the holdin ment outside the holding, except in the case of a dis		ain for a minimum period of 30 days			
Part I:							
— Box re	eference I.8	3.: Provide the code of territory as appearing in Part	1 of Annex I to Regulation (EU) No 2	206/2010.			
	eference I.1 96/2010.	3.: The assembly centre, if any, must fulfil the conditi	ions for its approval, as laid down in F	art 5 of Annex I to Regulation (EU)			
		5.: Registration number (railway wagons or containe g and reloading, the consignor must inform the BIP of		or name (ship) is to be provided. In			
— Box re	eference I.1	9.: Use the appropriate HS code: 01.02, 01.04.10, 0	01.04.20 or 01.06.19.				
— Box re	eference I.2	23.: For containers or boxes, the container number a	nd the seal number (if applicable) sh	ould be included.			
		8.: Identification system: Specify the identification sys rting country. The individual number must permit tra		nder). The ear tag includes the ISO			
Age: I	months.						
Sex (I	VI = male, I	F = female, C = castrated).					
Speci	<i>es</i> : Select t	the species amongst those listed for the following fa	milies:				
Antilo	capridae:	Antilocapra spp.;					
Bovid	ae:	Addax spp., Aepyceros spp., Alcelaphus spp., Ami laphus spp., Budorcas spp., Capra spp. (excluding (including Beatragus), Dorcatragus spp., Gazella s Madoqua spp., Naemorhedus spp. (including Nemo spp., Oryx spp., Ourebia spp., Ovibos spp., Ovis s Pseudois spp., Pseudoryx spp., Raphicerus spp., F Sylvicapra spp., Syncerus spp., Taurotragus spp.,	Capra hircus), Cephalophus spp., Co spp., Hemitragus spp., Hippotragus s orhaedus and Capricornis), Neotragus spp. (excluding Ovis aries), Pantholop ledunca spp., Rupicapra spp., Saiga s	nnochaetes spp., Damaliscus spp. pp., Kobus spp., Litocranius spp., spp., Oreamnos spp., Oreotragus s spp., Pelea spp., Procapra spp., pp., Sigmoceros-Alecelaphus spp.,			
Came	lidae:	Camelus spp., Lama spp., Vicugna spp.					
Cervio	lae:	Alces spp., Axis-Hyelaphus spp., Blastocerus spp Hippocamelus spp., Hydropotes spp., Mazama sp spp., Pudu spp., Rangifer spp.					
Giraffi	dae:	<i>Giraffa</i> spp., Okapia spp.					
Hippo	potamidae:	Hexaprotodon-Choeropsis spp., Hippopotamus spp	<b>'</b> ,				
Mosch	nidae:	Moschus spp.					
Tragu	lidae:	Hyemoschus spp., Tragulus-Moschiola spp.,					
Rhino	cerotidae:	Ceratotherium spp., Dicerorhinus spp., Diceros spp	o., <i>Rhinoceros</i> spp.				
Eleph	antidae:	Elephas spp., Loxodonta spp., as appropriate.					

со	OUNTRY Model RUM								
11.	Health information	II.a. Certificate reference number	II.b.						
Ра	Part II:								
(1)	<sup>1</sup> ) Code of the territory as it appears in Part 1 of Annex I to Regulation (EU) No 206/2010.								
(2)	Keep as appropriate.								
(3)	In this case the health certificate has to be accompanied by the officia I to Regulation (EU) No 206/2010 (model "CAM").	l document on quarantine and test cor	nditions laid down in Part 2 of Annex						
(4)	<sup>4</sup> ) Officially tuberculosis/brucellosis free regions or herds recognised as equivalent to the requirements laid down in Annex A to Directive 64/432/EEC and which appear in column 6 of Part 1 of Annex I to Regulation (EU) No 206/2010, with the entry "VII" as regards tuberculosis, "VIII", as regards brucellosis.								
(5)	5) Tests carried out in accordance with the protocols that, for the disease concerned, are described in Part 6 of Annex I to Regulation (EU) No 206/2010. However for the tuberculin test a result of an increase in skin fold thickness of 2mm or more, or clinical signs of such as oedema, exudation, necrosis, pain and/or inflammation shall be deemed to be positive.								
(6)	Supplementary guarantees to be provided when required in column 5 "A". Tests for Bluetongue and for Epizootic-haemorrhagic-disease i								
(7)	7) Date of loading. Imports of these animals shall not be allowed when the animals were loaded either prior to the date of authorisation for exportation to the Union of the third country, territory or part thereof referred to in boxes I.7. and I.8., or during a period where restrictive measures have been adopted by the Union against imports of these animals from this third country, territory or part thereof.								
(8)	When required by the EU Member State of destination.								
Of	ficial veterinarian								
	Name (in capital letters):	Qualification and	title:						
	Date:	Signature:							
	Stamp:								
L									

	Model SUI					
	COUNTRY	Veterinary certificate to EU				
	I.1. Consignor	I.2. Certificate reference number I.2.a.				
	Name	I.3. Central Competent Authority				
	Address	I.4. Local Competent Authority				
	Tel. No	1.4. Local Competent Authonity				
ut	I.5. Consignee	1.6.				
nme	Name					
nsig	Address					
l cor	Postal code					
chec	Tel. No					
Part I: Details of dispatched consignment	I.7. Country ISO I.8. Region Code of origin	I.9. Country of ISO I.10. Region of Code destination code destination				
ils o	I.11. Place of origin	1.12.				
l: Deta	Name Approval number Address					
Part	Name Approval number Address					
	Name Approval number Address					
	I.13. Place of loading	I.14. Date of departure time of departure				
	Address Approval number					
	I.15. Means of transport Aeroplane Ship Railway wagon	I.16. Entry BIP in EU				
	Road vehicle Other	I.17. No(s) of CITES				
	Identification: Documentary references:					
	I.18. Description of commodity	I.19. Commodity code (HS code)				
		I.20. Quantity				
	l.21.	I.22. Number of packages				
	I.23. Identification of container/seal number	1.24.				
	I.25. Commodities certified for:					
	Breeding Fattenin	ng Slaughter				
	l.26.	I.27. For import or admission into EU				
	I.28. Identification of the commodities					
	Species Identification (Scientific name) system	Identification Age Sex number				

II. II.1.	Health information	II.a. Certificate reference number	
			II.b.
	Public Health Attestation		
	I, the undersigned official veterin	arian, hereby certify, that the animals describe	d in this certificate:
ation	case of brucellosis, for t	ich has been free from any official prohibition c he last 30 days in the case of anthrax and for th en in contact with animals from holdings which	he past six months in the case of rabies and,
Part II: Certification	II.1.2 have not received:		
ö H	<ul> <li>any stilbene or thyro</li> </ul>	static substances,	
Part		enic, gestagenic or β - agonist substances for p d in Directive 96/22/EC).	urposes other than therapeutic or zootechnic
II.2.	Animal Health attestation		
	I, the undersigned official veterin	narian, hereby certify, that the animals described	d above meet the following requirements:
	II.2.1 they come from the territ	tory with code: (1) which	n, at the date of issuing this certificate:
		I months from foot-and-mouth disease, for 12 r er, swine vesicular disease and vesicular exa	
		st 12 months, no vaccination against these dis als vaccinated against these diseases are not p	
		ne territory described under point II.2.1 since bi d without contact with cloven-hoofed animals im	
	dispatch, and, during thi	ne holding described under boxes reference I.1 s period, in the holding(s) and in an area with a outbreak of the diseases referred to in point II.2	10 km radius around the holding(s) of origin,
1	vaccinated against the c	be killed under a national programme for the e liseases referred to in point II.2.1 and they have n test for porcine brucellosis with negative resu	been subjected within the past 30 days to a
(²) (³) [I		ed within the past 30 days to a test for swine ibodies with negative results in both cases]	vesicular disease antibodies and a test for
(²) (⁴) [I	.2.4 C they have been subject negative results]	ed within the past 30 days to a buffered Bruce	ella antigen test for porcine brucellosis with
	II.2.5 they come from holdings	s which:	
		Inder a national control and eradication prog Teschen disease), and	ramme for brucellosis, porcine enteroviral
	(b) are included in an o	fficial system for notification of these diseases;	
	II.2.6 they are dispatched from dispatched to the Union	n the holding described under boxes reference	I.11 and I.13 directly to the Union and, until
	(a) they did not come in described in this cer	n contact with other cloven-hoofed animals not tificate, and	t complying with the health requirements as
		place where, or around which within a 10 km ra ak of any of the diseases referred to in point II.2	

COUNT	RY			Model SU				
11.	Health	information	II.a. Certificate reference number	II.b.				
	II.2.7 any transport vehicles or containers in which they were loaded were cleaned and disinfected before loading with an officially authorised disinfectant;							
	II.2.8	they were examined by an official veterinarian within 24 hours of loading and showed no clinical sign of disease;						
	II.2.9 they have been loaded for dispatch to the Union on							
II.3.	Anima	I transport attestation						
	time of		arian, hereby certify, that the animals describe th the relevant provisions of Regulation (EC) he intended transport.					
(²) ( <sup>6</sup> ) [II.	4. Specif	ic requirements						
	II.4.1	Aujeszky's disease is not	tifiable in the country referred to in box referen	ce I.7;				
	II.4.2		rmation, no clinical, pathological or serologica nonths in the holding(s) of origin referred to in l d the holding(s);					
	II.4.3	the animals referred to in	box reference I.28:					
			r exportation, have remained since birth in 13 or they have remained in this holding for th					
			n accommodation approved by the competer export, without direct or indirect contact with of					
			d to an ELISA test for the presence of gI antil ith negative results; and, all animals in isolation					
			nated against Aujeszky's disease and have not s not been vaccinated during the previous 12 r					
(²) (	<sup>8</sup> ) [II.4.4		]]	(further requirements and/or tests)				
Notes								
			tic Suidae ( <i>Babyrousa</i> spp., <i>Hylochoerus</i> spp o., <i>Pecari</i> spp., <i>Tayassu</i> spp.) and Tapiridae ( <i>Ta</i>					
			veyed without delay to the holding of destina outside the holding, except in the case of a dis					

СС	COUNTRY Model SUI								
II.	Health information	II.a. Certificate reference numbe	er II.b.						
Pa	Part I:								
_	- Box reference I.8: Provide the code of territory as appearing in Part 1 of Annex I to Regulation (EU) No 206/2010.								
_	Box reference I.13: The assembly centr Regulation (EU) No 206/2010.	e, if any, must fulfil the condition	ns for its approval, as laid down in Part 5 of Annex I to						
_	provided. In case of unloading and reloa	ding, the consignor must inform th	d lorries), flight number (aircraft) or name (ship) is to be he BIP of entry into the Union.						
_	Box reference I.19: Use the appropriate I								
_	Box reference I.23: For containers or box Box reference I.28: <i>Identification system</i>		e seal number (if applicable) should be included.						
		tracing of their premises of origin	n. Specify the identification system (such as tag, tattoos,						
			dividual number must permit tracing of their premises of						
_	Box reference I.28: Age: months.								
—	Box reference I.28: Sex (M = male, F = fe	emale, $C = castrated$ ).							
_	Box reference I.28: <i>Species</i> .								
Pa	rt II:								
(1)	Code of the territory as it appears in Part	t 1 of Annex I to Regulation (EU) N	No 206/2010.						
(²)	Keep as appropriate.								
.,	with the entry ' <b>B</b> '.	•	G' of Part 1 of Annex I to Regulation (EU) No 206/2010,						
(4)	with the entry 'C'.		G' of Part 1 of Annex I to Regulation (EU) No 206/2010,						
( <sup>5</sup> )	for exportation to the Union of the third	country, territory or part thereof r	imals were loaded either prior to the date of authorisation eferred to in boxes I.7 and I.8, or during a period where f Suidae animals from this third country, territory or part						
( <sup>6</sup> )	When required by the EU Member State	of destination, in accordance with	h Decision 2008/185/EC.						
(7)	To be carried out according to the stand 4 months, the test used shall be the who		ecision 2008/185/EC. In the case of animals aged over						
(8)	Further requirements requested by Finla	nd in respect of transmissible gas	stro-enteritis.						
Of	icial veterinarian								
	Name (in capital letters):		Qualification and title:						
	Date:		Signature:						
	Stamp:								

	со	UNTRY				Veterinary ce	rtificate to EU
	l.1.	Consignor		I.2. Certifica	ate reference num	ber I.2.a.	
		Name		I.3. Central Competent Authority			
		Address					
		Tel. No		I.4. Local C	ompetent Authori	.y	
int	I.5.	Consignee		I.6.			
nme		Name					
nsig		Address					
d co		Postal code					
che		Tel. No				1	
Part I: Details of dispatched consignment	1.7.	Country ISO I.8. Region of origin code of origin	Code	I.9. Country destina	/ of ISO tion code	I.10. Region of destination	Code
ils o	I.11.	Place of origin		I.12.			
I: Deta		Name Approval numb Address	er				
Part		Name Approval numb Address					
	Name Approval number Address						
	I.13	Place of loading Address Approval numb	I.14. Date of departure time of departure				
	l.15	Means of transport Aeroplane Ship Railwa	I.16. Entry BIP in EU				
		Road vehicle		l.17. No(s) of CITES			
		Identification: Documentary references:		1.17. NO(S) 01	CITES		
	I.18	. Description of commodity			I.19. Commodit	y code (HS code)	01.06.19
					1.2	0. Quantity	
	I.21				1.2	2. Number of packag	es
	1.23. Identification of container/seal number				1.2	24.	
	I.25	. Commodities certified for:					
	Breeding Fattening				\$	Slaughter	
	1.26.			I.27. For imp	ort or admission i	nto EU	
	1.28	. Identification of the commodities		1			
		Species Identification (Scientific name) system		Identificatior number	1	Age	Sex

Model CAM Specific animal health attestation for animals quarantined in St. Pierre and Miquelon prior to introduction into the Union

	COUNTI	RY						Model CAN	
	Ш.	Health	information	П.	a. Certificate reference num	lber	II.b.		
	II.1.	Quarantine conditions attestation							
fication		(date (d Part 7 d Union a	dd/mm/yyyy) of of Annex I to Re	. released on entry (²)) in the gulation (EU) N period they hav	a, hereby certify, that the an 	nm/yyyy) have Pierre and Mic days	been resident from quelon under the co before being release	nditions provided for in ed for exportation to the	
Part II: Certification		II.1.1. Brucellosis:							
			(a) <i>B. abortus</i> least 42 da		nation Test (SAT) and Rose	Bengal Test (F	RBT) within two days	after arrival and after at	
			(b) <i>B. ovis</i> : Co	mplement Fixa	tion Test (CFT) within two d	ays after arriva	al and after at least 4	2 days	
			(c) <i>B. melitens</i>	sis: SAT and RE	3T within two days after arriv	val and after at	t least 42 days		
		II.1.2.	Bluetongue an	id Epizootic hae	emorrhagic disease				
			(⁵) either	[two tests us 21 days]	ing Bluetongue competitive	e Elisa test wit	hin two days after a	arrival and after at least	
			( <sup>5</sup> ) or		een quarantined for more the of Bluetongue vectors ( <i>C</i>				
		II.1.3.	Tuberculosis						
					est according to annex B to er arrival and after at least 4			ne and avian tuberculin	
		II.1.4.		th disease: ELI d after at least -	SA test for the detection of 42 days	f antibodies ar	nd a virus neutralizat	ton test within two days	
		II.1.5.	Rinderpest: co	mpetitive ELIS	A test within two days after a	arrival and afte	er at least 42 days		
		II.1.6.	Vesicular stom	atitis: ELISA or	virus- neutralisation test wi	thin two days a	after arrival and after	at least 42 days	
		II.1.7.	Rift valley feve	r: an ELISA tes	t or a virus neutralisation te	st within two da	ays after arrival and a	after at least 42 days	
		II.1.8.	Lumpy skin dis	sease: ELISA o	r virus neutralisation test wi	thin two days a	after arrival and after	at least 42 days	
		II.1.9.	Crimean Cong 42 days	jo haemorrhagi	c fever: ELISA or virus neuti	ralisation test v	within two days after	arrival and after at least	
		II.1.10.	Surra: blood m	iicroscopy withi	n two days after arrival and	after at least 4	2 days		
		II.1.11.	Malignant cata	arrhal fever: imn	nunofluorescence test withi	n two days afte	er arrival and after at	least 42 days	
	II.2.	Supple	mentary guara	antees					
		II.2.1		s: AGID test or of destination)	ELISA within two days after ( <sup>5</sup> )	arrival and afte	er at least 42 days (W	Vhen required by the EU	

### \_\_\_\_

Π.	Health information	II.a. Certificate reference number	II.b.					
1.3.	Treatments	I						
	They have been subjected	d to:						
	II.3.1. an internal and external antiparasitic treatment during the quarantine period							
	II.3.2.							
	( <sup>5</sup> ) <i>either</i> [a	a treatment with streptomycin 25mg/kg]						
	., .		pira spp. (specify					
	.,	ng/kg	pria spp. (specily					
		inst rabies (if requested) on nd lot), and with the test result	(dd/mm/yyyy) using vaccine]					
lotes	3							
his c	certificate is meant for live anim	als of the family Camelidae.						
art I:	:							
– Bo	ox reference I 8: Provide the cc	ode of territory as appearing in Part 1 of Annex I	to Begulation (EU) No 206/2010					
			its approval, as laid down in Part 5 of Annex I to					
	egulation (EU) No 206/2010.							
		number (railway wagons or container and lorrie nd reloading, the consignor must inform the BIP	es), flight number (aircraft) or name (ship) is to be P of entry into the Union.					
– Во	ox reference I.23: For container	rs or boxes, the container number and the seal r	number (if applicable) should be included.					
– Во	ox reference I.28: Identification	system: The animals must bear:						
_		h permits tracing of their premises of origin. bonder) and the anatomic place used in the a	Specify the identification system (such as tag animal.					
	<ul> <li>An ear tag that includes the origin.</li> </ul>	ISO code of the exporting country. The individua	al number must permit tracing of their premises of					
– Во	ox reference I.28: Age: months.							
— Во	ox reference I.28: <i>Sex</i> (M = mal	le, $F = female$ , $C = castrated$ ).						
– Во	ox reference I.28: <i>Species</i> : Sele	ect amongst ' <i>Camelus</i> spp.', ' <i>Lama</i> spp.', ' <i>Vicugi</i>	<i>na</i> spp.' as appropriate.					
Part II	l:							
1) Ar	nimal health certificate for non ( f Annex I to Regulation (EU) No		t to the Union (model 'RUM') as laid down in Part 2					
of	ate in which the last animal in a	a group entered the quarantine facility.						
	ests performed in accordance v	with the methods described in Chapter 2 of Part	7 of Annex I to Regulation (EU) No 206/2010.					
²) Da								
2) Da 3) Te	esults of the tests performed m	nust be attached in original to this health attestat	tion.					
<sup>2</sup> ) Da <sup>3</sup> ) Te <sup>4</sup> ) Re	esults of the tests performed m	nust be attached in original to this health attesta	tion.					

# COUNTRY

COUNTRY Mod					
Ш.	Health information	II.a. Certificate reference number	II.b.		
Official veterinarian					
	Name (in capital letters):	Qualification	and title:		
	Date:	Signature:			
	Stamp				

### PART 3

#### Addendum for transport of animals by sea

(To be completed and attached to the veterinary certificate when transport to the Union frontier includes, even for part of the journey, transportation by ship.)

#### Declaration by the master of the ship

at any place outside (exporting country) en	remained on board the ship during the voyage in the Union and that the ship did not call route to the Union other than:
Done at	on
(Port of arrival)	(Date of arrival)
	(signature of master)
(stamp)	
	(name in capital letters and title)

#### PART 4

#### Addendum for transport of animals by air

(To be completed and attached to the veterinary certificate when transport to the Union frontier includes, even for part of the journey, transportation by air.)

Declaration by the captain of the aircraft	
I, the undersigned, captain of the aircraft (name ), declare that the crate or container and the area around the crate or container containing the animals referred to in the attached veterinary certificate No has been sprayed with insecticide before departure.	
on	
(Airport of departure) (Date of departure)	
(signature of captain)	
(stamp)	
(name in capital letters and title)	

### PART 5

#### Conditions for the approval of assembly centres (referred to in Article 4)

In order to be approved, assembly centres must meet the following requirements:

- I. They must be supervised by an official veterinarian.
- II. They must each be situated at the centre of an area of at least 20 km in diameter in which, according to official findings, there has been no case of foot-and-mouth disease for at least a period of 30 days prior to their use as approved assembly centres.

- III. They must, before each use as approved assembly centres, be cleansed and disinfected with a disinfectant officially authorised in the exporting country as effective for the control of foot-and-mouth disease.
- IV. They must have, taking into account their animal capacity:
  - (a) a facility dedicated exclusively for use as an assembly centre;
  - (b) appropriate facilities, that are easy to clean and disinfect, for loading, unloading and adequate housing of a suitable standard for the animals, for watering and feeding them, and for giving them any necessary treatment;
  - (c) appropriate facilities for inspection and isolation;
  - (d) appropriate equipment for cleaning and disinfecting rooms and trucks;
  - (e) an appropriate storage area for fodder, litter and manure;
  - (f) an appropriate system for collecting and disposal of waste water;
  - (g) an office for the official veterinarian.
- When operating, they must have sufficient veterinarians to carry out all duties set out in Part 5;
- VI. They must only admit animals that are individually identified so as to guarantee traceability. To this end, when animals are admitted the owner or the person in charge of the centre must ensure that the animals are properly identified and accompanied by health documents or certificates for the species and categories involved.

In addition, the owner or the person in charge of the assembly centre must record on a register or in a data base, and retain for at least three years the name of the owner, the origin of the animals, the dates of entry and exit, the identification number of the animals or registration number of the herd of origin and the holding of destination, and, the registration number of the carrier and the registration number of the lorry delivering or collecting animals from that assembly centre.

- VII. All animals passing through the assembly centre must fulfil the health conditions established for the introduction of the relevant category of animal into the Union.
- VIII. Animals to be introduced into the Union which pass through an assembly centre must, within six days of arrival at the assembly centre, be loaded and dispatched directly to the border of the exporting country:
  - (a) without coming into contact with cloven-hoofed animals other than animals which fulfil the health conditions established for the introduction of the relevant category of animal into the Union;
  - (b) segregated into consignments so that no consignment contains both animals for breeding or production and animals for immediate slaughter;
  - (c) in transport vehicles or containers which have first been cleansed and disinfected with a disinfectant officially authorised in the exporting country as effective for the control of foot-and-mouth disease and which are so constructed that faeces, urine, litter or fodder cannot flow or fall out during transportation.

- IX. Where the conditions for the export of animals to the Union require that a test is carried out within a specified period before loading, that period must include any period of assembly, up to six days, from the date of arrival of the animals at the approved assembly centre.
- X. The exporting third country must designate the centres which are approved for animals for breeding and production and those centres which are approved for animals for slaughter and must notify the Commission and the competent central authorities of the Member States of the names and addresses of such premises. That information must be updated regularly.
- XI. The exporting third country shall determine the procedure for official supervision of approved assembly centres and shall ensure that such supervision is carried out.
- XII. The approved assembly centres must be regularly inspected by the competent authority of the third country in order to check that the requirements for approval set out in points I to XI continue to be fulfilled.

If those inspections show that those conditions are no longer complied with, the approval of the centre must be suspended. The approval may be restored only when the competent authority of the third country is satisfied that the centre fully complies with the conditions set out in points I to XI.

#### PART 6

### Protocols for the standardisation of materials and testing procedures

(referred to in Article 5)

#### Tuberculosis (TBL)

The single intradermal tuberculin test using bovine tuberculin shall be carried out according to Annex B to Directive 64/432/EEC. In the case of Suidae animals, the single intradermal tuberculin test using avian tuberculin shall be carried out according to Annex B to 64/432/EEC, except that the site of injection shall be the loose skin at the base of the ear.

### ▼<u>M2</u>

#### Brucellosis (Brucella abortus) (BRL)

The serum agglutination test, complement fixation test, buffered brucella antigen test, enzyme-linked immunosorbent assays (ELISA) and fluorescence polarisation assay (FPA) shall be carried out according to Annex C to Directive 64/432/EEC.

### ▼<u>C1</u>

### Brucellosis (Brucella melitensis) (BRL)

Tests shall be carried out according to Annex C to Directive 91/68/EEC.

#### Enzootic Bovine Leukosis (EBL)

The agar gel immuno-diffusion test and the enzyme linked immuno-absorbent assay test (ELISA) shall be carried out according to paragraphs A and C of Chapter II of Annex D to Directive 64/432/EEC.

#### Bluetongue (BTG)

A. The blocking or competitive ELISA test shall be carried out according to the following protocol:

The competitive ELISA using monoclonal antibody 3-17-A3 is capable of identifying antibodies to all known serotypes of bluetongue virus (BTV).

The principle of the test is the interruption of the reaction between BTV antigen and a group-specific monoclonal antibody (3-17-A3) by the addition of test serum. Antibodies to BTV present in the test serum block the reactivity of the monoclonal antibody (Mab) and result in a reduction in the expected colour development after the addition of enzyme labelled antimouse antibody, and chromogen/ substrate. Sera can be tested at a single dilution of 1:5 (spot test – Appendix 1) or may be titrated (serum titration – Appendix 2) to give dilution end-point. Inhibition values higher than 50 % may be regarded as positive.

#### Material and Reagents:

- 1. Appropriate ELISA microtitre plates.
- 2. Antigen: supplied as a cell extracted concentrate, prepared as described below, and stored at either -20 °C or -70 °C.
- 3. Blocking buffer: phosphate buffered saline (PBS) containing 0,3 % BTV negative adult bovine serum, 0,1 % (v/v) Tween-20 (supplied as polyoxyethylene sorbiton monolaurate syrup) in PBS.
- Monoclonal antibody: 3-17-A3 (supplied as hybridoma tissue-culture supernatant) directed against the group-specific polypeptide VP7, stored at - 20 °C or freeze-dried and diluted 1/100 with blocking buffer before use.
- 5. Conjugate: rabbit anti-mouse globulin (adsorbed and eluted) conjugated to horseradish peroxidase and kept in the dark at 4 °C.
- Chromogen and substrate: Orthophenylene diamine (OPD-chromogen) at a final concentration of 0,4 mg/ml in sterile distilled water. Hydrogen peroxide (30 %w/v-substrate) 0,05 % v/v added immediately before use (5µl H<sub>2</sub> O<sub>2</sub> per 10 ml OPD). (*Handle OPD with care - wear rubber* gloves - suspected mutagen).
- 7. 1 Molar sulphuric acid: 26,6 ml of acid added to 473,4 ml of distilled water. (*Remember Acid must be added to water, never water to acid.*)
- 8. Orbital shaker.
- 9. ELISA plate reader (the test may be read visually).

#### Test format

Cc: conjugate control (no serum/ no monoclonal antibody); C++: strong positive serum control; C+: weak positive serum control; C-: negative serum control; Cm: monoclonal antibody control (no serum).

#### APPENDIX 1:

Spot dilution (1:5) format (40 sera/plate)

	Con	trols					Test	Sera				
	1	2	3	4	5	6	7	8	9	10	11	12
A	Ce	C-	1	2	3	4	5	6	7	8	9	10
В	Cc	C-	1	2	3	4	5	6	7	8	9	10

	Con	trols					Test	Sera				
	1	2	3	4	5	6	7	8	9	10	11	12
С	C++	C++										
D	C++	C++										
Е	C+	C+										
F	C+	C+										
G	Cm	Cm										40
Н	Cm	Cm										40

### APPENDIX 2:

Serum titration format (10 sera/plate)

	Con	trols					Test	Sera				
	1	2	3	4	5	6	7	8	9	10	11	12
А	Cc	C-	1:5									1:5
В	Cc	C-	1:10									1:10
С	C++	C++	1:20									1:20
D	C++	C++	1:40									1:40
Е	C+	C+	1:80									1:80
F	C+	C+	1:160									1:160
G	Cm	Cm	1:320									1:320
Н	Cm	Cm	1:640									1:640

Test protocol:

Conjugate control	Wells 1A and 1B are a blank control consisting of BTV
(Cc):	antigen and conjugate. This may be used to blank the
	ELISA reader.

Mab controlColumns 1 and 2, rows G and H are the monoclonal<br/>antibody control and contain BTV antigen, monoclonal<br/>antibody and conjugate. These wells represent maximum<br/>colour. The mean of the optical density readings from<br/>this control represents the 0 % inhibition value.

Positive control Columns 1 and 2, rows C-D-E-F. These wells contain (C++, C+): BTV antigen, BTV strong and weak positive antiserum respectively, Mab and conjugate.

Negative control Wells 2A and 2B are the negative controls, which (C-): Contain BTV antigen, BTV negative antiserum, Mab and conjugate.

Test sera:	For large-scale serological surveys and rapid screening,
	sera may be tested at a single dilution of 1:5 (Appendix
	1). Alternatively, 10 sera may be tested over a dilution
	range from 1:5 to 1:640 (Appendix 2). This will give
	some indication of the titre of antibody in the test sera.

#### Procedure:

- 1. Dilute BTV antigen to pre-titrated concentration in PBS, sonicate briefly to disperse aggregated virus (if sonicator is not available, pipette vigorously) and add 50  $\mu$ l to all wells of the ELISA plate. Tap sides of plate to disperse antigen.
- Incubate at 37 °C for 60 minutes on an orbital shaker. Wash plates three times by flooding and emptying the wells with non-sterile PBS and blot dry on absorbent paper.
- 3. Control wells: Add 100 µl of blocking buffer to Cc wells. Add 50 ul of positive and negative control sera, at a dilution of 1:5 (10 µ l sera + 40 µl blocking buffer), to respective wells C-, C+ and C++. Add 50µl blocking buffer to Mab control wells.

Spot titration method: Add a 1:5 dilution of each test serum in blocking buffer to duplicate wells of columns 3 to 12 (10  $\mu$ l sera + 40  $\mu$ l blocking buffer),

or

Serum titration method: Prepare a two-fold dilution series of each test sample (1:5 to 1:640) in blocking buffer across eight wells of single columns 3 to 12.

- 4. Immediately after the addition of the test sera, dilute Mab 1:100 in blocking buffer and add 50  $\mu$ l to all wells of the plate except for the blank control.
- 5. Incubate at 37  $^{\circ}\mathrm{C}$  for 60 minutes on an orbital shaker. Wash three times with PBS and blot dry.
- 6. Dilute rabbit anti-mouse concentrate to 1/5 000 in blocking buffer and add 50  $\mu l$  to all wells of the plate.
- 7. Incubate at 37 °C for 60 minutes on an orbital shaker. Wash three times with PBS and blot dry.
- 8. Thaw the O-Phenylenediamine dihydrochloride (OPD) and immediately before use add 5 μl of 30 % hydrogen peroxide to each 10 ml of OPD. Add 50 μl to all wells of the plate. Allow colour to develop for approximately 10 minutes and stop the reaction with 1 Molar sulphuric acid (50 μl per well). Colour should develop in the Mab control wells and in those wells containing sera with no antibody to BTV.
- 9. Examine and record the plates either visually or using a spectrophotometric reader.

#### Analysis of results:

Using the software package print out the optical density (OD) values, and the percentage inhibition (PI) for test and control sera based on the mean value recorded in the antigen control wells. The date expressed as OD and PI values are used to determine whether the test has performed within acceptable limits. The upper control limits (UCL) and lower control limits (LCL) for the Mab control (antigen plus Mab in the absence of test sera) are between OD values 0,4 and 1.4. Any plate that fails to conform to the above criteria must be rejected.

If a computer software package is not available print out the OD values using the ELISA printer. Calculate the mean OD value for the antigen control wells, which is equivalent to the 100 % value. Determine the 50 % OD value and manually calculate the positivity and negativity of each sample.

Percentage inhibition (PI) value =  $100 - (OD \text{ of each test control/Mean OD of Cm}) \times 100.$ 

The duplicate negative control serum wells and the duplicate blank wells must record PI values between + 25 % and - 25 %, and between + 95 % and + 105 %, respectively. Failure to be within these limits does not invalidate the plate but does suggest that background colour is developing. The strong and weak positive control sera must record PI values between + 81 % and + 100 %, and between + 51 % and + 80 %, respectively.

The diagnostic threshold for test sera is 50% (PI 50% or OD 50%). Samples recording PI values >50% are recorded negative. Samples that record PI values above and below the threshold for the duplicate wells are considered doubtful; such samples may be re-tested in the spot test and/or titration. Positive samples may also be titrated to provide an indication of the degree of positivity.

Visual reading: Positive and negative samples are easily discernible by eye; weakly positive or strong negative samples may be more difficult to interpret by eye.

#### Preparation of BTV ELISA antigen:

- 1. Wash 40-60 roux of confluent BHK-21 cells three times with serum-free Eagle's medium and infect with bluetongue virus serotype 1 in serum-free Eagle's medium.
- 2. Incubate at 37 °C and examine daily for cytopathic effect (CPE).
- 3. When CPE are complete in 90 % to 100 % of the cell sheet of each roux, harvest the virus by shaking any still-attached cells from the glass.
- 4. Centrifuge at 2 000 to 3 000 rpm to pellet the cells.
- 5. Discard the supernatant and re-suspend the cells in approximately 30 ml of PBS containing 1 % 'Sarkosyl' and 2 ml phenylmethylsulphonyl fluoride (lysis buffer). This may cause the cells to form a gel and more lysis buffer may be added to reduce this effect. (NB: phenylmethylsulphonyl fluoride is harmful handle with extreme caution.)

- 6. Disrupt the cells for 60 seconds using an ultrasonic probe at an amplitude of 30 microns.
- 7. Centrifuge at 10 000 rpm for 10 minutes.
- 8. Store the supernatant at + 4 °C and re-suspend the remaining cell pellet in 10 to 20 ml of lysis buffer.
- 9. Sonicate and clarify, storing the supernatant at each stage, a total of three times.
- 10. Pool the supernatants and centrifuge at 24 000 rpm (100,000 g) for 120 minutes at + 4 °C over a 5 ml cushion of 40 % sucrose (w/v in PBS) using 30 ml Beckmann centrifuge tubes and an SW 28 rotor.
- 11. Discard the supernatant, drain the tubes thoroughly and re-suspend the pellet in PBS by sonication. Store the antigen in aliquots at 20 °C.

Titration of BTV ELISA antigen:

Bluetongue ELISA antigen is titrated by the indirect ELISA. Twofold dilutions of antigen are titrated against a constant dilution (1/100) monoclonal antibody 3-17-A3. The protocol is as follows:

- 1. Titrate a 1:20 dilution of BTV antigen in PBS across the microtitre plate in a twofold dilution series (50  $\mu$ l/well) using a multichannel pipette.
- 2. Incubate for one hour at 37 °C on an orbital shaker.
- 3. Wash plates three times with PBS.
- 4. Add 50  $\mu$ l of monoclonal antibody 3-17-A3 (diluted 1/100) to each well of the microtitre plate.
- 5. Incubate for one hour at 37 °C on an orbital shaker.
- 6. Wash plates three times with PBS.
- 7. Add 50  $\mu$ l of rabbit anti-mouse globulin conjugated to horseradish peroxidase, diluted to a pre-titrated optimal concentration, to each well of the microtitre plate.
- 8. Incubate for one hour at 37 °C on an orbital shaker.
- 9. Add substrate and chromogen as described previously. Stop the reaction after 10 minutes by the addition of 1 Molar sulphuric acid (50 µl/well).

In the competitive assay, the monoclonal antibody must be in excess, therefore a dilution of antigen is chosen which falls on the titration curve (not on the plateau region) which gives approximately 0,8 OD after 10 minutes.

B. The agar gel immuno-diffusion test shall be carried out according to the following protocol:

#### Antigen:

Precipitating antigen is prepared in any cell culture system that supports the rapid multiplication of a reference strain of bluetongue virus. BHK or Vero cells are recommended. Antigen is present in the supernatant fluid at the end of virus growth but requires 50 to 100-fold concentration to be effective. This may be achieved by any standard protein concentration procedure; virus in the antigen may be inactivated by the addition of 0,3 % (v/v) beta-propiolactone.

#### Known positive control serum:

Using the international reference serum and antigen a national standard serum is produced, standardised for optimal proportion against the international reference serum, freeze-dried and used as the known control serum in each test.

#### Test serum

- Procedure: 1 % agarose prepared in borate or sodium barbitol buffer, pH 8,5 to 9,0, is poured into a petri dish to a minimum depth of 3,0 mm. A test pattern of seven moisture-free wells, each 5,0 mm in diameter, is cut in the agar. The pattern consists of one centre well and six wells arranged round it in a circle of radius 3 cm. The central well is filled with the standard antigen. Peripheral wells 2, 4 and 6 are filled with known positive serum, wells 1, 3 and 5 are filled with test sera. The system is incubated for up to 72 hours at room temperature in a closed humid chamber.
- Interpretation: A test serum is positive if it forms a specific precipitin line with the antigen and forms a complete line of identity with the control serum. A test serum is negative if it does not form a specific line with the antigen and it does not bend the line of the control serum. Petri dishes must be examined against a dark background and using indirect illumination.

#### Epizootic haemorrhagic disease (EHD)

The agar gel immuno-diffusion test shall be carried out according to the following protocol:

#### Antigen:

Precipitating antigen is prepared in any cell culture system that supports the rapid multiplication of the appropriate serotype(s) of epizootic haemorrhagic disease virus. BHK or Vero cells are recommended. Antigen is present in the supernatant fluid at the end of virus growth but requires 50 to 100-fold concentration to be effective. This may be achieved by any standard protein concentration procedure; virus in the antigen may be inactivated by the addition of 0,3 % (v/v) beta-propiolactone.

#### Known positive control serum:

Using the international reference serum and antigen a national standard serum is produced, standardised for optimal proportion against the international reference serum, freeze-dried and used as the known control serum in each test.

Test serum	
Procedure:	1 % agarose prepared in borate or sodium barbitol buffer, pH 8,5 to 9,0, is poured into a petri dish to a minimum depth of 3,0 mm. A test pattern of seven moisture-free wells, each 5,0 mm in diameter, is cut in the agar. The pattern consists of one centre well and six wells arranged round it in a circle of radius 3 cm. The central well is filled with the standard antigen. Peripheral wells 2, 4 and 6 are filled with known positive serum, wells 1, 3 and 5 are filled with test sera. The system is incubated for up to 72 hours at room temperature in a closed humid chamber.
Interpretation:	A test serum is positive if it forms a specific precipitin line with the antigen and forms a complete line of identity with the control serum. A test serum is negative if it does not form a specific line with the antigen and it does not bend the line of the control serum. Petri dishes must be examined against a dark background and using indirect illumination.
	ovine rhinotracheitis (IBR) / infectious pustular vulvo-vaginitis (IPV)
A. The serum neu protocol:	atralisation test shall be carried out according to the following
Serum:	All sera are heat-inactivated at 56 °C for 30 minutes before use.
Procedure:	The constant virus-varying serum neutralisation test on microtitre plates employs MDBK or other susceptible cells. The Colorado, Oxford or any other reference strain of the virus is used at 100 TCID50 per 0,025 ml; inactivated undiluted serum samples are mixed with an equal volume (0,025 ml) of virus suspension. The virus/serum mixtures are incubated for 24 hours at 37 °C in the microtitre plates before the MDBK cells are added. Cells are used at a concentration which forms a complete monolayer after 24 hours.
Controls:	(i) virus infectivity assay, (ii) serum toxicity controls, (iii) uninoculated cell culture controls, (iv) reference antisera.
Interpretation:	The results of the neutralisation test and the titre of the virus used in the test are recorded after three to six days incubation at 37 °C. Serum titres are considered negative if there is no neutralisation at a dilution of 1/2 (undiluted serum).
B. Any other tes	t recognised in the framework of Decision 2004/558/EC (1).
I	Foot-and-mouth disease (FMD)
A. Collecting oes	ophageal/pharyngeal samples and testing shall be carried out

- A. Collecting oesophageal/pharyngeal samples and testing shall be carried out according to the following protocol:
  - Reagents: Prior to sampling, transport medium is prepared. Two ml volumes are dispensed in as many containers as there are animals to be sampled. The containers used must withstand freezing over solid  $CO_2$  or liquid

<sup>(&</sup>lt;sup>1</sup>) OJ L 249, 23.7.2004, p. 20.

nitrogen. Samples are obtained by the use of a specially-designed sputum collector or 'probang'. To obtain a sample the probang cup is passed through the mouth, over the dorsum of the tongue and down into the upper part of the oesophagus. Attempts are made to scrape the surface epithelium of the upper oesophagus and pharynx by movements directed laterally and dorsally. The probang is then withdrawn, if possible after the animal has swallowed. The cup must be full and contain a mixture of mucus, saliva, oesophageal fluid and cellular debris. Care must be taken to ensure that each specimen contains some visible cellular material. Very rough handling which causes bleeding must be avoided. Samples from some animals may be heavily contaminated with ruminal contents. Such samples must be discarded and the mouth of the animal flushed with water, or preferably physiological saline, before repeat sampling.

Treatmentof Each sample collected in the probang cup is examined for quality and 2 ml added to an equal volume of transport medium in a container which can withstand freezing. The containers are tightly closed, sealed, disinfected and labelled. The samples are kept cool (+ 4 °C) and examined within three to four hours or placed over dry ice (- 69 °C) or liquid nitrogen and kept frozen until examined. Between animals the probang is disinfected and washed in three changes of clean water.

Testing for FMD Samples are inoculated into cultures of primary bovine thyroid cell cultures using at least three tubes per sample. Other susceptible cells such as primary bovine or porcine kidney cells can be used but it must be kept in mind that for some strains of FMD virus they are less sensitive. The tubes are incubated at 37 °C on a roller apparatus and examined daily for 48 hours for the presence of a cytopathic effect (CPE). If negative, cultures are blind passaged onto new cultures and re-examined for 48 hours. The specificity of any CPE must be confirmed.

Recommended transport media:

- 1. 0,08M phosphate buffer pH 7,2 containing 0,01% bovine serum albumin, 0,002% phenol red and antibiotics.
- 2. Tissue culture medium (such as Eagle's MEM) containing 0,04 M Hepes buffer, 0,01 % bovine serum albumin and antibiotics, pH 7,2.
- Antibiotics (per ml final) must be added to the transport medium such as penicillin 1 000 IU, neomycin sulphate100 IU, polymyxin B sulphate50 IU, mycostatin100 IU.

B. The virus neutralisation test shall be carried out according to the following protocol:

Reagents: Stock FMDV antigen is prepared in cell cultures or on cattle tongues and stored at - 70 °C or less or at - 20 °C after the addition of 50 % glycerol. This is the stock antigen. FMDV is stable under these conditions and titres vary little over a period of months.

- Procedure: The test is carried out in flat-bottomed tissue culture grade microtitre plates using susceptible cells such as IB-RS-2, BHK-21 or calf kidney cells. Sera for the test are diluted 1/4 in serum-free cell culture medium with the addition of 100 IU/ml neomycin or other suitable antibiotics. Sera are inactivated at 56 °C for 30 minutes and 0.05 ml amounts are used to prepare a twofold series on microtitre plates using 0,05 ml diluting loops. Pre-titrated virus also diluted in serum-free culture medium and containing 100 TCID50/0.05 ml is then added to each well. Following incubation at 37 °C for one hour to allow neutralisation to take place, 0,05 ml of suspension cells containing 0,5 to  $1.0 \times 10^6$  cells per 1 ml in cell culture medium containing serum free of FMD antibody is added to each well and the plates are sealed. Plates are incubated at 37 °C. Monolayers are normally confluent within 24 hours. CPE is usually sufficiently advanced at 48 hours for a microscopic reading of the test. At this time a final microscopic reading may be made or the plates may be fixed and stained for macroscopic reading, for instance using 10 % formol-saline and 0,05 % methylene blue.
- Controls: Controls in each test include homologous antiserum of known titre, a cell control, a serum toxicity control, a medium control, and a virus titration from which the actual amount of virus in the test is calculated.
- Interpretation: Wells with evidence of CPE are considered to be infected and neutralisation titres are expressed as the reciprocal of the final dilution of serum present in the serum/virus mixtures at the 50 % end point estimated according to the Spearman-Karber method. (Karber, G., 1931, Archiv fuer Experimentelle Pathologie und Pharmokologie, 162, 480.). Tests are considered to be valid when the actual amount of virus used per well in the test is between 101,5 and 102,5 TCID50 and when the titre of the reference serum is within twofold of its expected titre, estimated from the mode of previous titrations. When the controls are outside these limits the tests are repeated. An end point titre of 1/11 or less is taken as negative.

- C. The detection and quantification of antibody by ELISA shall be carried out according to the following protocol:
  - Rabbit antisera to 146S antigen of seven types of foot-Reagents: and-mouth disease virus (FMDV) used at a predetermined optimum concentration in carbonate/bicarbonate buffer, pH 9,6. Antigens are prepared from selected strains of virus grown on monolayers of BHK-21 cells. The unpurified supernatants are used and pretitrated according to the protocol but without serum, to give a dilution which after the addition of an equal volume of PBST (phosphate buffered saline containing 0,05 % Tween-20 and phenol red indicator) would give an optical density reading of between 1,2 and 1,5. The viruses can be used inactivated. PBST is used as a diluent. Guinea-pig antisera are prepared by inoculating guinea pigs with 146S antigen of each serotype. A predetermined optimum concentration is prepared in PBST containing 10 % normal bovine serum and 5 % normal rabbit serum. Rabbit antiguinea-pig immunoglobulin conjugated to horseradish peroxidase is used at a predetermined optimum concentration in PBST containing 10 % normal bovine serum and 5 % normal rabbit serum. Test sera are diluted in PBST.

#### Procedure:

- 1. ELISA plates are coated with 50 μl of rabbit antiviral sera overnight in a humidity chamber at room temperature.
- 2. Fifty microlitres of a duplicate, twofold series of each test serum starting at 1/4 are prepared in U-bottomed multiwell plates (carrier plates). Fifty microlitres of a constant dose of antigen are added to each well and the mixtures are left overnight at 4 °C. The addition of the antigen reduces the starting serum dilution to 1/8.
- 3. The ELISA plates are washed five times with PBST.
- 4. Fifty microlitres of serum/antigen mixtures are then transferred from the carrier plates to the rabbit-serum-coated ELISA plates and incubated at 37 °C for one hour on a rotary shaker.
- 5. After washing, 50  $\mu$ l of guinea-pig antiserum to the antigen used in point 4 is added to each well. The plates are incubated at 37 °C for one hour a rotary shaker.
- The plates are washed and 50 μl of rabbit anti-guinea-pig immunoglobulin conjugated to horseradish peroxidase is added to each well. The plates are incubated at 37 °C for one hour on a rotary shaker.
- 7. The plates are washed and 50  $\mu l$  of orthophenylene diamine containing 0,05 % H\_2O\_2 (30 %) w/v is added to each well.
- 8. The reaction is stopped after 15 minutes with 1,25M H<sub>2</sub>SO<sub>4</sub>.

The plates are read spectrophotometrically at 492 nm on an ELISA reader linked to a microcomputer.

	Controls:	For each antigen used 40 wells contain no serum but contain antigen diluted in PBST. A duplicated twofold dilution series of homologous bovine reference antiserum. A duplicate twofold dilution series of negative bovine serum.
	Interpretation:	Antibody titres are expressed as the final dilution of tests serum giving $50\%$ of the mean OD value recorded in the virus control wells where test serum is absent. Titres in excess of $1/40$ are considered positive.
	References:	Hamblin C, Barnett ITR and Hedger RS (1986) 'A new enzyme-linked immunosorbent assay (ELISA) for the detection of antibodies against foot-and-mouth disease virus. I. Development and method of ELISA.' Journal of Immunological Methods, 93, 115 to 121.11.
	А	ujeszky's disease (AJD)
A.	The serum neutralis protocol:	sation test shall be carried out according to the following

- Serum: All sera are heat-inactivated at 56 °C for 30 minutes before use.
- Procedure: The constant virus-varying serum neutralisation test on microtitre plates employs Vero or other sensitive cell systems. Aujeszky's disease virus is used at 100 TCID50 per 0,025 ml; inactivated undiluted serum samples are mixed with an equal volume (0,025 ml) of virus suspension. The virus/serum mixtures are incubated for two hours at 37 °C in the microtitre plates before the appropriate cells are added. Cells are used at a concentration which forms a complete monolayer after 24 hours.
- Controls: (i) virus infectivity assay, (ii) serum toxicity controls, (iii) uninoculated cell culture controls, (iv) reference antisera.
- Interpretation: The results of the neutralisation test and the titre of the virus used in the test are recorded after three to seven days incubation at 37 °C. Serum titres less than 1/2 (undiluted serum) are considered negative.
- B. Any other test recognised in the framework of Decision 2008/185/EC (1).

### Transmissible gastro-enteritis (TGE)

The serum neutralisation test shall be carried out according to the following protocol:

- Serum: All sera are heat-inactivated at 56 °C for 30 minutes before use.
- Procedure: The constant virus-varying serum neutralisation test on microtitre plates employs A72 (dog tumour) cells or other sensitive cell systems. TGE virus is used at 100 TCID50 per 0,025 ml; inactivated undiluted serum samples are mixed with

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(1) OJ L 59, 4.3.2008, p. 19.

an equal volume (0,025 ml) of virus suspension. The virus/serum mixtures are incubated for 30 to 60 minutes at 37 °C in the microtitre plates before the appropriate cells are added. Cells are used at a concentration which forms a complete monolayer after 24 hours. Each cell receives 0,1 ml of cell suspension.

Controls: (i) virus infectivity assay, (ii) serum toxicity controls, (iii) uninoculated cell culture controls, (iv) reference antisera.

Interpretation: The results of the neutralisation test and the titre of the virus used in the test are recorded after three to five days incubation at 37 °C. Serum titres less than 1/2 (final dilution) are considered negative. If undiluted serum samples are toxic to the tissue cultures, these sera may be diluted 1/2 before being used in the test. This is equivalent to 1/4 final dilution of serum. Serum titres of less than 1/4 (final dilution) are considered negative in these cases.

#### Swine vesicular disease (SVD)

Tests for swine vesicular disease (SVD) shall be carried out according to Decision 2000/428/EC (<sup>1</sup>).

#### Classical swine fever (CSF)

Tests for classical swine fever (CSF) shall be carried out according to Decision 2002/106/EC (<sup>2</sup>).

The performance of tests for CSF must follow the guidelines set out in the relevant chapter of the OIE Manual of Diagnostic Tests and Vaccines for Terrestrial Animals.

The evaluation of sensitivity and specificity of the serological test for CSF must be carried out in a national laboratory with a quality assurance scheme in place. Tests employed must be shown to recognise a range of weak and strong positive reference sera and allow detection of antibody in early phase and convalescence.

#### PART 7

Import and quarantine animal health conditions for animals imported into St. Pierre and Miquelon within a period of less than six months prior to introduction into the Union

(referred to in Article 6)

Animal species covered

Taxon								
ORDER	FAMILY	GENUS AND SPECIES						
Artiodactyla	Camelidae	Camelus spp., Lama spp., Vicugna spp.						

(<sup>1</sup>) OJ L 167, 7.7.2000, p. 22.

<sup>(&</sup>lt;sup>2</sup>) OJ L 39, 9.2.2002, p. 71.

### CHAPTER 1

#### **Residence** and quarantine

- 1. Animals imported into St. Pierre and Miquelon must reside in an authorised quarantine station for a minimum period of 60 days before being dispatched for introduction into the Union. This period may be increased due to testing requirements for individual species. In addition the animals must comply with the following requirements:
  - (a) Separate consignments may enter the quarantine station. However, upon entry in the quarantine station all animals of the same species in the quarantine facility must be considered as a single group, and referred to as such. The quarantine period must commence for the whole group at the time when the last animal entered the quarantine facility.
  - (b) Within the quarantine station each specific group of animals must be maintained in isolation, with no direct or indirect contact with any other animals, including those from other consignments that may be present.

Each consignment must be kept in the approved quarantine station and protected from vector insects.

- (c) If, during the period of quarantine, the isolation of a group of animals is not maintained and contact is made with other animals, the quarantine period must begin again for the same duration as initially prescribed on entry into the quarantine station.
- (d) Animals to be introduced into the Union which pass through the quarantine station must be loaded and dispatched directly to the Union:
  - without coming into contact with animals other than animals which fulfil the health conditions established for the introduction of the relevant category of animal into the Union;
  - segregated into consignments so that no consignment can came in contact with animals not eligible for importation into the Union;
  - (iii) in transport vehicles or containers which have first been cleansed and disinfected with a disinfectant officially authorised in St. Pierre and Miquelon as effective in the control of the diseases referred to in Chapter 2 and which are so constructed that faeces, urine, litter or fodder cannot flow or fall out of the vehicle or container during transportation.
- The quarantine premises must at least meet the minimum standards laid down in Annex B to Directive 91/496/EEC (<sup>1</sup>), and the following conditions:
  - (a) they must be supervised by an official veterinarian;
  - (b) they must be situated at the centre of an area of at least 20 km in diameter in which, according to official findings, for at least 30 days prior to their use as a quarantine station there has been no case of footand-mouth disease;

<sup>(1)</sup> OJ L 268, 24.9.1991, p. 56.

- (c) they must, before being used as a quarantine station, be cleansed and disinfected with a disinfectant officially authorised in St Pierre et Miquelon as effective in the control of the diseases referred to in Chapter 2;
- (d) they must operate, taking into account their animal capacity:
  - a facility dedicated exclusively for the quarantine of animals, including adequate housing to a suitable standard for the animals;
  - (ii) appropriate facilities, that:
    - are easy to thouroughly clean and disinfect,
    - include facilities for safe loading and unloading,
    - are able to fulfil all watering and feeding requirements for the animals,
    - allow any necessary veterinary treatment to be easily administered;
  - (iii) appropriate facilities for inspection and isolation;
  - (iv) appropriate equipment for cleaning and disinfecting rooms and transport vehicles;
  - (v) an appropriate storage area for fodder, litter and manure;
  - (vi) an appropriate system for collecting waste water;
  - (vii) an office for the official veterinarian;
- (e) when operating, they must have sufficient veterinarians to carry out all duties;
- (f) they must only admit animals that are individually identified so as to guarantee traceability. To this end, when animals are admitted the owner or the person in charge of the quarantine station must ensure that the animals are properly identified and accompanied by health certificates for the species and categories involved. In addition, the owner or the person in charge of the quarantine station must record on a register or in a data base, and retain for at least three years, the name of the owner, the origin of the animals in the consignment, the dates of entry and exit of the animals in the consignment, the identification number of the animals in the consignment and their place of destination;
- (g) the competent authority must determine the procedure for official supervision of the quarantine station and must ensure that such supervision is carried out; this supervision must include regular inspections in order to ascertain that the requirements for approval continue to be fulfilled. In case of failure and suspension, the approval may only be restored when the competent authority is satisfied that the quarantine premises are in full compliance with all the conditions set out in points (a) to (g).

#### CHAPTER 2

#### Animal health tests

1. GENERAL REQUIREMENTS

The animals must be subjected to the following tests carried out on samples of blood taken, if not specified otherwise, not earlier than 21 days from the date of commencement of the isolation period.

The laboratory tests must be carried out in an approved laboratory in the Union and all laboratory tests and their results, vaccinations and treatments must be enclosed with the health certificate.

In order to keep animal interventions to a minimum, sampling, tests and any vaccinations must be grouped as far as is possible whilst respecting the minimum time intervals required by the testing protocols set out in Part 2 of this Chapter.

#### 2. SPECIFIC REQUIREMENTS

#### 2.1 CAMELIDAE

- 2.1.1 Tuberculosis
  - (a) Test to be used: comparative intradermal reaction test using Bovine purified protein derivative (PPD) and Avian PPD conforming to the standards for the manufacture of bovine and avian tuberculins as described in point 2.1.2 of Annex B of Directive 64/432/EEC.

The test must be executed in the area behind the shoulder (axillary region) following the technique described in point 2.2.4 of Annex B of Directive 64/432/EEC.

(b) **Timing**: the animals must be tested within two days from the date of arrival in the quarantine station and 42 days from the date of the first test.

#### (c) Interpretation of tests:

the reaction shall be considered:

- negative if the increased skin thickness is less than 2 mm.
- positive if the increased skin thickness is more than 4 mm.
- inconclusive if the increased skin thickness to the bovine PPD is between 2mm and 4 mm, or more than 4 mm but less then the reaction to the avian PPD.

#### (d) Options for action following testing:

If an animal presents a positive result to the intradermal-reaction to the bovine PPD, that animal shall be excluded from the group and the other animals shall be re-tested starting at least 42 days from the date of the first positive test was administered and this shall be considered as the first test described in (b).

If more than one animal of the group presents a positive result, the whole group shall be rejected for exportation to the Union.

If one or more animals of the same group present an inconclusive reaction, the whole group shall be re-tested starting at least 42 days from the date of the first test was administered and it shall be considered as the first test described in (b).

#### 2.1.2 Brucellosis

#### (a) Test to be used:

- (i) Brucella abortus: Rose Bengal test (RBT) and Serum agglutination test (SAT) as described respectively in points 2.5 and 2.6 of Annex C to Directive 64/432/EEC. In the case of a positive result, a complement-fixation test shall be performed for confirmation as described in Part 6 of Annex I to Regulation (EU) No 206/2010.
- (ii) Brucella melitensis: RBT and SAT as described respectively in points 2.5 and 2.6 of Annex C to Directive 64/432/EEC. In the case of a positive result, a complement-fixation test following the method described in Annex C to Directive 91/68/EEC shall be performed for confirmation.
- (iii) *Brucella ovis*: Complement fixation test as described in Annex D to Directive 91/68/EEC
- (b) **Timing**: the animals have to be tested within two days from the date of their arrival in the quarantine station and 42 days from the date of the first test.

#### (c) Interpretation of tests:

A positive reaction to the tests shall be as defined in Annex C to Directive 64/432/EEC.

#### (d) Options for action following testing:

Animals tested positive to one of the tests shall be excluded from the group and the other animals shall be re-tested starting at least 42 days from the date the first positive test was performed: this shall be considered as the first test described in (b).

Only the animals that tested negative to two consecutive tests performed as described in (b) shall be allowed for the introduction to the Union.

#### 2.1.3 Bluetongue and Epizootic haemorrhagic disease (EHD)

(a) **Test to be** used: agar gel immunodiffusion (AGID) test as described in Part 6 of Annex I to Regulation (EU) No 206/2010.

In case of a positive reaction the animals shall be tested with competitive ELISA test as described in Part 6 of Annex I to Regulation (EU) No 206/2010 to discriminate between the two diseases.

### (b) Timing:

The animals must be tested with negative result to two tests: the first within two days from the date of their arrival in the quarantine station and the second at least 21 days from date of the first test.

#### (c) Options for action following testing:

(i) Bluetongue

If one or more animals tested positive to the ELISA as described in Part 6 of Annex I to Regulation (EU) No 206/2010, the positive animal/animals shall be excluded from the group, and all the remaining animals in the group must be quarantined for 100 days starting from the date on which the samples for the positive test were collected. The group shall only be considered free of the bluetongue disease if regular checks carried out by official veterinarians throughout the duration of the quarantine period fail to reveal clinical symptoms of disease, and the quarantine station remains free of bluetongue vectors (*Culicoides*).

If a further animal presents clinical symptoms of bluetongue disease during the quarantine period as described in the first subparagraph, all the animals in the group shall be rejected for introduction into the Union.

(ii) Epizootic haemorrhagic disease (EHD)

If one or more animals tested positive reveal the presence of antibodies to the EHD virus during confirmatory ELISA testing, the animal(s) shall be considered positive and shall be excluded from the group, and the whole group shall be subject to repeat testing beginning at least 21 days from the date of the initial positive diagnosis and again at least 21 days from the date of the repeat test, both with negative results.

If any additional animals are tested positive during either or both of the two tests carried out for repeat testing, the whole group of animals shall be rejected for introduction into the Union.

#### 2.1.4 Foot-and-Mouth Disease (FMD)

- (a) Test to be used: Diagnostic tests (probang and serology) using ELISA and (Virus Neutralisation) (VN) techniques in accordance with the Protocols described in Part 6 of Annex I to Regulation (EU) No 206/2010.
- (b) **Timing**: the animals shall be tested with negative results to two tests: the first within two days from the date of their arrival in the quarantine station and the second at least 42 days from the date of the first test.
- (c) Options for action following testing: If any animal tests positive for the FMD virus, then none of the animals present in the quarantine station shall be considered as eligible for introduction into the Union.

**Note:** Any detection of antibodies to structural or not structural proteins of FMD virus shall be considered as a result of previous infection of FMD irrespective of the vaccination status.

#### 2.1.5 Rinderpest

- (a) Test to be used: The competitive ELISA test as described in the OIE Manual of Diagnostic Tests and Vaccines for Terrestrial Animals, latest version, is the prescribed test for international trade and is test of choice. Serum neutralisation test, or other recognised tests in accordance with the protocols described in relevant sections of the OIE manual may also be used.
- (b) **Timing**: the animals shall be tested twice: the first within two days from the date of their arrival in the quarantine station and the second at least 42 days from the date of the first test.
- (c) Options for action following testing: If any animal tests positive for the Rinderpest virus, then none of the animals present in the quarantine station shall be considered as eligible for introduction into the Union.

### 2.1.6 Vesicular stomatitis

- (a) **Test to be used**: ELISA, virus neutralisation test, or other recognised test in accordance with the protocols described in the relevant sections of the OIE manual.
- (b) Timing: the animals shall be tested twice: the first within two days from the date of their arrival in the quarantine station and the second at least 42 days from the date of the first test.
- (c) Options for action following testing: If any animal tests positive for vesicular stomatitis virus, then none of the animals present in the quarantine station shall be considered as eligible for introduction into the Union.

#### 2.1.7 Rift valley fever

- (a) **Test to be used**: ELISA, virus neutralisation test, or other recognised test in accordance with the protocols described in relevant sections of the OIE manual.
- (b) **Timing**: the animals shall be tested twice: the first within two days from the date of their arrival in the quarantine station and the second at least 42 days from the date of the first test.
- (c) Options for action following testing: If any animal displays evidence of exposure to rift valley fever agent, then none of the animals present in the quarantine station shall be considered as eligible for introduction into the Union.

#### 2.1.8 Lumpy skin disease

- (a) Test to be used: Serology using ELISA, virus neutralisation test, or other recognised test in accordance with the protocols described in relevant sections of the OIE manual.
- (b) **Timing**: the animals shall be tested twice: the first within two days from the date of their arrival in the quarantine station and the second at least 42 days from the date of the first test.

- (c) Options for action following testing: If any animal displays evidence of exposure to lumpy skin disease, then none of the animals present in the quarantine station shall be considered as eligible for introduction into the Union.
- 2.1.9 Crimean congo haemorrhagic fever
  - (a) **Test to be used**: ELISA, virus neutralisation test, Immunofluorescence test or other recognised test.
  - (b) **Timing**: the animals shall be tested twice: the first within two days from the date of their arrival in the quarantine station and the second at least 42 days from the date of the first test.
  - (c) **Options for action following testing**: If any animal displays evidence of exposure to crimean congo haemorrhagic fever agent, then none of the animals present in the quarantine station shall be considered as eligible for introduction into the Union.
- 2.1.10 Surra (Trypanosoma evansi (T. evansi))
  - (a) **Test to be used**: The parasitic agent can be identified in concentrated blood samples in accordance with the protocols described in relevant sections of the OIE manual.
  - (b) Timing: the animals shall be tested twice: the first within two days from the date of their arrival in the quarantine station and the second at least 42 days from the date of the first test.
  - (c) Options for action following testing: If *T. evansi* is detected in any animal in the consignment, then that animal shall be considered not eligible for introduction into the Union. The remaining animals of the group shall then undergo internal and external antiparasitic treatment using suitable agents that are effective against *T. evansi*.
- 2.1.11 Malignant catarrhal fever
  - (a) Test to be used: Detection of viral DNA based on identification by immunofluorescence or immunocytochemistry using the protocols described in relevant sections of the OIE manual.
  - (b) **Timing**: the animals shall be tested twice: the first within two days from the date of their arrival in the quarantine station and the second at least 42 days from the date of the first test.
  - (c) **Options for action following testing**: If any animal displays evidence of exposure to MCF, then none of the animals present in the quarantine station shall be considered as eligible for introduction into the Union.
- 2.1.12 Rabies

**Vaccination**: Rabies vaccination may be carried out when requested by the Member State of destination and the animal shall be blood sampled and a serum neutralisation test for antibodies carried out.

- 2.1.13 *Enzootic bovine leucosis.* (only in the case where the animals are destined for an officially enzootic-bovine-leucosis free Member State or region, as referred to in Article 2(2)(k) of Directive 64/432/EEC)
  - (a) **Test to be used**: AGID or blocking ELISA, in accordance with the protocols described in the OIE manual, latest version.

- (b) **Timing**: the animals shall be tested twice: the first within two days from the date of their arrival in the quarantine station and the second at least 42 days from the date of the first test.
- (c) **Options for action following testing**: animals tested positive to the test described in (a) shall be excluded from the group of animals in the quarantine facility and the other animals shall be re-tested starting at least 21 days from the date of the first positive test was performed: this shall be considered as the first test described in (b).

Only the animals that tested negative to two consecutive tests performed as described in (b) shall be considered eligible for introduction into the Union.

▼<u>M2</u>

### ANNEX II

# FRESH MEAT

# PART 1

# List of third countries, territories and parts thereof (1)

ISO code and name of	Code of Tomitom	ode of Territory Description of third country, territory or part thereof	Veterinary certificate		Specific	Classing data (2)	Omening data (3)
third country	Code of Territory	Description of third country, territory of part thereof	Model(s)	SG	conditions	Closing date ( <sup>2</sup> )	Opening date ( <sup>3</sup> )
1	2	3	4	5	6	7	8
AL – Albania	AL-0	Whole country	_				
AR – Argentina	AR-0	Whole country	EQU				
	AR-1	-1 The Provinces of: Buenos Aires, Catamarca, Corrientes (except the departments of Berón de Astrada, Capital, Empedrado, General Paz, Itati, Mbucuruyá, San Cosme and San Luís del Palmar) Entre Ríos,	BOV	A	1		18 March 2005
	La Rioja, Mendoza, Misiones, Part of Neuquén (excluding territory included in AR Part of Río Negro (excluding territory included in AR San Juan, San Luis, Santa Fe, Tucuman, Cordoba, La Pampa, Santiago del Estero, Chaco, Formosa, Jujuy and Salta, excluding the buffer of 25 Km from the border with Bolivia and Paraguay extends from the Santa Catalina District in the Provinc	Mendoza, Misiones, Part of Neuquén (excluding territory included in AR-4), Part of Río Negro (excluding territory included in AR-4), San Juan, San Luis,	RUF	A	1		1 December 2007
		Tucuman, Cordoba, La Pampa,	RUW	A	1		1 August 2010

▼	Μ	2

1	2	3	4	5	6	7	8
	AR-2	Chubut, Santa Cruz and Tierra del Fuego	BOV, OVI, RUW, RUF				1 March 2002
	AR-3	Corrientes: the departments of Berón de Astrada, Capital, Empedrado, General Paz, Itati, Mbucuruyá, San Cosme and San Luís del Palmar	BOV RUF	А	1		1 December 2007
	AR-4	Part of Río Negro (except: in Avellaneda the zone located north of the Provincial road 7 and east of the Provincial road 250, in Conesa the zone located east of the Provincial road 2, in El Cuy the zone located north of the Provincial road 7 from its intersection with the Provincial road 66 to the border with the Department of Avellaneda, and in San Antonio the zone located east of the Provincial roads 250 and 2) Part of Neuquén (except in Confluencia the zone located east of the Provincial road 17, and in Picun Leufú the zone located east of the Provincial road 17)	BOV, OVI, RUW, RUF				1 August 2008
AU – Australia	AU-0	Whole country	BOV, OVI, POR, EQU, RUF, RUW, SUF, SUW				
BA – Bosnia and Herzegovina	BA-0	Whole country	_				
BH – Bahrain	BH-0	Whole country	_				
BR – Brazil	BR-0	Whole country	EQU				
	BR-1	State of Minas Gerais State of Espírito Santo; State of Goiás; State of Mato Grosso State of Rio Grande Do Sul, State of Mato Grosso Do Sul (except for the designated high surveillance zone of 15 Km from the external borders in the municipalities of Porto Murtinho, Caracol, Bela Vista, Antônio João, Ponta	BOV	A and H	1		1 December 2008

▼ <u>M2</u>								
	1	2	3	4	5	6	7	8
			Porã, Aral Moreira, Coronel Sapucaia, Paranhos, Sete Quedas, Japorã, and Mundo Novo and the designated high surveillance zone in the municipalities of Corumbá and Ladário).					
		BR-2	State of Santa Catarina	BOV	A and H	1		31 January 2008
		BR-3	States of Paraná and São Paulo	BOV	A and H	1		1 August 2008
▼ <u>M7</u>								
	BW — Botswana	BW-0	Whole country	EQU, EQW				
		BW-1	The veterinary disease control zones 3c, 4b, 5, 6, 8, 9 and 18, except the intensive surveillance zone in zone 6 between the border with Zimbabwe and the highway A1	BOV, OVI, RUF, RUW	F	1	11 May 2011	26 June 2012
		BW-2	The veterinary disease control zones 10, 11, 13 and 14	BOV, OVI, RUF, RUW	F	1		7 March 2002
		BW-3	The veterinary disease control zone 12	BOV, OVI, RUF, RUW	F	1	20 October 2008	20 January 2009
		BW-4	The veterinary disease control zone 4a, except the intensive surveillance buffer zone of 10 km along the boundary with the foot-and-mouth disease vaccination zone and wildlife management areas	BOV	F	1		18 February 2011
▼ <u>M2</u>								
	BY – Belarus	BY-0	Whole country					
	BZ – Belize	BZ-0	Whole country	BOV, EQU				

▼M2

▼<u>M2</u>

1	2	3	4	5	6	7	8
CA – Canada	CA-0	Whole country	BOV, OVI, POR, EQU, SUF, SUW, RUF, RUW	G			
CH – Switzerland	CH-0	Whole country	*				
CL – Chile	CL-0	Whole country	BOV, OVI, POR, EQU, RUF, RUW, SUF				
CN – China	CN-0	Whole country	_				
CO – Colombia	CO-0	Whole country	EQU				
CR – Costa Rica	CR-0	Whole country	BOV, EQU				
CU – Cuba	CU-0	Whole country	BOV, EQU				
DZ – Algeria	DZ-0	Whole country	_				
ET – Ethiopia	ET-0	Whole country	_				
FK – Falkland Islands	FK-0	Whole country	BOV, OVI, EQU				
GL – Greenland	GL-0	Whole country	BOV, OVI, EQU, RUF, RUW				
GT – Guatemala	GT-0	Whole country	BOV, EQU				
HK – Hong Kong	НК-0	Whole country	_				
HN – Honduras	HN-0	Whole country	BOV, EQU				
HR – Croatia	HR-0	Whole country	BOV, OVI, EQU, RUF, RUW				
IL – Israel	IL-0	Whole country	_				

1	2	3	4	5	6	7	8
IN – India	IN-0	Whole country	_				
IS – Iceland	IS-0	Whole country	BOV, OVI, EQU, RUF, RUW				
KE – Kenya	KE-0	Whole country	_				
MA – Morocco	MA-0	Whole country	EQU				
ME – Montenegro	ME-0	Whole country	BOV, OVI, EQU				
MG – Madagascar	MG-0	Whole country	_				
MK – Former Yugoslav Republic of Macedonia ( <sup>4</sup> )	МК-0	Whole country	OVI, EQU				
MU – Mauritius	MU-0	Whole country	_				
MX – Mexico	MX-0	Whole country	BOV, EQU				
NA – Namibia	NA-0	Whole country	EQU, EQW				
	NA-1	South of the cordon fences which extend from Palgrave Point in the west to Gam in the east	BOV, OVI,RUF, RUW	F and J	1		
NC – New Caledonia	NC-0	Whole country	BOV, RUF, RUW				
NI – Nicaragua	NI-0	Whole country	_				
NZ – New Zealand	NZ-0	Whole country	BOV, OVI, POR, EQU, RUF, RUW, SUF, SUW				
PA – Panama	PA-0	Whole country	BOV, EQU				

1	2	3	4	5	6	7	8
5							
PY – Paraguay	РҮ-0	Whole country	EQU				
	РҮ-1	Whole country except the designated high surveillance zone of 15 km from the external borders	BOV	А	1	18 September 2011	1 August 2003
2							
RS – Serbia ( <sup>5</sup> )	RS-0	Whole country	BOV, OVI, EQU				
RU – Russia	RU-0	Whole country	_				
	RU-1	Region of Murmansk, Yamolo-Nenets autonomous area	RUF				
SV – El Salvador	SV-0	Whole country	—				
SZ – Swaziland	SZ-0	Whole country	EQU, EQW				
	SZ-1	Area west of the 'red line' fences which extends northwards from the river Usutu to the frontier with South Africa west of Nkalashane,	BOV, RUF, RUW	F	1		
	SZ-2	The veterinary foot and mouth disease surveillance and vaccination control areas as gazetted as a Statutory Instrument under legal notice number 51 of 2001	BOV, RUF, RUW	F	1		4 August 2003
TH – Thailand	TH-0	Whole country	—				
TN – Tunisia	TN-0	Whole country	_				
TR – Turkey	TR-0	Whole country	_				
	TR-1	The provinces of Amasya, Ankara, Aydin, Balikesir, Bursa, Cankiri, Corum, Denizli, Izmir, Kastamonu, Kutahya, Manisa, Usak, Yozgat and Kirikkale	EQU				
UA – Ukraine	UA-0	Whole country	_				

V IVIZ								
	1	2	3	4	5	6	7	8
	US – United States	US-0	Whole country	BOV, OVI, POR, EQU,SUF, SUW, RUF, RUW	G			
	UY – Uruguay	UY-0	Whole country	EQU				
				BOV,	А	1		1 November 2001
				OVI	А	1		
▼ <u>M3</u>								
	ZA - South Africa	ZA-0	Whole country	EQU, EQW				
		ZA-1	<ul> <li>The whole country except:</li> <li>the part of the foot-and-mouth disease control area situated in the veterinary regions of Mpumalanga and Northern provinces, in the district of Ingwavuma of the veterinary region of Natal and in the border area with Botswana east of longitude 28°, and</li> <li>the district of Camperdown, in the province of KwaZulu-Natal.</li> </ul>	BOV, OVI, RUF, RUW	F	1	11 February 2011	
▼ <u>M2</u>								
	ZW – Zimbabwe	ZW-0	Whole country	_				

Footnotes:

(1) Without prejudice to specific certification requirements provided for in Union agreements with third countries.

(2) Meat from animals slaughtered on or before the date set out in column 7 may be imported into the Union for 90 days from that date. Consignments carried on vessels on the high seas may be imported into the Union if certified before the date set out in column 7 for 40 days from that date. (N.B.: no date in column 7 means that there are no time restrictions).

(3) Only meat from animals slaughtered on or after the date set out in column 8 may be imported into the Union (no date in column 8 means that there are no time restrictions).

(4) The former Yugoslav Republic of Macedonia; provisional code that does not prejudge in any way the definitive nomenclature for this country, which will be agreed following the conclusion of negotiations currently taking place on this subject in the United Nations.

(5) Not including Kosovo which is at present under international administration pursuant to United Nations Security Council Resolution 1244 of 10 June 1999

\* = Requirements as in accordance with the Agreement between the European Community and the Swiss Confederation on trade in agricultural products (OJ L 114, 30.4.2002, p. 132).

- = No certificates are laid down and fresh meat imports are prohibited (except for those species where indicated in the line comprising the entry for the whole country).

'1' Category restrictions:

No offal is authorised for introduction into the Union (except, in the case of bovine species, diaphragm and masseter muscles).

#### PART 2

#### Models of veterinary certificates

Model(s):

- <sup>'</sup>BOV': Model of veterinary certificate for fresh meat, including minced meat, of domestic bovine animals (including *Bison* and *Bubalus* species and their cross-breeds).
- 'OVI': Model of veterinary certificate for fresh meat, including minced meat, of domestic ovine animals (*Ovis aries*) and domestic caprine animals (*Capra hircus*).
- 'POR': Model of veterinary certificate for fresh meat, including minced meat, of domestic porcine animals (*Sus scrofa*).
- 'EQU': Model of veterinary certificate for fresh meat, excluding minced meat, of domestic solipeds (*Equus caballus, Equus asinus* and their crossbreeds).
- <sup>(RUF)</sup> Model of veterinary certificate for fresh meat, excluding offal and minced meat, of farmed non-domestic animals of the order Artiodactyla (excluding bovine animals (including *Bison* and *Bubalus* species and their cross-breeds), *Ovis aries, Capra hircus*, Suidae and Tayassuidae), and of the families Rhinocerotidae and Elephantidae.
- 'RUW': Model of veterinary certificate for fresh meat, excluding offal and minced meat, of wild non-domestic animals of the order Artiodactyla (excluding bovine animals (including *Bison* and *Bubalus* species and their cross-breeds), *Ovis aries, Capra hircus*, Suidae and Tayassuidae), and of the families Rhinocerotidae and Elephantidae.
- 'SUF': Model of veterinary certificate for fresh meat, excluding offal and minced meat, of farmed non-domestic animals belonging to the Suidae, Tayassuidae, or Tapiridae families.
- 'SUW': Model of veterinary certificate for fresh meat, excluding offal and minced meat, of wild non-domestic animals belonging to the Suidae, Tayassuidae, or Tapiridae families.
- 'EQW': Model of veterinary certificate for fresh meat, excluding offal and minced meat, of wild solipeds belonging to the subgenus *Hippotigris* (zebra).
- SG (Supplementary guarantees)
- 'A': guarantees regarding the maturation, pH measurement and boning of fresh meat, excluding offal, certified according to the models of veterinary certificates BOV (point II.2.6), OVI (point II.2.6), RUF (point II.2.7) and RUW (point II.2.4).
- <sup>c</sup>C<sup>'</sup>: guarantees regarding the laboratory test for classical-swine-fever in the carcases from which fresh meat was obtained, certified according to the model of veterinary certificate SUW (point II.2.3 B).
- <sup>(D)</sup>: guarantees regarding swill feed on holding(s) of animals from which fresh meat certified was obtained according to the model of veterinary certificate POR (point II.2.3 d).
- 'E': guarantees regarding tuberculosis test in the animals from where fresh meat certified was obtained, according to the model of veterinary certificate BOV (point II.2.4 d).
- 'F': guarantees regarding the maturation and de-boning of fresh meat, excluding offal, certified according to the models of veterinary certificates BOV (point II.2.6), OVI (point II.2.6), RUF (point II.2.6) and RUW (point II.2.7).

- 'G': guarantees regarding 1, exclusion of offals and spinal cord; and 2, testing and origin of cervid animals in relation to chronic wasting disease as referred to in the models of veterinary certificates RUF (point II.1.7) and RUW (point II.1.8).
- <sup>c</sup>H<sup>2</sup>: supplementary guarantees required for Brazil. Concerning vaccination programmes, as the State of Santa Catarina in Brazil does not vaccinate against foot and mouth disease, the reference to a vaccination programme is not applicable for meat coming from animals originating and slaughtered in that State.
- <sup>c</sup>J<sup>2</sup>: guarantees regarding the movement of bovine, ovine and caprine animals from holdings to the slaughterhouse, which allow them to pass via an assembly centre (including markets) before being transported directly to slaughter.

## Model BOV

cou	NTRY					Veterinary certificate to EL				
	l.1.	Consignor		I.2. Certificat	te reference No	l.2.a.				
		Name		I.3. Central competent authority						
		Address			-	·				
t		Tel.		I.4. Local co	mpetent authority					
dispatched consignment	l.5.	Consignee		I.6.						
onsiç		Name								
o pa		Address								
atch		Postal code								
disp		Tel.								
ls of	1.7.	Country of origin ISO code	I.8. Region of origin Code	I.9. Country destination		I.10. Region of Code destination				
Detai				destinati						
Part I: Details of	l.11.	Place of origin		l.12.						
Ра		Name	Approval number							
		Address								
	113	Place of loading		I.14. Date of c	leparture					
		That's of fouring								
	l.15.	Means of transport		I.16. Entry BIF	° in EU					
		Aeroplane Ship Ship Road vehicle Other								
		Identification		l.17.						
		Documentary references								
	1.18.	Description of commodity			I.19. Commodity	code (HS code)				
						I.20. Quantity				
	1.21.	Temperature of product				I.22. Number of packages				
		Ambient 🔲	Chilled	Frozen 🔲						
	1.23.	Seal/Container No				I.24. Type of packaging				
	1.25.	Commodities certified for:								
		Human consumption 🗖								
	1.26.			I.27. For impo	rt or admission ir	nto EU				
	1.28	Identification of the commodities								
		Species Nature	of Treatment	Annroval numbe	r of establishmen	ts Number of Net				
		(scientific name) commod				packages weight				
			Aban	earaing	, p.a 001					

	COUNT	RY			Model BOV						
	11.	Health information		II.a. Certificate reference number	II.b.						
	II.1.	Public Health Attestation	n								
		(EC) No 852/2004, (EC) I	al veterinarian declare that I am aw No 853/2004, (EC) No 854/2004 and roduced in accordance with those req	(EC) No 999/2001 and certify that the							
Part II: Certification	II.1.1.	<ol> <li>the [meat] [minced meat] (<sup>1</sup>) comes from (an) establishment(s) implementing a programme based on the HACCP principles in accordar with Regulation (EC) No 852/2004;</li> </ol>									
II: Cer	II.1.2.	the meat has been obtained in compliance with Section I of Annex III to Regulation (EC) No 853/2004;									
Part			t has been produced in compliance wit ture of not more than – 18 °C;]	th Section V of Annex III to Regulation	(EC) No 853/2004 and frozen to an						
			n found fit for human consumption foll on I and Chapters I and IX of Section								
			cass or parts of the carcass have beer to Regulation (EC) No 854/2004;]	n marked with a health mark in accord	ance with Chapter III of Section I of						
			kages of [meat] [minced meat] ( <sup>1</sup> ) have I to Regulation (EC) No 853/2004;]	e been marked with an identification m	ark in accordance with Section I of						
II.1.6. the [meat] [minced meat] ( <sup>1</sup> ) satisfies the relevant criteria set out in Regulation (EC) No 2073/2005 on microbiol foodstuffs;											
II.1.7. the guarantees covering live animals and products thereof provided by the residue plans submitted in accordanc 96/23/EC, and in particular Article 29 thereof, are fulfilled;											
			meat] ( <sup>1</sup> ) has been stored and transp nex III to Regulation (EC) No 853/2004		requirements of Sections I and V						
		II.1.9. with regard to bovi	ne spongiform encephalopathy (BSE):								
		( <sup>1</sup> ) <i>either</i> [II.1.9.1	. for imports from a country or a 2007/453/EC:	region with a negligible BSE risk	and listed as such in Decision						
			<ul> <li>(a) the country or region is classified country or region posing a negling</li> </ul>	ed in accordance with Article 5(2) of ligible BSE risk;	Regulation (EC) No 999/2001 as a						
			(b) the animals from which the bovi slaughtered in a country with a	ne meat or minced meat was derived negligible BSE risk ( <sup>13</sup> );	were born, continuously reared and						
		(	<sup>1</sup> ) [(c) if in the country or region there	have been BSE indigenous cases:							
<ul> <li>(<sup>1</sup>) <i>either</i> [the animals were born after the date from which the ban on the feeding of ruminan and-bone meal and greaves derived from ruminants had been enforced.]</li> <li>(<sup>1</sup>) <i>or</i> [the bovine meat or minced meat does not contain and is not derived from specified as defined in Annex V to Regulation (EC) No 999/2001, or mechanically sep obtained from bones of bovine animals.]]]</li> </ul>											
										( <sup>1</sup> ) or [II.1.9.2.	for imports from a country or a 2007/453/EC:
			<ul> <li>(a) the country or region is classifie country or region posing a contri</li> </ul>	ed in accordance with Article 5(2) of I rolled BSE risk;	Regulation (EC) No 999/2001 as a						

II.	Health information		II.a. Certificate reference number	II.b.
		(b) the animals from which the h	bovine meat or minced meat was deriv	
		stunning by means of gas inje	ected into the cranial cavity or killed by central nervous tissue by means of a	the same method or slaughtered by
		( <sup>1</sup> ) either [(c) the bovine meat or minced defined in Annex V to Regu bones of bovine animals.]	meat does not contain and is not deri Ilation (EC) No 999/2001, or mechanic	
		quarters contain no specific ganglia. The carcasses or	es or half carcasses cut into no mo ed risk material other than the verte wholesale cuts of carcasses of bo ied by a blue stripe on the label	bral column, including dorsal roc ovine animals containing vertebra
	( <sup>1</sup> ) or [II.1.9.3.	for imports from a country or a region whic (EC) No 999/2001 or has been categorised Decision 2007/453/EC:		
		(a) the country or region has not been cate has been categorised as a country or		of Regulation (EC) No 999/2001 c
		<ul><li>(b) the animals from which the bovine mea greaves derived from ruminants;</li></ul>	at or minced meat was derived have no	ot been fed meat-and-bone meal o
		<ul> <li>(c) the animals from which the bovine meal means of gas injected into the cranial stunning of central nervous tissue by r cavity;</li> </ul>	I cavity or killed by the same method	or slaughtered by laceration after
	( <sup>1</sup> ) eithe	r [(d) the bovine meat or minced meat was	not derived from:	
		(i) specified risk material as defined i	n Annex V to Regulation (EC) No 999	/2001;
		(ii) nervous and lymphatic tissues exp	posed during the deboning process;	
		(iii) mechanically separated meat obta	ined from bones of bovine animals.]	
	( <sup>1</sup> ) or	[(d) the carcasses, half carcasses or half c no specified risk material other than wholesale cuts of carcasses of bovir stripe on the label referred to in Regu	the vertebral column, including dors ne animals containing vertebral colum	al root ganglia. The carcasses c
	Parli	Ifils the requirements of Regulation (EC) No lament and of the Council as regards spec den of certain meat and eggs;]		
II.2.	Animal Health att	estation		
	I, the undersigned	official veterinarian, hereby certify, that the t	fresh meat described in Part I:	
	II.2.1. has be	en obtained in the territory/ies with code:	( <sup>2</sup> ) which, a	t the date of issuing this certificate
		is been free for 12 months from rinderpest, a ace, and	and during the same period no vaccina	tion against this disease has take
		is been free for 12 months from foot-and-mout is taken place;]	th disease, and during the same period	no vaccination against this disease
		s been considered free from foot-and-mouth erwards, and authorised to export this meat		

COUN	ITRY			Model BOV				
П.	Health info	ormation	II.a. Certificate reference number	II.b.				
	( <sup>1</sup> ) ( <sup>5</sup> ) or	[(b) vaccination programmes against foot animals;]	and-mouth disease are being officially carried ou	t and controlled in domestic bovine				
	( <sup>1</sup> ) ( <sup>6</sup> ) or	vaccination programme is controlled	amme against foot and mouth disease and from I by the competent veterinary authority through and which also demonstrates the absence of foot	a regular serological surveillance				
	( <sup>1</sup> ) ( <sup>6</sup> ) or		lled by the competent veterinary authority	th disease, and during the same period no vaccination against this disease the competent veterinary authority through a regular surveillance infection;]				
	II.2.2.	has been obtained from animals that:						
		( <sup>1</sup> ) <i>either</i> [have remained in the territor slaughter;]	y described under point II.2.1 since birth, or for a	t least the last three months before				
			(dd/mm/yyyy) into the territory des $(^2)$ that at that date was authorised to imp					
		( <sup>1</sup> ) or [have been introduced on Member State	(dd/mm/yyyy) into the territory descr ;].	ibed under point II.2.1, from the EU				
	II.2.3.	has been obtained from animals coming	from holdings in which:					
		(a) None of the animals present therein	n have been vaccinated against [foot-and-mouth o	disease or] ( <sup>7</sup> ) rinderpest, and				
	( <sup>1</sup> ) either	[(b) in these holdings, and in the holding mouth disease or rinderpest during	is situated in their vicinity within 10 km, there has the previous 30 days,]	been no case/outbreak of foot-and-				
	( <sup>1</sup> ) ( <sup>8</sup> ) or		mal health reasons and where, in these holdings on no case/outbreak of foot-and-mouth disease o					
		(c) they have remained for at least 40	days before direct dispatch to the slaughterhouse	ə;]				
	( <sup>1</sup> ) ( <sup>14</sup> ) or		0 days before passing through one assembly c into contact with animals of a different health					
	( <sup>1</sup> ) ( <sup>9</sup> ) or		mal health reasons and where, in these holdings on no case/outbreak of foot-and-mouth disease o					
		(c) they have remained for at least 40	days before direct dispatch to the slaughterhous	ə;]				
	(1) (6)	[(d) animals have not been introduced of	during the last 3 months from areas not approved	by the EU;				
		(e) animals are identified and registered	I in the national System of Identification and Certif	ication of Origin for bovine animals;				
		official report, in TRACES (10) and	as approved holdings, following a favourable co inspections are regularly carried out by the comp n Regulation (EU) No 206/2010 are respected.]					
	II.2.4. has	s been obtained from animals which:						
	(a)		in vehicles, cleaned and disinfected before loadi did not comply with the conditions referred to in p					

	RY				Model BO
II.	Health	info	ormatior	on II.a. Certificate reference number II.b.	
		(b)		e slaughterhouse, have passed ante-mortem health inspection during the 24 hours before slaughter and, in p on no evidence of the diseases referred to in point II.2.1,	oarticular, have
		(c)		been slaughtered on (dd/mm/yyyy) or between (dd/mm/yyyy) (mm/yyyy) ( <sup>11</sup> );	/y) and
	(1) (12)	[(d)	have	e reacted negatively to an official intra-dermal tuberculosis test carried out within 3 months before slaughte	ər;]
	(1) (6)	[(e)		e slaughterhouse have been kept prior to slaughter completely separate from animals the meat of which is n Union].	ot intended for
	II.2.5.	refe imp	rred to ortation	n obtained in an establishment around which, within a radius of 10 km, there has been no case/outbreak of to in point II.2.1 during the previous 30 days or, in the event of a case/outbreak of disease, the preparati on to the Union has been authorised only after slaughter of all animals present, removal of all meat, and the fection of the establishment under the control of an official veterinarian;	on of meat for
	II.2.6.				
		(1) 6	either	[has been obtained and prepared without contact with other meats not complying with the conditions r certificate.]	required in this
		( <sup>1</sup> ) ( <sup>1</sup>	<sup>3</sup> ) or	[contains [boneless meat] [and] [minced meat] ( <sup>1</sup> ), obtained only from de-boned meat other than offal that from carcasses in which the main accessible lymphatic glands have been removed, which have been maturation at a temperature above + 2 °C for at least 24 hours before the bones were removed and in value of the meat was below 6.0 when tested electronically in the middle of the longissimus-dors maturation and before de-boning, and	n submitted to which the pH
				has been kept strictly separate from meat not conforming to the requirements referred to in this certifi stages of its production, de-boning and storage until it has been packed in boxes or cartons for furt dedicated areas.]	
		( <sup>1</sup> ) (	( <sup>9</sup> ) or	[contains [boneless meat] [and] [minced meat] ( <sup>1</sup> ), obtained only from de-boned meat other than offal tha from carcasses in which the main accessible lymphatic glands have been removed, which have been maturation at a temperature above + 2 °C for at least 24 hours before the bones were removed, and	
				has been kept strictly separate from meat not conforming to the requirements referred to in this certifi stages of its production, de-boning and storage until it has been packed in boxes or cartons for furt dedicated areas.]	
II.3.	Anima	al we	elfare a	attestation	
				ed official veterinarian, hereby certify, that the fresh meat described in Part I derives from animals which hav house before and at the time of slaughter or killing in accordance with the relevant provisions of Union leg	
Notes					
	ertificate preeds).	is	meant	t for fresh meat, including minced meat, of domestic bovine animals (including Bison and Bubalus spe	ecies and their
Fresh r	meat m	eans	all ani	nimal parts fit for human consumption whether fresh, chilled or frozen.	
Part I					
— Box	referer	nce I	.8: Prov	rovide the code of territory as appearing in Part 1 of Annex II to Regulation (EU) No 206/2010.	
— Box	referer	nce I	.11: Pla	Place of origin: name and address of the dispatch establishment.	
				Registration number (railway wagons or container and lorries), flight number (aircraft) or name (ship) is to b nd reloading, the consignor must inform the BIP of entry into the Union.	pe provided. In

co	OUNTRY Model BOV							
١١.	Health information	II.a. Certificate reference number	II.b.					
-	Box reference I.20: Indicate total gross weight and total net weight.							
-	- Box reference I.23: For containers or boxes, the container number and the seal number (if applicable) must be included.							
-	Box reference I.28: Nature of commodity: Indicate "carcass-whole",	"carcass-side", "carcass-quarters", "cu	ts", "offal" or "minced meat".					
	Minced meat is deboned meat that has been minced into fragment (including the adjoining fatty tissues) except heart muscle.	ts and that must have been prepared	exclusively from striated muscle					
-	Box reference I.28: Treatment type: If appropriate, indicate "debone	d"; "bone in"; "matured"						
Pa	rt II:							
(1)	Keep as appropriate.							
(2)	Code of the territory as it appears in Part 1 of Annex II to Regulate	on (EU) No 206/2010.						
(3)	The number of bovine carcasses or wholesale cuts of carcasses, t number where removal of the vertebral column is not required must b 2 (1) of Regulation (EC) No 136/2004.							
(4)	Delete if the consignment is not intended for introduction into Finlar	nd or Sweden.						
(5)	Only matured de-boned meat fulfilling the supplementary guarantee	s referred to in footnote (8).						
(6)	Supplementary guarantees regarding import of matured de-boned me to Regulation (EU) No 206/2010 with the entry "H".	eat to be provided when required in co	olumn 5 "SG" of Part 1 of Annex II					
(7)	) Delete when the exporting country carries out vaccination against foot-and-mouth disease with serotypes A, O or C, and this country is allowed to import into the Union matured de-boned meat which fulfils the supplementary guarantees described, in footnote ( <sup>8</sup> ).							
( <sup>8</sup> )	) Supplementary guarantees regarding meats from matured de-boned meat to be provided when required in column 5 "SG" of Part 1 of Annex II to Regulation (EU) No 206/2010, with the entry "A".							
( <sup>9</sup> )	Supplementary guarantees regarding meats from matured de-boned II to Regulation (EU) No 206/2010, with the entry "F". The matured de days after the date of slaughter of the animals.							
(10	) The list of approved holdings provided by the competent authority authority. The Commission will ensure that this list of approved ho integrated computerised veterinary system (TRACES).							
(11	) Date or dates of slaughter. Imports of this meat shall not be allow authorisation for importation into the Union of the third country, terri where restrictive measures have been adopted by the Union again	itory or part thereof referred to in box	es I.7 and I.8, or during a period					
(12	) Supplementary guarantees concerning tuberculosis test, to be provid (EU) No 206/2010, with the entry "E". Intra-dermal tuberculosis test to 64/432/EEC.							
(13	) List of countries in the Annex to Decision 2007/453/EC.							
(14	) Alternative guarantee may be provided when allowed for by the No 206/2010.	entry "J" in column 5 "SG" of Part 1	of Annex II to Regulation (EU)					
Of	ficial veterinarian							
	Name (in capital letters):	Qualifica	tion and title:					
	Date:	Signature	9:					
	Stamp:							

Model OVI

cou	NTRY								Veterinary certi	ficate to El
	l.1.	Consignor		1.2.	Certifica	ate ref	erence No		I.2.a.	
		Name		I.3. Central competent authority						
		Address	1.0.							
, T		Tel.		1.4.	Local c	ompet	ent authority			
um	1.5.	Consignee		I.6.						
onsiç		Name								
D D		Address				_				
atche		Postal code								
dispa		Tel.								
Part I: Details of dispatched consignment	1.7.	Country of origin ISO code I.8. Region of origin	Code	1.9.	Country destina	r of tion	ISO code	l.10.	Region of destination	n Code
Det		Place of evicin								
art:	1. 1 1.	Place of origin		l.12.						
å		Name Approval number Address								
	I.13.	Place of loading	l.14.	Date of	depar	ture				
	I.15. Means of transport					IP in E	U			
		Road vehicle Other I Identification		l.17.						
		Documentary references								
	l.18.	Description of commodity				I.19	. Commodity	code (	(HS code)	
								1.20. 0	Quantity	
									-	
	1.21.	Temperature of product						1.22. N	lumber of packages	
		Ambient Chilled		Froze	n 🗖					
	1.23.	Seal/Container No						I.24. T	Type of packaging	
	1.25.	Commodities certified for:								
		Human consumption 🔲								
	1.26.			- I.27.	For imp	ort or	admission ir	nto EU		
	1.28.	Identification of the commodities		1						
		Species Nature of Treatment (scientific name) commodity type	Abatte			er of e ng plar	establishmen nt Col	ts d store	Number of packages	Net weight

		NTRY					Model OVI				
	Ш.	Hea	lth informat	tion			II.a. Certificate refe	erence number	II.b.		
	11.1.	Public	c Health A	ttestation							
I, the undersigned official veterinarian, declare that I am aware of the relevant requirements of Regulations (EC) N (EC) No 852/2004, (EC) No 853/2004, (EC) No 854/2004 and (EC) No 999/2001 and certify that the meat of domest caprine animals described in Part I was produced in accordance with those requirements, in particular that:											
Part II: Certification		II.1.1.			at] ( <sup>1</sup> ) comes from (an) ation (EC) No 852/2004;	establishm	ent(s) implementing	g a programme b	ased on the HACC	次 principles in	
(1) II.1.2. the meat has been obtained in compliance with the conditions set out in Section I of Annex III to Regulation										) No 853/2004;	
( <sup>1</sup> ) II.1.3. [the minced meat has been produced in compliance with Section V of Annex III to Regulation (EC) No 853/2004 and froz internal temperature of not more than - 18 °C;]									nd frozen to an		
	-	II.1.4.			nd fit for human consum and Chapters II and IX of					ccordance with	
		ll.1.5.	( <sup>1</sup> ) either		or parts of the carcass ha egulation (EC) No 854/20		marked with a healt	h mark in accorda	ance with Chapter II	I of Section I of	
	( <sup>1</sup> ) or [the packages of [meat] [minced meat] ( <sup>1</sup> ) have been marked with an identification mark in accordance with Sect Annex II to Regulation (EC) No 853/2004;]								vith Section I of		
	II.1.6. the [meat] [minced meat] ( <sup>1</sup> ) satisfies the relevant criteria set out in Regulation (EC) No 2073/2005 on microbiological c foodstuffs;							gical criteria for			
	II.1.7. the guarantees covering live animals and products thereof provided by the residue plans submitted in accordance with Di 96/23/EC, and in particular Article 29 thereof, are fulfilled;							e with Directive			
		II.1.8.			t] ( <sup>1</sup> ) has been stored ar to Regulation (EC) No 8		rted in accordance	with the relevant	requirements of Se	ections I and V	
		ll.1.9.	with regar	rd to bovine sp	oongiform encephalopathy	/ (BSE):					
	(1)	either [	II.1.9.1. for	<sup>,</sup> imports from	a country or a region wit	h a negligi	ble BSE risk and lis	sted as such in D	ecision 2007/453/E0	D:	
					y or region is classified in negligible BSE risk;	accordanc	e with Article 5(2) o	of Regulation (EC)	No 999/2001 as a c	ountry or region	
					ls from which the meat c ith negligible BSE risk; ( <sup>2</sup> )		meat was derived v	were born, continu	lously reared and s	aughtered in a	
			( <sup>1</sup> ) [(	(c) if in the co	ountry or region there hav	ve been B§	SE indigenous case	s:			
				( <sup>1</sup> ) either	[the animals were born a meal and greaves derive				g of ruminants with	meat-and-bone	
				( <sup>1</sup> ) or	[the meat or minced me Annex V to Regulation (E ovine or caprine animals	EC) No 999					
	(1)	) or	[11.1.9.2.	for imports fro	m a country or a region v	with a cont	trolled BSE risk and	d listed as such in	Decision 2007/453	/EC:	
					/ or region is classified in controlled BSE risk;	accordanc	e with Article 5(2) o	f Regulation (EC)	No 999/2001 as a c	ountry or region	
	(b) animals from which the meat or minced meat was derived have not been slaughtered after stunning by means of injected into the cranial cavity or killed by the same method or slaughtered by laceration after stunning of cer nervous tissue by means of an elongated rod-shaped instrument introduced into the cranial cavity;										

COUNT	'RY					Model OVI				
П.	Health i	nform	ation		II.a. Certificate reference number	ll.b				
		(1)	either	[(c) the meat or minced meat does not conta Regulation (EC) No 999/2001, or mecha animals.]						
		(1)	or	(c) the carcasses, half carcasses or half ca no specified risk material other than the						
	( <sup>1</sup> ) or	[11.	.1.9.3.	for imports from a country or a region whic (EC) No 999/2001 or has been categorised Decision 2007/453/EC:						
				<ul> <li>(a) the country or region has not been categorised as a country or region</li> </ul>		ed in accordance with Article 5(2) of Regulation (EC) No 999/2001 or $_{\rm N}$ with undetermined BSE risk;				
				<ul><li>(b) the animals from which the meat or min derived from ruminants;</li></ul>	ced meat was derived have not been f	ed meat-and-bone meal or greaves				
				(c) the animals from which the meat or mining of gas injected into the cranial cavity or central nervous tissue by means of an	killed by the same method or slaught	ered by laceration after stunning of				
		(1)	either	[(d) the meat or minced meat was not deriv	ed from:					
				(i) specified risk material as defined in	Annex V to Regulation (EC) No 999/2	2001;				
				(ii) nervous and lymphatic tissues expo	sed during the deboning process;					
				(iii) mechanically separated meat obtain	ed from bones of domestic ovine or c	aprine animals.]				
		(1)	or	(d) the carcasses, half carcasses or half ca no specified risk material other than the						
II.2.	Animal	Healt	th atte	estation						
	l, the u	ndersi	igned	official veterinarian, hereby certify, that the f	resh meat described in Part I:					
	∥.2.1.	has	been	obtained in the territory/ies with code:	( <sup>3</sup> ) which, at the date of iss	uing this certificate:				
			has be and	een free for 12 months from rinderpest, and d	uring the same period no vaccination a	gainst this disease has taken place,				
	( <sup>1</sup> ) eithei			een free for 12 months from foot-and-mouth aken place;]	disease, and during the same period	no vaccination against this disease				
	( <sup>1</sup> ) or	[(b)	break	een considered free from foot-and-mouth dis s afterwards, and authorised to export this r m/yyyy);]						
	( <sup>1</sup> ) ( <sup>4</sup> ) or	[(b)	vaccir anima	nation programmes against foot-and-mouth d ls;]	disease are being officially carried out	and controlled in domestic bovine				
	II.2.2.	has	been	obtained from animals that:						
		( <sup>1</sup> ) e	either	[have remained in the territory described slaughter;]	under point II.2.1 since birth, or for at	least the last three months before				
		( <sup>1</sup> ) d	or	[have been introduced on territory with code ( <sup>3</sup> ) that at that dat						
		( <sup>1</sup> ) d	or	[have been introduced on Member State	. (dd/mm/yyyy) into the territory descrit	bed under point II.2.1, from the EU				

cou	NTRY		1	Model OVI
١١.	Healt	h information	II.a. Certificate reference number	ll.b.
	II.2.3.	has been obtained from animals coming from holdings:		
		(a) in which none of the animals present therein have be	een vaccinated against [foot-and-mouth	n disease or] ( <sup>5</sup> ) rinderpest,
		(b) not subject to prohibition as a result of an outbreak of	of ovine or caprine brucellosis during t	he previous six weeks, and
	( <sup>1</sup> ) either	[(c) in and around which, in an area of 10 km radius, th during the previous 30 days;]	ere has been no case/outbreak of for	ot-and-mouth disease or rinderpest
	( <sup>1</sup> ) ( <sup>4</sup> ) or	[(c) where there is no official restriction for health reason case/outbreak of foot-and-mouth disease or rinderper		f 50 km radius, there has been no
		(d) where they have remained for at least 40 days befor	e direct dispatch to the slaughterhous	e;]
	( <sup>1</sup> ) ( <sup>8</sup> ) or	[(d) where they have remained for at least 40 days be veterinary authority without coming into contact with a slaughterhouse;]		
	II.2.4.	has been obtained from animals which:		
		<ul> <li>(a) have been transported from their holdings in vehicles without contact with other animals which did not corr</li> </ul>	s, cleaned and disinfected before loadi ply with the requirements set out in p	ing, to an approved slaughterhouse oints II.2.1, II.2.2 and II.2.3,
		(b) at the slaughterhouse, have passed ante-mortem hea shown no evidence of the diseases referred to in po		re slaughter and, in particular, have
		(c) have been slaughtered on (dd/mm/yyyy)	or between (dd/mm/yyyy	/) and(dd/mm/yyyy) ( <sup>6</sup> );
	II.2.5.	has been obtained in an establishment around which, wi referred to in point II.2.1 during the previous 30 days or importation into the Union has been authorised only after and disinfection of the establishment under the control o	, in the event of a case/outbreak of d slaughter of all animals present, remov	isease, the preparation of meat for
	II.2.6.			
	( <sup>1</sup> ) either	[has been obtained and prepared without contact with o	ther meats not complying with the co	nditions required in this certificate.]
	( <sup>1</sup> ) ( <sup>4</sup> ) or	[contains [boneless meat] [and] [minced meat] ( <sup>1</sup> ), obta carcasses in which the main accessible lymphatic glar temperature above + 2 °C for at least 24 hours before th 6.0 when tested electronically in the middle of the long	nds have been removed, which have ne bones were removed and in which t	been submitted to maturation at a he pH value of the meat was below
		has been kept strictly separate from meat not conform production, de-boning and storage until it has been pac		
	( <sup>1</sup> )( <sup>7</sup> ) or	[contains [boneless meat] [and] [minced meat] ( <sup>1</sup> ), obta carcasses in which the main accessible lymphatic glar temperature above + 2 °C for at least 24 hours before	nds have been removed, which have	
		has been kept strictly separate from meat not conform production, de-boning and storage until it has been pac		
II.3.	Animal	welfare attestation		
		dersigned official veterinarian, hereby certify, that the fresh ghterhouse before and at the time of slaughter or killing i		

COUNTRY		Model OVI					
II. Health information	II.a. Certificate reference number	II.b.					
Notes		<u></u>					
This certificate is meant for fresh meat, including minced meat, of do	This certificate is meant for fresh meat, including minced meat, of domestic ovine animals (Ovis aries) and caprine animals (Capra hircus).						
Fresh meat means all animal parts fit for human consumption whether fr	resh, chilled or frozen.						
Part I:							
- Box reference I.8: Provide the code of territory as appearing in Part	1 of Annex II to Regulation (EU) No 2	206/2010.					
- Box reference I.11: Place of origin: name and address of the dispate	h establishment.						
<ul> <li>Box reference I.15: Registration number (railway wagons or contained case of unloading and reloading, the consignor must inform the BIP</li> </ul>		r name (ship) is to be provided. In					
<ul> <li>Box reference I.19: Use the appropriate HS code: 02.04, 02.06 or 05. column 5 "SG" of Part 1 of Annex II to Regulation (EU) No 206/2010</li> </ul>							
- Box reference I.20: Indicate total gross weight and total net weight.							
- Box reference I.23: For containers or boxes, the container number a	nd the seal number (if applicable) sho	uld be included.					
<ul> <li>Box reference I.28: Nature of commodity: Indicate "carcass-whole", "c meat is de-boned meat that has been minced into fragments and that adjoining fatty tissues) except heart muscle.</li> </ul>							
<ul> <li>Box reference I.28: Treatment type: If appropriate, indicate "de-bone freezing (mm/yy) of the cuts/pieces.</li> </ul>	əd"; 'bone in"; "matured" and/or "minc	ed". If frozen, indicate the date of					
Part II:							
( <sup>1</sup> ) Keep as appropriate.							
( <sup>2</sup> ) List of countries in the Annex to Decision 2007/453/EC.							
(3) Code of the territory as it appears in Part 1 of Annex II to Regulation	n (EU) No 206/2010.						
( <sup>4</sup> ) Supplementary guarantees regarding meats from matured de-boned n to Regulation (EU) No 206/2010, with the entry "A".	neat to be provided when required in a	olumn 5 "SG" of Part 1 of Annex II					
( <sup>5</sup> ) Delete when the exporting country carries out vaccination against authorised to import into the Union matured de-boned meat which fu							
( <sup>6</sup> ) Date or dates of slaughter. Imports of this meat shall not be allow authorisation for importation into the Union of the third country, territor restrictive measures have been adopted by the Union against import	y or part thereof referred to in boxes I.	7 and I.8, or during a period where					
( <sup>7</sup> ) Supplementary guarantees regarding meats from matured de-boned n to Regulation (EU) No 206/2010, with the entry "F". The matured de-b days after the date of slaughter of the animals.	neat to be provided when required in o coned meat shall not be authorised for	olumn 5 "SG" of Part 1 of Annex II r importation into the Union until 21					
( <sup>8</sup> ) Alternative guarantee may be provided when allowed for by the (EU) No 206/2010.	∋ entry " <b>J</b> " in column 5 "SG" of F	art 1 of Annex II to Regulation					
Official veterinarian							
Name (in capital letters):	Qualification and title:	:					
Date: Signature:							
Stamp:							

			Mode	del POR					
	COUNTRY			1			Veterinary certi	ficate to El	
	I.1. Consignor		I.2. Certifica	ate reference	e numbe	r I.2.a.			
	Name			I.3. Central Competent Authority					
	Address					-			
ent	Tel. No			I.4. Local Competent Authority					
ŭuŭ	I.5. Consignee			I.6.					
nsiç	Name								
o p	Address								
tche	Postal code								
spa	Tel. No								
Part I: Details of dispatched consignment	I.7. Country ISO of origin code	.8. Region of origin	Code	I.9. Country destina		ISO code	I.10. Region of destination	Code	
Deta	I.11. Place of origin			l.12.		·			
Ë		Approval number							
Ра	Address								
	I.13. Place of loading	I.14. Date of departure							
	I.15. Means of transport		I.16. Entry BIP in EU						
	Aeroplane Ship	Railway wago	on 🗌						
	Road vehicle Other								
	Identification:			l.17.					
	Documentary references:								
	I.18. Description of commodity			I.19. Commodity code (HS code)					
						1.00	<b>0</b>		
						1.20.	Quantity		
	I.21. Temperature of product					I.22.	Number of packages		
	Ambient	Chiled		Frozen	1				
					1				
	I.23. Identification of container/sea	Inumber				I.24.	Type of packaging		
	I.25. Commodities certified for: Human consumption								
	1.26.	I.27. For import or admission into EU							
	I.28. Identification of the commodit	ies							
	Species Nature o		Арр	roval number e	stablishmer	its	Number	Net	
	(Scientific name) commodi	ty type	Abatto	r Cutting a	lant Colo	store	of packages	weight	
			Aballo	r Cutting p		51018			

	COUNTRY					Model POR				
	П.	Health	information		II.a. Certificate reference number	II.b.				
	II.1.	Public	Health Attesta	ation						
		(EC) N	o 852/2004, (E	C) No 853/2	rian, declare that I am aware of the relevant rec 2004 and (EC) No 854/2004 and hereby certif ince with those requirements, in particular tha	y that the meat of domestic swine described				
fication		II.1.1			(1) comes from (an) establishment(s) implen with Regulation (EC) No 852/2004;	nenting a programme based on the HACCP				
Part II: Certification		II.1.2	the meat has No 853/2004;		n obtained in compliance with the conditions set out in Section I of Annex III to Regulation (EC)					
Parl		II.1.3			he requirements of Regulation (EC) No 2075/2005 laying down specific rules on official controls for at, and in particular:					
			(1) either	[has bee	n subjected to an examination by a digestion	method with negative results]				
			(1) or	[has bee No 2075/	en subjected to a freezing treatment in acc /2005;]	cordance with Annex II to Regulation (EC)				
			(1) or	holding c	ase of meat from domestic swine kept solely or category of holdings that has been officiall <i>Trichinella</i> in accordance with Annex IV to Re	y recognized by the competent authority as				
		(¹) II.1.4		d meat has been produced in accordance with Section V of Annex III to Regulation (EC) No 853/2004 and n internal temperature of not more than –18 $^\circ C;]$						
		II.1.5		vith Chapte	een found fit for human consumption following ante and post-mortem inspections carried out in Chapter II of Section I and Chapters IV and IX of Section IV of Annex I to Regulation (EC)					
		II.1.6 (	) either	-	e carcass or parts of the carcass have been marked with a health mark in accordance wi lapter III of Section I of Annex I to Regulation (EC) No 854/2004;]					
			(1) or		kages of [meat] [minced meat] (1) have be nce with Section I of Annex II to Regulation (Effective)					
		II.1.7	the [meat] [min criteria for foo	-	(1) satisfies the relevant criteria set out in Regu	lation (EC) No 2073/2005 on microbiological				
		II.1.8	•	•	live animals and products thereof provided by and in particular Article 29, are fulfilled.	the residue plans submitted in accordance				
		II.1.9			t] (') has been stored and transported in acc vely of Annex III to Regulation (EC) No 853/20					
.,.					of Regulation (EC) No 1688/2005 implementi erning Salmonella for consignments to Finland					
	II.2.	Anima	l Health attest	ation						
		I, the u	ndersigned offi	cial veterina	arian, hereby certify, that the fresh meat descr	bed in Part I :				
		II.2.1	has been obta	ained in the	territory/ies with code:	) which, at the date of issuing this certificate:				
			(1) either		been free for 12 months from foot-and-mou sical swine fever, swine vesicular disease, and					
			(1) or		nas been free for 12 months from rinderpest, Afri classical swine fever] (') and [swine vesicular o					

	Health	information		II.a. Certificate reference number	II.b.				
			[! 	as been considered free from [foot-and-mousswine vesicular disease] (1), sincead cases/outbreaks afterwards, and author Regulation (EC) No/, of	(dd/mm/yyyy), without havin orised to export this meat by Commissio				
				g the last 12 months no vaccination agains rts of domestic animals vaccinated agains pry;					
	II.2.2 has been obtained from animals that:								
	(1) either [have remained in the territory described under point II.2.1 since birth, or for at least the months before slaughter;]								
		(1) <i>or</i>	point II.2.	en introduced on					
		(1) <i>or</i>		en introduced on (do 1, from the EU Member State					
	II.2.3 has been obtained from animals coming from holdings:								
	<ul> <li>(a) in which none of the animals present therein have been vaccinated against the diseases point II.2.1,</li> </ul>								
		o case/outbreak of the diseases referred to							
		(c) that are weeks;	not subject	to prohibition as a result of an outbreak of	f porcine brucellosis during the previous si				
				ing has been received that pigs are not fed with catering waste, are subject to official controls the list established by the competent authority for the purpose of importing pig meat into the					
	II.2.4 has been obtained from animals that:								
		(a) have rem	ained separ	parate since birth from wild cloven-hoofed animals,					
				orted from their holdings in vehicles, cleaned and disinfected before loading, to an approved thout contact with other animals which did not comply with the conditions set out in points II.2.1					
				use, have passed ante-mortem health inspection during the 24 hours before slaughter and, i own no evidence of the diseases referred to in point II.2.1, and					
	<ul> <li>(d) have been slaughtered on</li></ul>								
	II.2.5	of the diseas preparation o	ses referred of meat for in	an establishment around which, within a radius of 10 km, there has been no case/outbreak ed to in point II.2.1 during the previous 40 days or, in the event of a case of disease, the r importation into the Union has been authorised only after slaughter of all animals present nd the total cleaning and disinfection of the establishment under the control of an officia					
	II.2.6 has been obtained and prepared without contact with other meats not complying with the conditions required certificate.								
.3.	Anima	I welfare atte	station						
				rian, hereby certify, that the fresh meat descri se before and at the time of slaughter or killir					

COUNTRY Model POR										
Π.	Health information	II.a. Certificate reference number	II.b.							
No	Notes									
This certificate is meant for fresh meat, including minced meat, of domestic swine (Sus scrofa).										
Fresh meat means all animal parts fit for human consumption whether fresh, chilled or frozen.										
Part I:										
   Pa	<ul> <li>Box reference I.8: Provide the code of territory as appearing in Part 1 of Annex II to Regulation (EU) No 206/2010.</li> <li>Box reference I.11: Place of origin: name and address of the dispatch establishment.</li> <li>Box reference I.15: Registration number (railway wagons or container and lorries), flight number (aircraft) or name (ship) is to be provided. In case of unloading and reloading, the consignor must inform the BIP of entry into the Union.</li> <li>Box reference I.19: Use the appropriate HS code: 02.03, 02.06, 02.09, 05.04 or 15.01.</li> <li>Box reference I.20: Indicate total gross weight and total net weight.</li> <li>Box reference I.23: For containers or boxes, the container number and the seal number (if applicable) should be included.</li> <li>Box reference I.28: <i>Nature of commodity</i>: Indicate 'carcass-whole', 'carcass-side', 'carcass-quarters', 'cuts' or 'minced meat'. Minced meat is deboned meat that has been minced into fragments and that must have been prepared exclusively from striated muscle (including the adjoining fatty tissues) except heart muscle.</li> <li>Box reference I.28: Treatment type: If appropriate, indicate 'deboned'; 'bone in'; 'matured' and/or 'minced'. If frozen, indicate the date of freezing (mm/yy) of the cuts/pieces.</li> </ul>									
	rt II:									
( <sup>1</sup> ) ( <sup>2</sup> )	Keep as appropriate. Delete if the consignment is not intende	ed for import into Finland or Sweden								
() ( <sup>3</sup> )	-	rt 1 of Annex II to Regulation (EU) No 206/201	0.							
( <sup>4</sup> )	Supplementary guarantees to be provi	ded when required in column 5 'SG' of Part 1								
	with the entry 'D'.									
		Catering waste means: all waste from food intended for human consumption from restaurants, catering facilities or kitchens, including industrial kitchens and household kitchens of the farmer or persons tending pigs.								
(5)	) Date or dates of slaughter. Imports of this meat shall not be allowed when obtained from animals slaughtered either prior to the date of authorisation for importation into the Union of the third country, territory or part thereof referred to in boxes I.7 and I.8, or during a period where restrictive measures have been adopted by the Union against imports of this meat from this third country, territory or part thereof.									
Official veterinarian										
	Name (in capital letters):	Qualification	n and title:							
	Date:	Signature:								
		orginature.								
	Stamp:									

			Mod	el EQU								
	COUNTRY			1			Veterinary cer	tificate to EU				
Part I: Details of dispatched consignment	I.1. Consignor	I.2. Certificate reference number I.2.a.										
	Name	I.3. Central Competent Authority										
	Address											
	Tel. No	I.4. Local Competent Authority										
	I.5. Consignee	I.6.										
	Name											
	Address											
	Postal code											
	Tel. No											
ails of d	I.7. Country ISO I. of origin code	8. Region of origin	Code	I.9. Country destina		ISO code	I.10. Region of destination	Code				
Deta	I.11. Place of origin			I.12.								
Ë		pproval number										
Ъ	Address	Address										
	I.13. Place of loading	I.14. Date of departure										
	I.15. Means of transport			I.16. Entry BIP in EU								
	Aeroplane Ship	Aeroplane Ship Railway wagon										
	Road vehicle 🗌 Other											
	Identification:	Identification:				1.17.						
	Documentary references:	Documentary references:										
	I.18. Description of commodity						I.19. Commodity code (HS code)					
				I.20. Quantity								
		I.21. Temperature of product					I.22. Number of packages					
	Ambient	Ambient Chiled					Frozen					
	I.23. Identification of container/seal	number			1.24.	I.24. Type of packaging						
	I.25. Commodities certified for: Human consumption											
	1.26.	.26.					I.27. For import or admission into EU					
	I.28. Identification of the commoditie	es	I									
	Species Nati (Scientific name) comr	umber establishments Number Net of packages weight				Net weight						
		Abati	Cutting plant Cold store									

.. . . . . . .

	COUNT	RY			Model EQU				
	П.	Health	information	II.a. Certificate reference number	II.b.				
-	II.1.	Public Health Attestation							
		(EC) N	o 852/2004, (EC	al veterinarian, declare that I am aware of the relevant req ) No 853/2004 and (EC) No 854/2004 and hereby certify t in accordance with those requirements, in particular that	hat the meat of domestic solipeds described				
ification		II.1.1		es from (an) establishment(s) implementing a progra h Regulation (EC) No 852/2004;	mme based on the HACCP principles in				
Part II: Certification		II.1.2	the meat has b No 853/2004;	een obtained in compliance with the conditions set out	in Section I of Annex III to Regulation (EC)				
Par		II.1.3		the requirements of Regulation (EC) No 2075/2005 lay n meat, and in particular, has been subject to an exami					
		II.1.4		been found fit for human consumption following ante a th Chapter II of Section I and Chapters III and IX of					
		II.1.5	(1) either	[the carcass or parts of the carcass have been mark Chapter III of Section I of Annex I to Regulation (EC) No					
			(1) or	[the packages of meat have been marked with an identi Annex II to Regulation (EC) No 853/2004;]	fication mark in accordance with Section I of				
		II.1.6	the meat satisfies the relevant criteria set out in Regulation (EC) No 2073/2005 on microbiological criteria for foodstuffs;						
		II.1.7		covering live animals and products thereof provided by 6/23/EC, and in particular Article 29 thereof, are fulfilled					
		II.1.8		een stored and transported in accordance with the relev ) No 853/2004.	rant requirements of Section I of Annex III to				
	II.2.	Anima	l Health attesta	tion					
		I, the u	ndersigned offic	ial veterinarian, hereby certify, that the fresh meat descri	bed in Part I:				
		II.2.1	has been obtai	ned in the territory/ies with code:	(²);				
		II.2.2	has been obtai	ned from domestic solipeds, which:					
			(1) either	[have remained in the territory described under point I months before slaughter;]	.2.1 since birth, or for at least the last three				
			(1) <i>or</i>	[have been introduced on					
			(1) <i>or</i>	[have been introduced on(dd/ point II.2.1, from the EU Member State					
		II.2.3	which, within a previous 40 da has been auth		n animals which were slaughtered on				

### COUNTRY

COUNTR	COUNTRY Model EQ									
II.	Health information	II.a. Certificate reference number	II.b.							
	II.2.4 has been obtained and prepared without contact with other meats not complying with the conditions required in this certificate.									
II.3.	Animal welfare attestation									
	I, the undersigned official veterinarian, hereby certify that the fresh meat described in this certificate derives from animals which have been treated in the slaughterhouse before and at the time of slaughter or killing in accordance with the relevant provisions of Union legislation.									
Notes										
This cert breeds).	This certificate is meant for fresh meat, excluding minced meat, of domestic solipeds (Equus caballus, Equus asinus and their cross- breeds).									
Fresh me	eat means all animal parts fit for hu	man consumption whether fresh, chilled	l or frozen.							
Part I:										
— Boxi	reference I.8: Provide the code of te	erritory as appearing in Part 1 of Annex	II to Regulation (EU) No 206/2010.							
— Boxi	reference I.11: Place of origin: nam	e and address of the dispatch establish	ment.							
		r (railway wagons or container and lorr ading, the consignor must inform the BI	ies), flight number (aircraft) or name (ship) is to be P of entry into the Union.							
— Box	reference I.19: Use the appropriate	HS code: 02.05, 02.06 or 05.04.	-							
— Box i	reference I.20: Indicate total gross	weight and total net weight.								
— Box ı	reference I.23: For containers or bo	xes, the container number and the seal	number (if applicable) should be included.							
— Box ı	reference I.28: Nature of commodit	y: Indicate 'carcass-whole', 'carcass-sic	le', 'carcass-quarters' or 'cuts'.							
	reference I.28: <i>Treatment type</i> : If a ing (mm/yy) of the cuts/pieces.	appropriate, indicate 'deboned'; 'bone	in' and/or 'matured'. If frozen, indicate the date of							
Part II:										
(1) Keep	as appropriate.									
(²) Code	e of the territory as it appears in Pa	t 1 of Annex II to Regulation (EU) No 20	06/2010.							
for in	nportation into the Union of the thir	d country, territory or part thereof referr	slaughtered either prior to the date of authorisation ed to in boxes I.7 and I.8, or during a period where eat from this third country, territory or part thereof.							
Official v	eterinarian									
	Name (in capital letters):	Quali	fication and title:							
	Date:	Signa	ature:							
	Stamp:									

		Mod	del RUF					
	COUNTRY		1	Veterinary certificate to EU				
	I.1. Consignor		I.2. Certificate reference	ce number I.2.a.				
	Name		I.3. Central Competent Authority					
	Address							
ent	Tel. No		I.4. Local Competent A	Authority				
Jum	I.5. Consignee		I.6.					
nsiç	Name							
d co	Address							
tche	Postal code							
spat	Tel. No							
Part I: Details of dispatched consignment	I.7. Country ISO I.8. Reg of origin code of or	ion Code rigin	I.9. Country of destination	ISO I.10. Region of Code destination				
Deta	I.11. Place of origin		1.12.					
Ξt	Name Approva	Inumber						
Pai	Address							
	I.13. Place of loading		I.14. Date of departure					
	I.15. Means of transport		I.16. Entry BIP in EU					
		Railway wagon 🗌						
	Road vehicle 🗌 Other 🗌							
	Identification:		1.17.					
	Documentary references:							
	I.18. Description of commodity		I.19. Commodity code (HS code)					
				I.20. Quantity				
	I.21. Temperature of product			I.22. Number of packages				
	Ambient Chiled		Frozen					
	I.23. Identification of container/seal number			I.24. Type of packaging				
	I.25. Commodities certified for: Human consumption							
	1.26.		I.27. For import or admission into EU					
	I.28. Identification of the commodities		1					
	Species Nature of (Scientific name) commodity	Treatment App type Abatto	oroval number establishme ir Cutting plant Col	nts Number Net of packages weight Id store				

	COUN	TRY						Model RUF	
	П.	Health	information		II.a. Certificate reference	e number	II.b.		
	II.1.	Public	Health Attestation						
ification		No 178 the me and the	3/2002, (EC)   at of farmed eir cross-bree	No 852/200- animals of t eds), <i>Ovis a</i>	erinarian, declare that I a 4, (EC) No 853/2004, (EC he order Artiodactyla (exc <i>rries, Capra hircus,</i> Suida I was produced in accorda	) No 854/2004 and Iuding bovine anim e and Tayassuidae	l (EC) No 999/2001 a nals (including <i>Bison</i> e), and of the familie	and hereby certify that and <i>Bubalus</i> species s Rhinocerotidae and	
Part II: Certification		II.1.1			(an) establishment(s) impl ion (EC) No 852/2004;	lementing a progra	amme based on the	HACCP principles in	
Part II		II.1.2	the meat has No 853/2004		ned in accordance with the	e conditions set out	in Section III of Anne	x III to Regulation (EC)	
		II.1.3		with Chapte	d fit for human consumpti er II of Section I and Cha				
		II.1.4	(1) either		cass or parts of the carca III of Section I of Annex I to			rk in accordance with	
			(1) or		ckages of meat have bee I of Annex II to Regulation			k in accordance with	
		II.1.5	the meat sa foodstuffs;	tisfies the r	elevant criteria set out in	Regulation (EC) N	o 2073/2005 on micr	robiological criteria for	
		II.1.6	•	•	live animals and products and in particular Article 29 t			bmitted in accordance	
	(1)	(²) [II.1.7	with regard t	o Chronic W	asting Disease (CWD):				
			animals white other diagno	ch have bee ostic method	or is derived exclusively fi en examined for Chronic V d recognised by the comp lerd where Chronic Wastin	Vasting Disease by betent authority with	/ histopathology, imn h negative results ar	nunohistochemistry or nd is not derived from	
		II.1.8	the meat has Regulation (I		d and transported in accord 2004.	dance with the relev	vant requirements of \$	Section I of Annex III to	
	II.2.	Anima	I Health attes	station					
		I, the u	ndersigned of	ficial veterina	arian, hereby certify, that th	e fresh meat descri	bed in Part I:		
		II.2.1	has been ob	tained in the	territory/ies with code:	(3)	which, at the date of	issuing this certificate:	
				n free for 12 n place, and	months from rinderpest, a	nd during the same	e period no vaccinatio	on against this disease	
		(1) either		n free for 12 ase has take	months from foot-and-mou en place;]	uth disease, and du	ring the same period	no vaccination against	
Part II: C		(1) or	having h	ad cases/ou	d free from foot-and-mout tbreaks afterwards, and au (dd/mm/yy	thorised to export th			
		(1) (4) or		ion program c bovine anir	mes against foot-and-mo mals;]	uth disease are be	eing officially carried	out and controlled in	

co	UNTRY			Model RUF				
Ш.	Health	information	II.a. Certificate reference number	II.b.				
	II.2.2	has been obtained from a	animals that:					
			nained in the territory described under point I before slaughter;]	I.2.1 since birth, or for at least the last three				
		point II.2	en introduced on(dd/ .1, from the territory with code this fresh meat into the Union;]					
	II.2.3	has been obtained from a	animals coming from holdings:					
		<ul> <li>(a) in which none of t or] (<sup>5</sup>) rinderpest,</li> </ul>	he animals present therein have been va	ccinated against [foot-and-mouth disease				
		(b) where regular veterinary inspections are carried out to diagnose diseases transmissible to humans or animals and, these holdings are not subject to prohibition as a result of an outbreak of brucellosis during the previous six weeks, and						
	(1) either	[(c) in and around which rinderpest during the	in an area of 10 km radius, there has been no previous 30 days,]	case/outbreak of foot-and-mouth disease or				
	(1) (4) or		cial restriction for health reasons and in and ar tbreak of foot-and-mouth disease or rinderpes					
		(d) where the animals have remained for at least 40 days before direct dispatch to the slaughterhous						
	II.2.4	has been obtained from a	animals:					
	(1) either	[(a) which have been transported from their holdings in vehicles, cleaned and disinfected before loading, to a approved slaughterhouse, without contact with other animals which did not comply with the conditions mentione above,						
			terhouse, have passed ante-mortem health inspection during the 24 hours before slaughte have shown no evidence of the diseases referred to in point II.2.1, and					
			ughtered on(dd/mm/yyyy) (6);]	n/yyyy) or between				
	(1) <i>or</i>		aughtered on the holding of origin, followir olding, who has provided a written statement					
			unacceptable risk would have been posed to t of the animals to an slaughterhouse,	he welfare of the animals or to their handlers				
		<ul> <li>the holding had animals,</li> </ul>	been inspected and authorised by the com	petent authority for the slaughter of game				
			passed the ante-mortem health inspection de shown no evidence of the diseases referred	<b>o</b>				
		<ul> <li>the animals were (dd/mm/yyyy), (<sup>6</sup>)</li> </ul>	e slaughtered between)	(dd/mm/yyyy) and				
		<ul> <li>the bleeding of the second second</li></ul>	ne animals was performed correctly, and					
		<ul> <li>the slaughtered a</li> </ul>	animals were eviscerated within three hours of	the time of slaughter, and				
		where more than one	ch have been transported to the approved sla b hour elapsed since the time of slaughter, a to rival of the vehicle used for the transport;]					
	(1) (7) II.2.5	[has been obtained from hoofed animals;]	animals that have remained since birth or for t	the last 3 months separate from wild cloven-				

	Health	information		II.a. Certificate reference number	II.b.
	II.2.6	of the disea preparation	ises referred of meat for in all meat, and	to in point II.2.1 during the previous 30 on mortation into the Union has been author	lius of 10 km, there has been no case/outbread days or, in the event of a case of disease, the rised only after slaughter of all animals present e establishment under the control of an officia
	II.2.7				
		(1) either	[has bee required		ith other meats not complying with the conditior
		(1) (4) or	carcasse submitte removed	es in which the main accessible lymphatic d to maturation at a temperature above + 2	ned meat other than offal that was obtained fror c glands have been removed, which have bee 2 °C for at least 24 hours before the bones wer vas below 6.0 when tested electronically in th ation and before de-boning, and
			certificat		conforming to the requirements set out in th coning and storage until it has been packed reas.]
		( <sup>1</sup> ) ( <sup>8</sup> ) or	carcasse	es in which the main accessible lymphatic d to maturation at a temperature above + 2	ned meat other than offal that was obtained from c glands have been removed, which have bee 2 °C for at least 24 hours before the bones wer
			certificat		conforming to the requirements set out in th poning and storage until it has been packed reas.]
This c anima	ertificate is Is (includin	g <i>Bison</i> and B	<i>Bubalus</i> spec	•	pra hircus, Suidae and Tayassuidae), and of th
<sup>-</sup> his c Inima amilie	ertificate is Ils (includin es Rhinocer	g <i>Bison</i> and <i>B</i> otidae and Ele	<i>Bubalus</i> spec ephantidae, t	ies and their cross-breeds), Ovis aries, Ca	<i>pra hircus,</i> Suidae and Tayassuidae), and of th th or for the last three months in farms.
This c Inima amilie Fresh	ertificate is Ils (includin es Rhinocer meat mear	g <i>Bison</i> and <i>B</i> otidae and Ele	<i>Bubalus</i> spec ephantidae, t	ies and their cross-breeds), Ovis aries, Ca that are domestically kept or bred since bir	<i>pra hircus,</i> Suidae and Tayassuidae), and of th th or for the last three months in farms.
This c anima amilie Fresh <b>Part I</b> :	ertificate is Ils (includin es Rhinocer meat mear	g <i>Bison</i> and <i>B</i> rotidae and Ele ns all animal p	Bubalus spec ephantidae, t arts fit for hu	ies and their cross-breeds), Ovis aries, Ca that are domestically kept or bred since bir	<i>pra hircus</i> , Suidae and Tayassuidae), and of th th or for the last three months in farms. frozen.
anima amilie Fresh <b>Part I</b> : — Bo	ertificate is ils (includin es Rhinocer meat mear : : :	g <i>Bison</i> and <i>B</i> rotidae and Ele ns all animal pa e I.8: Provide t	Bubalus spec ephantidae, t arts fit for hu the code of to	ies and their cross-breeds), <i>Ovis aries, Ca</i> hat are domestically kept or bred since bir man consumption whether fresh, chilled or	r frozen. o Regulation (EU) No 206/2010.
This c anima amilie Fresh <b>Part I</b> : — Bo — Bo — Bo	ertificate is ils (includin es Rhinocer meat mear : cox reference cox reference cox reference	g <i>Bison</i> and <i>B</i> otidae and Ele is all animal p e I.8: Provide t e I.11: Place o e I.15: Registr	Bubalus spec ephantidae, t arts fit for hu the code of to f origin: nam ation numbe	ies and their cross-breeds), Ovis aries, Ca that are domestically kept or bred since bir man consumption whether fresh, chilled or erritory as appearing in Part 1 of Annex II to e and address of the dispatch establishme	<i>pra hircus</i> , Suidae and Tayassuidae), and of th th or for the last three months in farms. frozen. D Regulation (EU) No 206/2010. ent. ), flight number (aircraft) or name (ship) is to b
This c anima amilie Fresh Part I: — Bo — Bo pr	ertificate is ils (includin es Rhinocer meat mear cox reference cox reference cox reference cov reference covided. In c	g <i>Bison</i> and <i>B</i> otidae and Ele as all animal p e I.8: Provide t e I.11: Place o e I.15: Registr case of unload	Bubalus spec ephantidae, f arts fit for hu the code of to of origin: nam ration numbe ling and reloa	ies and their cross-breeds), Ovis aries, Ca that are domestically kept or bred since bir man consumption whether fresh, chilled or erritory as appearing in Part 1 of Annex II to e and address of the dispatch establishme r (railway wagons or container and lorries)	<i>pra hircus</i> , Suidae and Tayassuidae), and of th th or for the last three months in farms. frozen. D Regulation (EU) No 206/2010. ent. ), flight number (aircraft) or name (ship) is to b
This c anima amilie - Tresh - Bo - Bo pr - Bo	ertificate is ils (includin es Rhinocer meat mear cox reference cox reference cox reference cox reference cox reference cox reference	g <i>Bison</i> and <i>B</i> rotidae and Ela ns all animal p e I.8: Provide t e I.11: Place o e I.15: Registr rase of unload e I.19: Use the	Bubalus spec ephantidae, f arts fit for hui the code of to f origin: nam ation numbe ling and reloa	ies and their cross-breeds), <i>Ovis aries, Ca</i> that are domestically kept or bred since bir man consumption whether fresh, chilled or erritory as appearing in Part 1 of Annex II to e and address of the dispatch establishme r (railway wagons or container and lorries) ading, the consignor must inform the BIP o	<i>pra hircus</i> , Suidae and Tayassuidae), and of th th or for the last three months in farms. frozen. D Regulation (EU) No 206/2010. ent. ), flight number (aircraft) or name (ship) is to b
Fhis c anima amilia Fresh - Bo - Bo pr - Bo - Bo - Bo	ertificate is ils (includin es Rhinocer meat mear cox reference ox reference ox reference ox reference ox reference ox reference	g <i>Bison</i> and <i>B</i> otidae and Ele is all animal p e I.8: Provide t e I.11: Place o e I.15: Registr ase of unload e I.19: Use the e I.20: Indicate	Bubalus spec ephantidae, f arts fit for hui the code of to of origin: nam ration numbe ling and reloa e appropriate e total gross f	ies and their cross-breeds), <i>Ovis aries, Ca</i> that are domestically kept or bred since bir man consumption whether fresh, chilled or erritory as appearing in Part 1 of Annex II to e and address of the dispatch establishme r (railway wagons or container and lorries) ading, the consignor must inform the BIP o HS code: 02.06, 02.08.90 or 05.04.	<i>pra hircus</i> , Suidae and Tayassuidae), and of th th or for the last three months in farms. frozen. D Regulation (EU) No 206/2010. ent. ), flight number (aircraft) or name (ship) is to b f entry into the Union.
Fhis c anima amilia Fresh Part I: Ba Ba P Ba Ba Ba Ba Ba Ba Ba Ba Ba Ba Ba Ba Ba	ertificate is ils (includin as Rhinocer meat mear cox reference cox reference cox reference cox reference cox reference cox reference cox reference cox reference	g <i>Bison</i> and <i>B</i> otidae and Ele as all animal p e I.8: Provide t e I.11: Place o e I.15: Registr case of unload e I.19: Use the e I.20: Indicate e I.23: For con	Bubalus spec ephantidae, f arts fit for hui the code of te of origin: nam ration numbe ling and reloa e appropriate e total gross ttainers or bo	ies and their cross-breeds), <i>Ovis aries, Ca</i> that are domestically kept or bred since bir man consumption whether fresh, chilled or erritory as appearing in Part 1 of Annex II to e and address of the dispatch establishme r (railway wagons or container and lorries) ading, the consignor must inform the BIP o HS code: 02.06, 02.08.90 or 05.04. weight and total net weight.	<i>pra hircus</i> , Suidae and Tayassuidae), and of th th or for the last three months in farms. frozen. b Regulation (EU) No 206/2010. ent. ), flight number (aircraft) or name (ship) is to b f entry into the Union.

cc	COUNTRY Model RUF								
Π.	Health information	II.a. Certificate reference number	II.b.						
Pa	rt II:								
(1) (2)	<ol> <li>Keep as appropriate.</li> <li>Supplementary guarantees regarding fresh meat obtained from cervids to be provided when required in column 5 'SG' of Part 1 of Annex II to Regulation (EU) No 206/2010, with the entry 'G'.</li> </ol>								
( <sup>3</sup> )	Code of the territory as it appears in Part 1 of Annex II to Regulation (EU) No 206/2010.								
(4)	Part 1 of Annex II to Regulation (EU)	No 206/2010 with the entry 'A'.	provided when required in column 5 'SG' of						
(5)			disease with serotypes A, O or C, and this e supplementary guarantees described under						
( <sup>6</sup> )	date of authorisation for importation int during a period where restrictive measu territory or part thereof.	o the Union of the third country, territory or p ures have been adopted by the Union again:	ed from animals slaughtered either prior to the part thereof referred to in boxes I.7 and I.8, or st imports of this meat from this third country,						
(7) (8)		neats from matured de-boned meat to be prov 2010, with the entry ' <b>F</b> '. The matured de-bone	vided when required in column 5 'SG' of Part 1 d meat shall not be authorised for importation						
Of	ficial veterinarian								
	Name (in capital letters):	Qualificatio	on and title:						
	Date:	Signature:							
	Stamp:								

	co	Mode	I RUW Veterinary certificate to EU				
		Consignor	I.2. Certificate reference number I.2.a.				
		Name					
		Address	I.3. Central Competent Authority				
Ŧ		Tel. No	I.4. Local Competent Authority				
mer	1.5.	Consignee	1.6.				
sign		Name					
con		Address					
hed		Postal code					
pato		Tel. No					
Part I: Details of dispatched consignment	I.7.	Country ISO I.8. Region Code of origin	I.9. Country of ISO I.10. Region of Code destination code destination				
t I: Deta	I.11.	Place of origin	I.12.				
		Name Approval number					
Pai		Address					
	I.13	. Place of loading	I.14. Date of departure				
		· · · · · · · · · · · · · · · · · · ·					
	l.15	. Means of transport Aeroplane Ship Railway wagon	I.16. Entry BIP in EU				
		Road vehicle Other					
		Identification: Documentary references:	l.17.				
	l.18	. Description of commodity	I.19. Commodity code (HS code)				
			I.20. Quantity				
	I.21	. Temperature of product	I.22. Number of packages				
		Ambient Chiled	Frozen				
	1.23	. Identification of container/seal number	I.24. Type of packaging				
	I.25	Commodities certified for:	· · · · · · · · · · · · · · · · · · ·				
	1.26		I.27. For import or admission into EU				
	1.28	L Identification of the commodities					
			roval number establishments Number Net of packages weight r Cutting plant Cold store				

	COUNTRY			Model RUW			
	II. Hea	alth information	II.a. Certificate reference number	II.b.			
ation	ll.1. Put	lic Health Attestation					
	No anii <i>Ovi</i>	178/2002, (EC) No 852/2004 nals of the order Artiodactyla s aries, Capra hircus, Suida	erinarian, declare that I am aware of the re I, (EC) No 853/2004 and (EC) No 854/2004 a (excluding bovine animals (including <i>Bison</i> ar e and Tayassuidae), and of the families Rhin ace with those requirements, in particular that:	nd hereby certify that the fresh meat of wild ad <i>Bubalus</i> species and their cross-breeds), locerotidae and Elephantidae described in			
Part II: Certification	II.1.	1 the meat comes from accordance with Regula	(an) establishment(s) implementing a progra tion (EC) No 852/2004;	amme based on the HACCP principles in			
Part II:	II.1.	2 the meat has been obta 853/2004, and in particu	ained in compliance with the conditions set o lar:	out in Section IV of Annex III to Regulation			
		(i) before skinning, it ha	is been stored and handled separately from ot	her food and not frozen;			
		and					
		(ii) after skinning, it has	undergone a final inspection as referred to in p	point II.1.4;			
	(1) II.1.		le species, the meat fulfils the requirements of controls for Trichinella in meat;]	Regulation (EC) No 2075/2005 laying down			
	II.1.		d fit for human consumption following a post-m I and Chapters VIII and IX of Section IV of An				
	II.1.	()	ase of large wild game, the carcass or parts of accordance with Chapter III of Section I of Ann				
		.,	kages of meat have been marked with an identi to Regulation (EC) No 853/2004;]	fication mark in accordance with Section I of			
	II.1.	6 the meat satisfies the r foodstuffs;	elevant criteria set out in Regulation (EC) N	o 2073/2005 on microbiological criteria for			
	II.1.	0	ring live animals and products thereof provided by the residue plans submitted in accordance EC, and in particular Article 29 thereof, are fulfilled.				
	(¹) (²) [II.1	8 with regard to Chronic W	asting Disease (CWD):				
		have been examined fo method recognised by th	is derived exclusively from meat, excluding offal r Chronic Wasting Disease by histopathology, le competent authority with negative results an asting Disease has been confirmed in the last t	immunohistochemistry or other diagnostic d is not derived from animals coming from a			
	II.1.	9 the meat has been store Regulation (EC) No 853/	d and transported in accordance with the relev 2004.	vant requirements of Section I of Annex III to			
	II.2. <b>Ani</b>	mal Health attestation					
	I, th	e undersigned official veterin	arian, hereby certify, that the fresh meat descri	bed in Part I:			
	II.2.	1 has been obtained in the	e territory/ies with code:	which, at the date of issuing this certificate:			
		<ul><li>(a) has been free for 12 has taken place, and</li></ul>	months from rinderpest, and during the same I	e period no vaccination against this disease			
	(1) either	[(b) has been free for 12 this disease has take	months from foot-and-mouth disease, and du an place;]	ring the same period no vaccination against			
l							

. Healt	h information	II.a. Certificate reference number	II.b.					
having had cases			e since(dd/mm/yyyy), withou d to export these animals by Commission Regulation yyy);]					
(1) (4) or		on programmes against foot-and-mouth dise bovine animals;]	ase are being officially carried out and controlled i					
II.2.2			ween(dd/mm/yyyy) an y referred to in point II.2.1, and the killing took place:					
		nce that exceeds 20 km from the borders of a co importing this fresh meat into the Union,	untry or part thereof, which is not authorised during th					
		(b) in an area where during the last 60 days, there has been no restrictions for the diseases referred to in point II.2.1;						
II.2.3	game-handlir diseases refe of meat for im	g establishment around which, within a radiu rred to in point II.2.1 during the previous 30 day	sported as soon as possible for chilling to an approve is of 10 km, there has been no case/outbreak of th ys or, in the event of a case of disease, the preparation nly after removal of all meat, and the total cleaning an ial veterinarian;					
II.2.4								
	(1) either	[has been obtained and prepared without cor required above.]	ntact with other meats not complying with the condition					
	(1) (4) or	carcasses in which the main accessible lyn submitted to maturation at a temperature ab	de-boned meat other than offal that was obtained fro nphatic glands have been removed, which have bee ove +2 °C for at least 24 hours before the bones we meat was below 6.0 when tested electronically in the raturation and before de-boning, and					
			t not conforming to the requirements set out in th n, de-boning and storage until it has been packed ated areas.]					
	(1) (6) <i>or</i>	carcasses in which the main accessible lyn	de-boned meat other than offal that was obtained fro nphatic glands have been removed, which have been ove $+2$ °C for at least 24 hours before the bones we					
			t not conforming to the requirements set out in th n, de-boning and storage until it has been packed ated areas.]					
otes								
nimals (includi	ng <i>Bison</i> and <i>Bu</i>		ild animals of the order Artiodactyla (excluding bovir ies, Capra hircus, Suidae and Tayassuidae), and of th					
esh meat mea	ans all animal pa	rts fit for human consumption whether fresh, ch	illed or frozen.					
ter importatio	n unskinned car	casses must be conveyed without delay to the	processing establishment of destination					

After importation, unskinned carcasses must be conveyed without delay to the processing establishment of destination.

\_\_\_\_

СС	COUNTRY Model RUV								
II.	Health information	II.a. Certificate reference number	II.b.						
Pa	Part I:								
	<ul> <li>Box reference I.8: Provide the code of territory as appearing in Part 1 of Annex II to Regulation (EU) No 206/2010.</li> <li>Box reference I.11: Place of origin: name and address of the dispatch establishment.</li> <li>Box reference I.15: Registration number (railway wagons or container and lorries), flight number (aircraft) or name (ship) is to be provided. In case of unloading and reloading, the consignor must inform the BIP of entry into the Union.</li> <li>Box reference I.19: Use the appropriate HS code: 02.01, 02.02, 02.04, 02.06, 02.08.90 or 05.04.</li> <li>Box reference I.20: Indicate total gross weight and total net weight.</li> <li>Box reference I.23: For containers or boxes, the container number and the seal number (if applicable) should be included.</li> <li>Box reference I.28: <i>Nature of commodity</i>: Indicate 'carcass-whole', 'carcass-side', 'carcass-quarters' or 'cuts'.</li> <li>Box reference I.28: <i>Treatment type</i>: If appropriate, indicate 'matured' or 'unskinned'. If frozen, indicate the date of freezing (mm/yy) of the cuts/pieces.</li> <li>Box reference I.28: <i>Abattoir</i>: any abattoir or game handling establishment.</li> </ul>								
Pa	rt II:								
(²)	of Annex II to Regulation (EU) No 206	2010, with the entry 'G'.	e provided when required in column 5 'SG' of Part 1						
( <sup>3</sup> )	, ,,	• • • •							
0	Part 1 of Annex II to Regulation (EU) The matured de-boned meat shall not	Supplementary guarantees regarding meat from matured de-boned meat to be provided when required in column 5 'SG' of Part 1 of Annex II to Regulation (EU) No 206/2010 with the entry 'A'. The matured de-boned meat shall not be authorised for importation into the Union until 21 days after the date of killing of the							
( <sup>5</sup> )	for importation into the Union of the thi	rd country, territory or part thereof refe	killed or hunted either prior to the date of authorisation prred to in boxes I.7 and I.8, or during a period where meat from this third country, territory or part thereof.						
(6)		10, with the entry 'F'. The matured de-	e provided when required in column 5 'SG' of Part 1 of boned meat shall not be allowed for importation into						
Off	ficial veterinarian								
	Name (in capital latters):	<b>•</b> ••	alification and title:						
	Name (in capital letters):	Qu	aincation and title:						
	Date:	Sig	nature:						
	Stamp:								

		.,			Mod	el SUF						
	COUNTR									Veterinary certif	ficate to EU	
	I.1. Cons	0				I.2. Certificate reference number I.2.a.						
	Nam					I.3. C	Central	Competer	t Authority	y		
	Addr	Address					ocal Cr	ompetent	Authority			
ent	Tel. N	Tel. No						Inpetent	Authority			
gnm	I.5. Cons	5. Consignee										
nsi	Nam	Name										
d co	Addr	Address										
tche	Posta	Postal code										
spa	Tel. N	Tel. No										
Part I: Details of dispatched consignment	I.7. Cour of ori		ISO code	I.8. Region of origin	Code		Country lestinat		ISO code	I.10. Region of destination	Code	
Deta	I.11. Place	e of origin				I.12.						
rt I:	Nam			Approval number								
Pa	Addr	ess						_				
	I.13. Place	e of loading				I.14. [	Date of o	departure				
	I.15. Mea	ns of transpo	ort			I.16. E	Entry BI	P in EU				
	Aero	plane 🗌	Shi	p 📃 🛛 Railway wage	on 🗌							
	Road	l vehicle 🗌	Othe	er 🗌								
	Ident	ification:				1.17.						
		imentary ref	erences:									
	I.18. Desc	ription of co	mmodity			I.19. Commodity code (HS code)						
							L		I.20.	Quantity		
	I.21. Tem	perature of p	roduct						1.22.	Number of packages		
		bient		Chiled 🗌								
						Frozen						
	I.23. Ident	ification of c	ontainer/se	eal number					1.24.	Type of packaging		
		modities cer an consump										
	I.26.					I.27. For import or admission into EU						
	I.28. Ident	ification of th	he commo	dities								
		ecies ific name)	Nature commo		App Abatto		mber e utting p	stablishme lant Co	ents Id store	Number of packages	Net weight	

	COUNT	FRY				Model SUF
	П.	Health	information		II.a. Certificate reference number	II.b.
-	II.1.	Public	Health Attes	tation		
Ition		(EC) N animal those	No 852/2004, ( Is belonging to requirements, i	EC) No 853 the Suidae in particular	3/2004 and (EC) No 854/2004 and hereby e, Tayassuidae, or Tapiridae families descrit that:	t provisions of Regulations (EC) No 178/2002, certify that the meat of farmed non-domestic bed in Part I was produced in accordance with
ITITICa		II.1.1			tion (EC) No 852/2004;	gramme based on the HACCP principles in
Part II: Certification		II.1.2	the meat has No 853/2004		ined in compliance with the conditions set o	out in Section III of Annex III to Regulation (EC)
<b>1</b>		II.1.3				i laying down specific rules on official controls amination by a digestion method with negative
		II.1.4		with, Chap		e and post-mortem inspections carried out in ( of Section IV of Annex I to Regulation (EC)
		II.1.5	(¹) either		cass or parts of the carcass have been m r III of Section I, of Annex I to Regulation (EC	narked with a health mark in accordance with ) No 854/2004;]
			(1) or		kages of meat have been marked with an ide I to Regulation (EC) No 853/2004;]	entification mark in accordance with Section I of
		II.1.6	the meat sa foodstuffs;	tisfies the r	relevant criteria set out in Regulation (EC)	No 2073/2005 on microbiological criteria for
		II.1.7			live animals and products thereof provided and in particular Article 29 thereof, are fulfil	l by the residue plans submitted in accordance led;
		II.1.8	the meat has Regulation (I			elevant requirements of Section I of Annex III to
	II.2.	Anima	Il Health attes	tation		
		I, the u	indersigned of	ficial veterin	narian, hereby certify, that the fresh meat dea	scribed in Part I:
		II.2.1	has been ob	tained in the	e territory/ies with code:	which, at the date of issuing this certificate:
			(1) either		been free for 12 months from foot-and-m sical swine fever, swine vesicular disease, a	outh disease, rinderpest, African swine fever, nd]
			(1) or		has been free for 12 months from rinderpest, , [classical swine fever] (1) and [swine vesicul	African swine fever, [foot-and-mouth disease] ('), ar disease] ('), and
					[swine vesicular disease] (1), since	outh disease] ( <sup>1</sup> ), [classical swine fever] ( <sup>1</sup> ) and 
				imp		nst these diseases have been carried out and nst these diseases are not permitted in this
		II.2.2	has been ob	tained from	animals that:	
			(1) either	-	emained in the territory described under poi before slaughter;]	nt II.2.1 since birth, or for at least the last three

Health	information		II.a. Certificate reference number	II.b.
	(1) <i>or</i>	point II.2		(dd/mm/yyyy) into the territory described unde
II.2.3	has been ob	tained from	animals coming from holdings:	
	(a) in which point II.2		he animals present therein have been va	accinated against the diseases referred to i
			n in an area of 10 km radius, there has been e previous 40 days,	no case/outbreak of the diseases referred to i
	and, the		are not subject to prohibition as a result of	se diseases transmissible to humans or animal f an outbreak of porcine brucellosis during th
II.2.4	has been ob	tained from	animals which:	
	(1) either	to a		chicles, cleaned and disinfected before loading with other animals which did not comply with th
		· ·	ughter and, in particular, have shown no evi	em health inspection during the 24 hours befor dence of the diseases referred to in point II.2.
			re been slaughtered on /mm/yyyy) and(dd/r	(dd/mm/yyyy) or between nm/yyyy) (³);]
	(1) or		e been slaughtered on the holding of origin, ponsible for the holding, who has provided a	following authorisation by an official veterinaria written statement that:
		_	in his opinion an unacceptable risk would h to their handlers by the transport of the anir	ave been posed to the welfare of the animals on nais to an slaughterhouse,
		_	the holding had been inspected and authori of game,	ised by the competent authority for the slaughte
		_	•	n health inspection during the 24 hours befor wn no evidence of the diseases referred to
		_	the animals were slaughtered between(dd/mm/yyyy), (3)	(dd/mm/yyyy) ar
		_	the bleeding of the animals was performed	correctly, and
		-	the slaughtered animals were eviscerated v	within three hours of the time of slaughter, and
		cor terr	ditions and, where more than one ho	he approved slaughterhouse under hygien ur elapsed since the time of slaughter, been found on the arrival of the vehicle use
II.2.5	has been ob	tained from	animals that have remained separate since	birth from wild cloven-hoofed animals;
II.2.6	of the disea preparation	ses referre of meat for Ill meat, an	d to in point II.2.1 during the previous 40 c importation into the Union has been author	ius of 10 km, there has been no case/outbrea days or, in the event of a case of disease, th ised only after slaughter of all animals preser e establishment under the control of an offici
II.2.7	has been ob certificate.	tained and <sub>l</sub>	prepared without contact with other meats no	ot complying with the requirements set out in th

COUNT	RY		Model SUF
II.	Health information	II.a. Certificate reference number	II.b.
II.3.		arian, hereby certify, that the fresh meat descri se before and at the time of slaughter or killin	
Notes			
	tificate is meant for fresh meat, ex e families that are domestically kept	cluding offal and minced meat, of wild anima or bred since birth in farms.	ls belonging to the Suidae, Tayassuidae, or
Fresh m	eat means all animal parts fit for hu	man consumption, whether fresh, chilled or fro	ozen.
Part I:			
<ul> <li>Box prov</li> <li>Box prov</li> <li>Box</li> <li>Box</li> <li>Box</li> <li>Box</li> <li>Box</li> <li>Box</li> <li>Box</li> <li>Cod</li> <li>Cod</li> <li>(3) Date of an period</li> </ul>	reference I.11: Place of origin: name reference I.15: Registration numbe ided. In case of unloading and reloa reference I.19: Use the appropriate reference I.20: Indicate total gross reference I.23: For containers or bo reference I.28: <i>Nature of commodit</i> reference I.28: <i>Treatment type</i> : If ap cuts/pieces. p as appropriate e of the territory as it appears in Par e or dates of slaughter. Imports of thi uthorisation for importation into the U	erritory as appearing in Part 1 of Annex II to Re e and address of the dispatch establishment. r (railway wagons or container and lorries), fli ading, the consignor must inform the BIP of er HS code: 02.03, 02.08.90 or 05.04. weight and total net weight. xes, the container number and the seal numb y: Indicate 'carcass-whole', 'carcass-side', 'car opropriate indicate deboned, or bone-in. If fro rt 1 of Annex II to Regulation (EU) No 206/201 s meat shall not be allowed when obtained fro Jnion of the third country, territory or part there been adopted by the Union against imports of	ight number (aircraft) or name (ship) is to be try into the Union. er (if applicable) should be included. rcass-quarters' or 'cuts'. zen, indicate the date of freezing (mm/yy) of 0. m animals slaughtered either prior to the date eof referred to in boxes I.7 and I.8, or during a
Official v	reterinarian		
	Name (in capital letters):	Qualification	n and title:
	Date:	Signature:	
	Stamp:		

	~~	UNTRY			Mode	elsuw			Votorinory contifi	ianto to Ell
		Consignor				I.2. Certifi	cate reference	ce numbe	Veterinary certifier I.2.a.	
		Name								
		Address				I.3. Centra	al Competen	t Authority	У	
Ħ		Tel. No				I.4. Local	Competent A	Authority		
nme	1.5.	Consignee				1.6.				
nsig		Name								
d co		Address								
tche		Postal code								
spat		Tel. No								
Part I: Details of dispatched consignment	I.7.	Country of origin	ISO code	I.8. Region of origin	Code	I.9. Count destin		ISO code	I.10. Region of destination	Code
Deta	I.11.	Place of origin				I.12.				
art I:		Name		Approval number						
Å		Address								
	I.13	. Place of loading				I.14. Date c	of departure			
	I.15	. Means of transpo	ort			I.16. Entry	BIP in EU			
		Aeroplane 🗌	Sh	ip 🗌 🛛 Railway wag	on 🗌					
		Road vehicle	Oth	er						
		Identification:				l.17.				
		Documentary ref	erences:							
	I.18	. Description of co	mmodity				I.19. Com	nmodity c	ode (HS code)	
								1.20.	Quantity	
	I.21	. Temperature of p	roduct					1.22.	Number of packages	
		Ambient		Chiled		Frozen [				
	1.23	. Identification of c	ontainer/s	eal number				1.24.	Type of packaging	
	1.25	. Commodities cer Human consump						1		
	1.26					I.27. For im	port or admi	ssion into	EU	
	1.28	. Identification of t	ne commo	dities		1				
	10	Species Scientific name)	Nature commo		Арр	roval number	establishme	ents	Number of packages	Net weight
	(0	Scientific hame)	comme	odity type	Abatto	ir Cutting	plant Col	d store	of packages	weight
					, iourio	Sutting				

Model SUW

	COUNT	RY				Model SUW
	Ш.	Health	information		II.a. Certificate reference number	II.b.
	II.1.	Public	Health Attesta	tion		
u		(EC) N the Su	lo 852/2004,(EC	) No 853/	arian declare that I am aware of the relevant requ 2004 and (EC) No 854/2004 and hereby certify ridae families described in Part I was produced	y that the meat of wild animals belonging to
rtificati		II.1.1			(an) establishment(s) implementing a progra tion (EC) No 852/2004;	mme based on the HACCP principles in
Part II: Certification		II.1.2	the meat has particular:	been obta	ained in accordance with Section IV of Annex	III to Regulation (EC) No 853/2004, an in
Ра			(i) before skin	ning, it ha	as been stored and handled separately from oth	ner food and not frozen;
			and			
			(ii) after skinni	ng, it has	undergone a final inspection as referred to in p	oint II.1.4;
		II.1.3			irements of Regulation (EC) No 2075/2005 lay nd in particular, has been subject to an exami	
		II.1.4			d fit for human consumption following a post-m n I and Chapters VIII and IX of Section IV of Anr	
		II.1.5	(1) either		cass or parts of the carcass have been mark III of Section I of Annex I to Regulation (EC) No	
			(1) <i>or</i>		kages of meat have been marked with an identi to Regulation (EC) No 853/2004;]	fication mark in accordance with Section I of
		II.1.6	the meat satis foodstuffs;	fies the r	elevant criteria set out in Regulation (EC) No	o 2073/2005 on microbiological criteria for
		II.1.7			live animals and products thereof provided by and in particular Article 29 thereof, are fulfilled.	
		II.1.8	the meat has b Regulation (EC		ed and transported in accordance with the relev /2004	rant requirements of Section I of Annex III to
	II.2.	Anima	I Health attesta	tion		
		I, the u	indersigned offic	ial veterin	arian, hereby certify, that the fresh meat descril	bed in Part I:
		II.2.1	has been obtai	ned in the	e territory/ies with code: (²) which, a	t the date of issuing this certificate:
			(1) either		been free for 12 months from foot-and-mout sical swine fever, swine vesicular disease, and ]	
			(1) or		has been free for 12 months from rinderpest, Afric [classical swine fever] (¹) and [swine vesicular d	
					has been considered free from [foot-and-mout [swine vesicular disease] ('), since cases/outbreaks afterwards, and authorised to (EU) No/, of	(dd/mm/yyyy), without having had export this meat by Commission Regulation
				imp	ng the last 12 months no vaccination against orts of domestic animals vaccinated against tory;	

	information		II.a. Certificate reference number	II.b.
II.2.2				en (dd/mm/yyyy) an rred to in point II.2.1, and the killing took place:
			eds 20 km from the borders of a cour s fresh meat into the Union,	try or part thereof, which is not authorised during thi
	(b) in an area point II.2.1		ing the last 60 days, there has be	een no restrictions for the diseases referred to i
II.2.3.A	centre, and im of 10 km, there in the event of	mediately at e has been r f a case of d	terwards] ( <sup>1</sup> ) to an approved game-h o case/outbreak of the diseases refe isease, the preparation of meat for i	sported within 12 hours for chilling [to a collectio andling establishment around which, within a radiu erred to in point II.2.1 during the previous 40 days o mportation into the Union has been authorised on a of the establishment under the control of an officia
(1) (4) [II.2.3.B	has been obta negative result		rcasses on which the following test f	or classical swine fever was carried out and provide
	(1) either	[virus isola	tion from blood (EDTA);]	
	(1) <i>or</i>	[virus isola	tion from samples of	
	(1) <i>or</i>	[immunofl	uorescence for viral antigen on samp	les of;
II.2.4	has been obta certificate.	ined and pre	pared without contact with other me	ats not complying with the conditions required in th
lotes	mont for fool		uling offel and minand most of wil	d opimals balancing to the Quideo. Tousequideo
his certificate is	s meant for fresh s that are killed c			d animals belonging to the Suidae, Tayassuidae, (
his certificate is apiridae families	s that are killed c	or hunted in t		
his certificate is apiridae families resh meat mear	s that are killed c ns all animal par	or hunted in t ts fit for hum	he wild. an consumption whether fresh, chille	
nis certificate is apiridae families resh meat mear fter importation	s that are killed c ns all animal par	or hunted in t ts fit for hum	he wild. an consumption whether fresh, chille	ed or frozen.
his certificate is apiridae families resh meat mear fter importation, <b>art I:</b>	s that are killed c ns all animal pari , unskinned carc	or hunted in t ts fit for hum casses must	he wild. an consumption whether fresh, chille	ed or frozen.
nis certificate is apiridae families resh meat mear fter importation <b>art I:</b> - Box reference	s that are killed c Is all animal part , unskinned carc e I.8: Provide the	or hunted in t ts fit for hum casses must e code of ter	he wild. an consumption whether fresh, chille be conveyed without delay to the pro	ed or frozen. ocessing establishment of destination. k II to Regulation (EU) No 206/2010.
nis certificate is apiridae families esh meat mear iter importation <b>art I:</b> - Box reference - Box reference - Box reference	s that are killed c Is all animal part , unskinned carc e I.8: Provide the e I.11: Place of c e I.15: Registrati	or hunted in t ts fit for hum casses must e code of ter origin: name ion number	he wild. an consumption whether fresh, chille be conveyed without delay to the pro ritory as appearing in Part 1 of Anner and address of the dispatch establis	ed or frozen. occessing establishment of destination. < II to Regulation (EU) No 206/2010. hment. rries), flight number (aircraft) or name (ship) is to b
nis certificate is piridae families esh meat mear iter importation, art I: - Box reference - Box reference - Box reference provided. In c	a that are killed c ns all animal part , unskinned carc e I.8: Provide the e I.11: Place of c e I.15: Registrat case of unloadin	or hunted in t ts fit for hum casses must e code of ter origin: name ion number g and reload	he wild. an consumption whether fresh, chille be conveyed without delay to the pro ritory as appearing in Part 1 of Anne: and address of the dispatch establis (railway wagons or container and loo	ed or frozen. occessing establishment of destination. < II to Regulation (EU) No 206/2010. hment. rries), flight number (aircraft) or name (ship) is to b
nis certificate is piridae families esh meat mear ter importation art I: • Box referenc • Box referenc provided. In c • Box referenc	a that are killed c ns all animal part , unskinned carc e I.8: Provide the e I.11: Place of c e I.15: Registrati case of unloadin e I.19: Use the a	or hunted in t ts fit for hum casses must e code of ter origin: name ion number g and reload	he wild. an consumption whether fresh, chille be conveyed without delay to the pro- ritory as appearing in Part 1 of Anne: and address of the dispatch establis (railway wagons or container and loo ing, the consignor must inform the E	ed or frozen. occessing establishment of destination. k II to Regulation (EU) No 206/2010. hment. rries), flight number (aircraft) or name (ship) is to b
his certificate is apiridae families resh meat mear fter importation <b>art I:</b> - Box reference - Box reference provided. In o - Box reference - Box reference - Box reference	a that are killed c as all animal part , unskinned carc e I.8: Provide the e I.11: Place of c e I.15: Registrat case of unloadin e I.19: Use the a e I.20: Indicate to	or hunted in t ts fit for hum casses must e code of ter origin: name ion number g and reload uppropriate H otal gross w	he wild. an consumption whether fresh, chille be conveyed without delay to the pro- ritory as appearing in Part 1 of Anne: and address of the dispatch establis (railway wagons or container and loo ing, the consignor must inform the E IS code: 02.03, 02.08.90 or 05.04. eight and total net weight.	ed or frozen. occessing establishment of destination. < II to Regulation (EU) No 206/2010. hment. rries), flight number (aircraft) or name (ship) is to b
his certificate is apiridae families resh meat mear fter importation <b>art I:</b> - Box reference - Box reference - Box reference - Box reference - Box reference - Box reference - Box reference	a that are killed c ns all animal part , unskinned carc e I.8: Provide the e I.11: Place of c e I.15: Registrati case of unloadin e I.19: Use the a e I.20: Indicate to e I.23: For conta	or hunted in t ts fit for hum casses must e code of ter origin: name ion number g and reload uppropriate H otal gross w iners or box	he wild. an consumption whether fresh, chille be conveyed without delay to the pro- ritory as appearing in Part 1 of Anne: and address of the dispatch establis (railway wagons or container and loo ing, the consignor must inform the E IS code: 02.03, 02.08.90 or 05.04. eight and total net weight.	ed or frozen. Decessing establishment of destination. (II to Regulation (EU) No 206/2010. hment. rries), flight number (aircraft) or name (ship) is to b IP of entry into the Union.
his certificate is apiridae families resh meat mear fter importation <b>art I:</b> - Box reference - Box reference	that are killed c s all animal part , unskinned carc e I.8: Provide the e I.11: Place of c e I.15: Registrat case of unloadin e I.19: Use the a e I.20: Indicate to e I.23: For conta e I.28: <i>Nature of</i> e I.28: <i>Treatmen</i>	or hunted in t ts fit for hum casses must e code of ter origin: name ion number g and reload uppropriate F otal gross w iners or box	he wild. an consumption whether fresh, chille be conveyed without delay to the pro- ritory as appearing in Part 1 of Annez and address of the dispatch establis (railway wagons or container and loo ing, the consignor must inform the E IS code: 02.03, 02.08.90 or 05.04. eight and total net weight. as, the container number and the sea Indicate 'carcass-whole', 'carcass-s	ocessing establishment of destination. « Il to Regulation (EU) No 206/2010. hment. rries), flight number (aircraft) or name (ship) is to b IP of entry into the Union.

со	UNTRY		Model SUW
II.	Health information	II.a. Certificate reference number	II.b.
(1) (2) (3)	for importation into the Union of the third where restrictive measures have been thereof. Supplementary guarantees to be provid with the entry ' <b>C</b> '. For such purpose, in	uthorised when obtained from anima I country, territory or part thereof refe adopted by the Union against impo ded when required in column 5 'SG' tests other than EDTA, the samples uple of at least one of the following	o 206/2010. Is killed or hunted either prior to the date of authorisation rred to in boxes reference 1.7 and 1.8, or during a period rts of this meat from this third country, territory or part of Part 1 of Annex II to Regulation (EU) No 206/2010, s to be used are a sample of tonsil and of spleen plus lymph nodes: retropharyngeal, parotid, mandibular or
Off	icial veterinarian		
	Name (in capital letters):		ualification and title:
	Date:	S	gnature:
	Stamp:		

	COUNTRY	Mod	el EQW	Veterinary certificate to EU
	I.1. Consignor		I.2. Certificate reference r	
	Name			
	Address		I.3. Central Competent Au	uthority
Ŧ	Tel. No		I.4. Local Competent Auth	nority
men	I.5. Consignee		1.6.	
sign	Name			
con	Address			
hed	Postal code			
patc	Tel. No			
Part I: Details of dispatched consignment	I.7. Country ISO of origin code	5		SO I.10. Region of Code destination
etai	I.11. Place of origin		I.12.	
t I: D	Name	Approval number		
Par	Address			
-	I.13. Place of loading		I.14. Date of departure	
	1.13. Flace of loading			
	I.15. Means of transport Aeroplane	Ship 🗌 Railway wagon 🗌	I.16. Entry BIP in EU	
	Road vehicle	Other		
	Identification: Documentary reference	s:	1.17.	
-	I.18. Description of commodi	ty	I.19. Commo	odity code (HS code)
				I.20. Quantity
	I.21. Temperature of product			I.22. Number of packages
	Ambient	Chiled	Frozen	
	I.23. Identification of containe	er/seal number		I.24. Type of packaging
-	I.25. Commodities certified for Human consumption	pr:		
	1.26.		I.27. For import or admission	on into EU
ŀ	I.28. Identification of the com	modities	I	
	Species (Scientific name)	Nature of Approval no commodity	umber establishments	Number Net of packages weight
		Abattoir C	Cutting plant Cold store	

				Model EC
П.	Health	information	II.a. Certificate reference number	II.b.
II.1.	Public	Health Attestation	 on	
	(EC) N	lo 852/2004, (EC)	veterinarian, declare that I am aware of the relevant No 853/2004 and (EC) No 854/2004 and hereby gris (zebra) described in Part I was produced in ac	certify that the meat of wild solipeds belonging
	II.1.1		s from (an) establishment(s) implementing a pro Regulation (EC) No 852/2004;	ogramme based on the HACCP principles in
	II.1.2	the meat was ob	tained in compliance with Section IV of Annex III to	Regulation (EC) No 853/2004;
	II.1.3		ne requirements of Regulation (EC) No 2075/2005 at, in particular, has been subject to an examination	
	II.1.4		en found fit for human consumption following a po f Section I and Chapters VIII and IX of Section IV of	•
	II.1.5	.,	[the carcass or parts of the carcass have been r Chapter III of Section I of Annex I to Regulation (EC	
		.,	[the packages of meat have been marked with an ic Annex II to Regulation (EC) No 853/2004;]	entification mark in accordance with Section I
	II.1.6	the meat satisfic foodstuffs;	es the relevant criteria set out in Regulation (EC	) No 2073/2005 on microbiological criteria fo
	II.1.7	0	covering live animals and products thereof provide /23/EC, and in particular Article 29 thereof, are fulfi	, ,
	II.1.8	the meat has be Regulation (EC)	en stored and transported in accordance with the No 853/2004.	elevant requirements of Section I of Annex III t
II.2.	Anima	l Health attestati	on	
	l, the u	ndersigned officia	l veterinarian, hereby certify, that the fresh meat de	scribed in Part I:
	II.2.1		ed from wild animals that were killed between (dd/mm/yyyy) (2) inside the territory/ies with	
		•••••		
	II.2.2	has been obtain centre, and immo of 10 km, there h the event of a ca	ed from wild animals which after killing were transp ediately afterwards] ( <sup>1</sup> ) to an approved game-hand has been no case/outbreak of African horse sickne se of such diseases, the preparation of meat for ex all meat, and the total cleaning and disinfection of t	orted within 12 hours for chilling [to a collection ing establishment around which, within a radiu as or glanders during the previous 40 days or, portation to the Union has been authorised on
	II.2.2 II.2.3	has been obtain centre, and imm of 10 km, there h the event of a ca after removal of a veterinarian;	ediately afterwards] (1) to an approved game-hand has been no case/outbreak of African horse sickne se of such diseases, the preparation of meat for ex	orted within 12 hours for chilling [to a collection ing establishment around which, within a radii ss or glanders during the previous 40 days or, portation to the Union has been authorised on the establishment under the control of an offici
		has been obtain centre, and imm of 10 km, there h the event of a ca after removal of a veterinarian; has been obtaine	ediately afterwards] (1) to an approved game-hand has been no case/outbreak of African horse sickne se of such diseases, the preparation of meat for ex all meat, and the total cleaning and disinfection of t	orted within 12 hours for chilling [to a collection ing establishment around which, within a radiu ss or glanders during the previous 40 days or, portation to the Union has been authorised on he establishment under the control of an offici
Notes		has been obtain centre, and imm of 10 km, there h the event of a ca after removal of a veterinarian; has been obtaine	ediately afterwards] (1) to an approved game-hand has been no case/outbreak of African horse sickne se of such diseases, the preparation of meat for ex all meat, and the total cleaning and disinfection of t	orted within 12 hours for chilling [to a collectic ing establishment around which, within a radiu ss or glanders during the previous 40 days or, portation to the Union has been authorised on he establishment under the control of an offici
This ce	II.2.3	has been obtain centre, and imm of 10 km, there h the event of a ca after removal of a veterinarian; has been obtaine certificate.	ediately afterwards] (1) to an approved game-hand has been no case/outbreak of African horse sickne se of such diseases, the preparation of meat for ex all meat, and the total cleaning and disinfection of t	norted within 12 hours for chilling [to a collectic ing establishment around which, within a radiu so or glanders during the previous 40 days or, portation to the Union has been authorised on he establishment under the control of an offici ot complying with the requirements set out in th
This ce (zebra).	II.2.3	has been obtain centre, and imm of 10 km, there h the event of a ca after removal of a veterinarian; has been obtaine certificate.	ediately afterwards] (1) to an approved game-hand has been no case/outbreak of African horse sicknes se of such diseases, the preparation of meat for ex all meat, and the total cleaning and disinfection of t ed and prepared without contact with other meats n	orted within 12 hours for chilling [to a collectio ing establishment around which, within a radiu so or glanders during the previous 40 days or, i portation to the Union has been authorised onl he establishment under the control of an officia of complying with the requirements set out in thi objects belonging to the subgenus <i>Hippotigri</i>

	Health information	II.a. Certificate reference number	II.b.
art I:			
		of territory as appearing in Part 1 of Annex I	Lto Begulation (ELI) No 206/2010
		name and address of the dispatch establish	
	•	and an and the second on the second sec	es), flight number (aircraft) or name (ship) is to be
pr	ovided. In case of unloading and r	reloading, the consignor must inform the BIF	P of entry into the Union.
	ox reference I.19: Use the appropr		
	ox reference I.20: Indicate total gro	0	
		or boxes, the container number and the seal	· · · · · · ·
		odity: Indicate 'carcass-whole', 'carcass-side	e, carcass-quarters or cuts. ed'. If frozen, indicate the date of freezing (mm/yy)
of	the cuts/pieces.		ed . If hozen, indicate the date of heezing (him/yy)
- Bo	ox reference I.28: <i>Abattoir</i> : any aba	attoir or game handling establishment.	
art II	:		
	eep as appropriate.		
foi	r importation into the Union of the	third country, territory or part thereof referre	led or hunted either prior to the date of authorisation ad to in boxes I.7 and I.8, or during a period where aat from this third country, territory or part thereof.
) Co	ode of the territory as it appears ir	n Part 1 of Annex II to Regulation (EU) No 20	6/2010.
fficia	ıl veterinarian		
	Name (in capital letters):	Qualif	ication and title:
	Date:	Signa	ture:
	Stamp:		

### ANNEX III

### Model TRANSIT/STORAGE

#### Veterinary certificate to EU

	COI	UNTRY	Veterinary certificate to EU
	I.1.	Consignor	I.2. Certificate reference number I.2.a.
		Name	I.3. Central Competent Authority
		Address	
ent		Tel. No	I.4. Local Competent Authority
gnm	1.5.	Consignee	I.6. Person responsible for the consignment in EU
onsi		Name	Name
∋q c		Address	Address
tche		Postal code	Postal code
lispe		Tel. No	Tel. No
Part I: Details of dispatched consignment	I.7.	Country ISO I.8. Region Code of origin code of origin	I.9. Country of ISO I.10. Region of Code destination code destination
Deta	I.11.	Place of origin	I.12. Place of destination
μ		Name Approval number	Custom warehouse Ship supplier
Pa		Address	Name Approval number
			Address Postal code
	I.13.	Place of loading	I.14. Date of departure
	I.15.	Means of transport	I.16. Entry BIP in EU
		Aeroplane Ship Railway wagon	
		Road vehicle Other	
		Identification: Documentary references:	I.17. No. (s) of CITES
	l.18.	Description of commodity	I.19. Commodity code (HS code)
			I.20. Quantity
	I.21.	. Temperature of product	I.22. Number of packages
		Ambient Chiled	Frozen
	1.23	. Identification of container/seal number	I.24. Type of packaging
	1.25	. Commodities certified for:	
		Human consumption	
	1.26	. For transit through EU to 3 rd Country	1.27.
		3rd country ISO code	
	1.28	. Identification of the commodities	
	(5	Scientific name) commodity type	Imber establishments Number Net of packages weight
			Cutting manufacturing plant/ plant

	TRY		Model TRANSIT/STORA
П.	Health information	II.a. Certificate reference number	II.b.
II.1.	Animal Health Attestation		
	I, the undersigned official veter	rinarian, hereby certify, that the fresh meat de	scribed in Part I:
		or region authorized for imports into the Unior he time of slaughter, and	n as laid down in Part 1 of Annex II to Regulatior
			n in the animal health attestation in the mode QW] (1) in Part 2 of Annex II to Regulation (EU)
		als which were slaughtered and processed	on (dd/mm/yyyy) o (dd/mm/yyyy) (²).
Notes This ce	rtificate is meant for transit and sto	prage in accordance with Article 12(4) or Artic	sle 13 of Directive 97/78/EC of:
This ce	rtificate is meant for transit and sto sh meat, including minced meat, o	-	cle 13 of Directive 97/78/EC of:
This ce	sh meat, including minced meat, o	-	
This ce — fres	sh meat, including minced meat, o domestic bovine animals (inclu	f:	oss-breeds) (Model 'BOV');
This ce — fres (1)	sh meat, including minced meat, o domestic bovine animals (inclu domestic ovine animals ( <i>Ovis</i> a	f: uding <i>Bubalus</i> and <i>Bison</i> species and their cro aries) or domestic caprine animals ( <i>Capra hir</i>	oss-breeds) (Model 'BOV');
This ce — fres (1) (2) (3)	sh meat, including minced meat, o domestic bovine animals (inclu domestic ovine animals ( <i>Ovis a</i> domestic porcine animals ( <i>Su</i> s	f: uding <i>Bubalus</i> and <i>Bison</i> species and their cro aries) or domestic caprine animals ( <i>Capra hird</i> <i>s scrofa</i> ) (Model 'POR');	oss-breeds) (Model 'BOV');
This ce — fres (1) (2) (3) — fres	sh meat, including minced meat, o domestic bovine animals (inclu domestic ovine animals ( <i>Ovis</i> domestic porcine animals ( <i>Sus</i> sh meat, excluding minced meat, o	f: uding <i>Bubalus</i> and <i>Bison</i> species and their cro aries) or domestic caprine animals ( <i>Capra hird</i> <i>s scrofa</i> ) (Model 'POR'); of:	oss-breeds) (Model 'BOV'); <i>cus</i> ) (Model 'OVI');
This ce — fres (1) (2) (3) — fres (4)	sh meat, including minced meat, o domestic bovine animals (inclu domestic ovine animals ( <i>Ovis a</i> domestic porcine animals ( <i>Sus</i> sh meat, excluding minced meat, o domestic solipeds ( <i>Equus cab</i>	f: uding <i>Bubalus</i> and <i>Bison</i> species and their cro aries) or domestic caprine animals ( <i>Capra hird</i> <i>s scrofa</i> ) (Model 'POR'); of: allus, Equus asinus and their cross-breeds) (I	oss-breeds) (Model 'BOV'); <i>cus</i> ) (Model 'OVI');
This ce — fres (1) (2) (3) — fres (4)	sh meat, including minced meat, o domestic bovine animals (inclu domestic ovine animals ( <i>Ovis a</i> domestic porcine animals ( <i>Sus</i> sh meat, excluding minced meat, o domestic solipeds ( <i>Equus cab</i> sh meat, excluding offal and mince farmed non-domestic animals their cross-breeds), <i>Ovis aries</i> ,	f: Juding <i>Bubalus</i> and <i>Bison</i> species and their cro aries) or domestic caprine animals ( <i>Capra hird</i> s scrofa) (Model 'POR'); of: <i>allus, Equus asinus</i> and their cross-breeds) (I ed meat, of: of the order Artiodactyla (excluding bovine ani	oss-breeds) (Model 'BOV'); <i>cus</i> ) (Model 'OVI'); Model 'EQU'); imals (including <i>Bison</i> and <i>Bubalus</i> species and
This ce — fres (1) (2) (3) — fres (4) — fres	sh meat, including minced meat, o domestic bovine animals (inclu domestic ovine animals ( <i>Ovis a</i> domestic porcine animals ( <i>Sus</i> sh meat, excluding minced meat, o domestic solipeds ( <i>Equus cab</i> sh meat, excluding offal and mince farmed non-domestic animals their cross-breeds), <i>Ovis aries</i> , (Model 'RUF'); wild non-domestic animals of	f: Juding Bubalus and Bison species and their cro aries) or domestic caprine animals (Capra hird s scrofa) (Model 'POR'); of: allus, Equus asinus and their cross-breeds) (I ed meat, of: of the order Artiodactyla (excluding bovine ani Capra hircus, Suidae and Tayassuidae), and o the order Artiodactyla (excluding bovine anin	oss-breeds) (Model 'BOV'); <i>cus</i> ) (Model 'OVI'); Model 'EQU'); imals (including <i>Bison</i> and <i>Bubalus</i> species and of the families Rhinocerotidae and Elephantidae nals (including <i>Bison</i> and <i>Bubalus</i> species and
This ce — fres (1) (2) (3) — fres (4) — fres (5)	sh meat, including minced meat, o domestic bovine animals (inclu domestic ovine animals ( <i>Ovis</i> a domestic porcine animals ( <i>Sus</i> sh meat, excluding minced meat, o domestic solipeds ( <i>Equus cab</i> sh meat, excluding offal and mince farmed non-domestic animals their cross-breeds), <i>Ovis aries</i> , (Model 'RUF'); wild non-domestic animals of their cross-breeds), <i>Ovis aries</i> , (Model 'RUW');	f: Juding Bubalus and Bison species and their cro aries) or domestic caprine animals (Capra hird s scrofa) (Model 'POR'); of: allus, Equus asinus and their cross-breeds) (I ed meat, of: of the order Artiodactyla (excluding bovine ani Capra hircus, Suidae and Tayassuidae), and o the order Artiodactyla (excluding bovine anin	oss-breeds) (Model 'BOV'); <i>cus</i> ) (Model 'OVI'); Model 'EQU'); imals (including <i>Bison</i> and <i>Bubalus</i> species and of the families Rhinocerotidae and Elephantidae nals (including <i>Bison</i> and <i>Bubalus</i> species and of the families Rhinocerotidae and Elephantidae
This ce — fres (1) (2) (3) — fres (4) — fres (5) (6)	sh meat, including minced meat, o domestic bovine animals (inclu domestic ovine animals ( <i>Ovis a</i> domestic porcine animals ( <i>Sus</i> sh meat, excluding minced meat, o domestic solipeds ( <i>Equus cab</i> sh meat, excluding offal and mince farmed non-domestic animals of their cross-breeds), <i>Ovis aries</i> , (Model 'RUF'); wild non-domestic animals of their cross-breeds), <i>Ovis aries</i> , (Model 'RUV'); farmed non-domestic animals	f: Juding Bubalus and Bison species and their cro aries) or domestic caprine animals (Capra hird as scrofa) (Model 'POR'); of: allus, Equus asinus and their cross-breeds) (I ad meat, of: of the order Artiodactyla (excluding bovine ani Capra hircus, Suidae and Tayassuidae), and of the order Artiodactyla (excluding bovine anin Capra hircus, Suidae and Tayassuidae), and of the order Artiodactyla (excluding bovine anin Capra hircus, Suidae and Tayassuidae), and of the order Artiodactyla (excluding bovine anin Capra hircus, Suidae and Tayassuidae), and of the order Artiodactyla (excluding bovine anin Capra hircus, Suidae and Tayassuidae), and of the order Artiodactyla (excluding bovine anin the order Artioda	oss-breeds) (Model 'BOV'); <i>cus</i> ) (Model 'OVI'); Model 'EQU'); imals (including <i>Bison</i> and <i>Bubalus</i> species and of the families Rhinocerotidae and Elephantidae nals (including <i>Bison</i> and <i>Bubalus</i> species and of the families Rhinocerotidae and Elephantidae
This ce — fres (1) (2) (3) — fres (4) — fres (5) (6) (7)	sh meat, including minced meat, o domestic bovine animals (inclu domestic ovine animals ( <i>Ovis a</i> domestic porcine animals ( <i>Sus</i> sh meat, excluding minced meat, o domestic solipeds ( <i>Equus cab</i> sh meat, excluding offal and mince farmed non-domestic animals their cross-breeds), <i>Ovis aries</i> , (Model 'RUF'); wild non-domestic animals of their cross-breeds), <i>Ovis aries</i> , (Model 'RUW'); farmed non-domestic animals wild non-domestic animals bel	f: uding <i>Bubalus</i> and <i>Bison</i> species and their cro aries) or domestic caprine animals ( <i>Capra hird</i> <i>s scrofa</i> ) (Model 'POR'); of: <i>allus</i> , <i>Equus asinus</i> and their cross-breeds) (I ed meat, of: of the order Artiodactyla (excluding bovine ani <i>Capra hircus</i> , Suidae and Tayassuidae), and of the order Artiodactyla (excluding bovine anin <i>Capra hircus</i> , Suidae and Tayassuidae), and of belonging to the Suidae, Tayassuidae, or Tapi	oss-breeds) (Model 'BOV'); <i>cus</i> ) (Model 'OVI'); Model 'EQU'); imals (including <i>Bison</i> and <i>Bubalus</i> species and of the families Rhinocerotidae and Elephantidae nals (including <i>Bison</i> and <i>Bubalus</i> species and of the families Rhinocerotidae and Elephantidae iridae families (Model 'SUF'); ae families (Model 'SUW');

COUNTRY Model TRANSIT/STORAGE						
II. Hea	alth information	II.a. Certificate reference num	per	II.b.		
Part I:						
<ul> <li>Box refere or ship cha</li> <li>Box refere provided. I</li> <li>Box refere</li> <li>Box refere</li> <li>Box refere</li> <li>Box refere</li> <li>Box refere</li> <li>Box refere</li> <li>Part II:</li> <li>(1) Keep as a (2) Date or da date of aut</li> </ul>	nce I.11: Place of origin: nam- nce I.12: Address (and appro- andler shall be included. In case of unloading and reloa ince I.19: Use the appropriate ince I.20: Indicate total gross in ince I.23: For containers or bo ince I.28: <i>Nature of commodit</i> , ince I.28: <i>Treatment type</i> : If from ppropriate. ttes of slaughter. Imports of the thorisation for exportation to the phere restrictive measures have	r (railway wagons or container a ading, the consignor must inform HS code: 02.01, 02.02, 02.03, 0 weight and total net weight. ixes, the container number and th y: Indicate 'carcass-whole', 'carca ozen, indicate the date of freezing is meat shall not be authorised w ne Union of the third country, territ	tablishment. nouse in a free nd lorries), flig the BIP of ent 2.04, 02.05, 0 ne seal numbe ass-side', 'card g (mm/yy) of th rhen obtained ory or part the	zone, free warehouse, customs warehouse pht number (aircraft) or name (ship) is to be ry into the Union. 2.06, 02.08.90, 02.09, 05.04 or 15.02. er (if applicable) should be included. cass-quarters', 'cuts', or 'minced meat'.		
Official veterin						
Nan Date	ne (in capital letters):		Qualification	and title:		
Date			Signature:			

### ANNEX IV

### ANIMALS REFERRED TO IN ARTICLE 1(1)(b)

### PART 1

### Lists of third countries, territories or parts thereof

### SECTION 1

### Parts of third countries or territories referred to in Article 7(2)

▼<u>M1</u>

Country/territory	Code of part of the country/territory	Description of part of the country/ territory
US – United States	US-A	The State of Hawaii (1)
( <sup>1</sup> ) Suspended from 5 May 2010.		

### ▼<u>C1</u>

### PART 2

### Tables of animals and the corresponding model veterinary certificates

Table 1					
'QUE': Model of veterinary certificate for consignments of queen bees and queen bumble bees (Apis mellifera and Bombus spp.),					
'BEE': Model of veterinary certificate for consignments of colonies of bumble bees (Bombus spp.)					
Order	Family	Genera/species			
Hymenoptera	Apidae	Apis mellifera, Bombus spp.			

	COUNTRY	Model QUE Veterinary certificate to EU				
	I.1. Consignor	I.2. Certificate reference number I.2.a.				
	Name					
	Address	I.3. Central Competent Authority				
Part I: Details of dispatched consignment	Tel. No	I.4. Local Competent Authority				
		1.6.				
	I.5. Consignee	1.0.				
	Name					
	Address					
	Postal code					
	Tel. No					
of dispa	I.7. Country ISO I.8. Region C of origin code of origin	Code I.9. Country of ISO I.10. Region of Code destination code destination				
ails	I.11. Place of origin	1.12.				
t I: Deta	Name Approval number Address					
Part	Name Approval number Address					
	Name Approval number Address					
	I.13. Place of loading	I.14. Date of departure time of departure				
	Address Approval number					
	I.15. Means of transport Aeroplane Ship Railway wagon	I.16. Entry BIP in EU				
	Road vehicle	I.17. No(s) of CITES				
	Identification: Documentary references:					
	I.18. Description of commodity	I.19. Commodity code (HS code) 01.06.90				
		I.20. Quantity				
	l.21.	I.22. Number of packages				
	I.23. Identification of container/seal number	1.24.				
	I.25. Commodities certified for: Breeding					
-	1.26.	I.27. For import or admission into EU				
	I.28. Identification of the commodities	1				
		Identification Identification system number				

	COUNTRY Model QUI										
	II.	Health	information	II.a. Certificate reference number	II.b.						
	II.1.	Animal Health attestation:									
u		I, the u	, the undersigned, hereby certify, that the animals referred to in Part I of this certificate meet the following requirements:								
		II.1.1	they come from the territory with code:(1) in which, American foulbrood, the small hive beetle (Activity turnida) and the Tropilaelaps mite ( <i>Tropilaelaps</i> spp.) are notifiable diseases/pests.								
icati		II.1.2	they:								
Certif			(a) come from a breedin	g apiary, which is supervised and controlled	by the competent authority;						
Part II: Certification			(b) come from an area which is not subject to any restrictions associated with an occurrence of American foulbrood, and where no such occurrence has taken place within at least 30 days prior to the issuance of the present certificate. Where an outbreak of American foulbrood has occurred previously, all hives within a radius of three kilometres have been checked by the competent authority and all infected hives burned or treated and inspected to the satisfaction of the said competent authority within 30 days following the last recorded case:								
			have been tested in t		umble bees) from which samples of the comb id down in the OIE Manual of Diagnostic Tests						
				at least 100 km radius which is not subject to a the least 100 km radius which is not subject to a the least 100 km radius of <i>Tropilaelaps</i> spp, and	any restrictions associated with the occurrence I where these infestations are absent;						
			()	ne from hives or colonies (in the case of bun show no clinical signs or suspicion of diseas	nble bees), which were inspected immediately se including infestations affecting bees;						
					and packaging do not contain the small hive tions, in particular <i>Tropilaelaps</i> spp., affecting						
	ood are new and have not been in contact with to prevent contamination with agents causing										
	Notes										
	Part I:										
	<ul> <li>Box reference I.20: Number of queen bees (Apis mellifera and Bombus spp.). Each queen bee may be accompanied by a maximur of 20 attendants.</li> </ul>										
	Part II:										
	(1) Code of the territory as it appears in Part 1 of Annex II or Section 1 of Part 1 of Annex IV to Regulation (EU) No 206/2010.										
	Official veterinarian /Official inspector										
		Name	(in capital letters):	Qualificati	on and title:						
		Date:		Signature:							
		Stamp	:								

				Mout	el BEE				
	со	UNTRY						Veterinary ce	ertificate to El
	1.1.	1. Consignor			I.2. Certific	ate reference	e numbe	r I.2.a.	
		Name	I.3. Central Competent Authority						
		Address							
nment		Tel. No	I.4. Local C	ompetent Al	utnority				
	l.5.	5. Consignee			I.6.				
		Name							
nsig		Address							
Part I: Details of dispatched consignment		Postal code							
		Tel. No							
f dispat	I.7.	Country ISO of origin code	I.8. Region of origin	Code	I.9. Country destina		ISO code	I.10. Region of destination	Code
ils o	I.11	. Place of origin	· · · ·		I.12.	·			
l: Deta		Name Address	Approval number						
Part		Name Approval number Address							
		Name Approval number Address							
I.13. Place of loading Address App			Approval number	I.14. Date of departure		time of departure			
	l.15	15. Means of transport			I.16. Entry BIP in EU				
		Aeroplane 🗌 Sh	ip 🗌 Railway wago	on 🗌					
		Road vehicle Oth	er						
		Identification: Documentary references:				I.17. No(s) of CITES			
	I.18	I.18. Description of commodity				I.19. Com	modity c	ode (HS code)	01.06.90
							I.20.	Quantity	
	I.21						1.22.	Number of packa	ges
	I.23. Identification of container/seal number			1.24.					
	1.25	i. Commodities certified for: Breeding							
	1.26.			I.27. For imp	ort or admis	sion into	EU		
	1.28	B. Identification of the commo	dities						
		Species (Scientific name)		Identif sys				Identificati number	on

	COUNTR	łY		Model B						
	П.	Health information	II.a. Certificate reference number	II.b.						
	II.1. Animal Health attestation:									
		I, the undersigned, hereby certify that:								
		II.1.1								
		(a) the bumble bees (Bombus spp.) referred to in Part I of this certificate have been bred and kept under a cont environment within a recognised establishment which is supervised and controlled by the competent auth								
		(b) the establishment referred to in Part I of this certificate was inspected immediately prior to dispatch and bumble bees and breeding stock show no clinical signs or suspicion of disease including infestations affect bees;								
		(c) all colonies for import into the Union have undergone detailed examination to ensure that all bumble b broodstock and packaging do not contain the small hive beetle (Aethina tumida) or its eggs and larvae or c infestations in particular Tropilaelaps spp., affecting bees;								
			combs, and all precautions have been taken	are new and have not been in contact with to prevent contamination with agents causing						
	Notes									
	Part I:									
	<ul> <li>Box reference I.20: Number of containers of bumble bees (<i>Bombus</i> spp.), each containing a colony of a maximum of 200 ac bumble bees.</li> </ul>									
	Official veterinarian /Official inspector									
Name (in capital letters): Qualification and title:				on and title:						
		Date:	Signature:							
		Stamp:								

### ANNEX V

#### Explanatory notes for completing the veterinary certificates

(referred to in Article 18)

(a) Veterinary certificates shall be issued by the exporting third country, based on the models set out in Part 2 of Annexes I, II and IV and Annex III according to the layout of the model that corresponds to the live animals/fresh meat 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.

If the Member State of destination imposes, for the live animals/fresh meat concerned, additional certification requirements, attestations to certify that those requirements are fulfilled shall also be incorporated in the original form of the veterinary certificate.

- (b) Where the model certificate states that certain statements shall be kept as appropriate, statements which are not relevant, may be crossed out and initialled and stamped by the certifying officer, or completely deleted from the certificate.
- (c) A separate and unique certificate must be provided for the live animals/fresh meat that are exported from a territory or territories of the same exporting country appearing in columns 2 and 3 of Part 1 of Annex I, II or IV which are consigned to the same destination and transported in the same railway wagon, lorry, aircraft or ship.
- (d) The original of each certificate shall consist of a single sheet of paper, or, where more text is required it must be in such a form that all sheets of paper required are part of an integrated whole and indivisible.
- (e) The veterinary certificate shall be drawn up in at least one of the official languages of the Member State of the border inspection post of introduction of the consignment into the Union and of the Member State of destination. However, those Member States may authorise the certificate to be drawn up in the official language of another Member State, and accompanied, if necessary, by an official translation.
- (f) If for reasons of identification of the items of the consignment (schedule in point I.28 of the model veterinary certificate), additional sheets of paper are attached to the certificate, those sheets of paper shall also be considered as forming part of the original of the certificate by the application of the signature and stamp of the certifying officer, on each of the pages.
- (g) When the certificate, including additional schedules referred to in (f), comprises more than one page, each page shall be numbered, (page number) of (total number of pages), at the end of the page and shall bear the certificate reference number that has been designated by the competent authority at the top of the pages.
- (h) The original of the certificate must be completed and signed by an official veterinarian or by another designated official inspector where this is provided for in the model veterinary certificate. In the case of live animals, the certificate must be completed and signed within 24 hours prior to loading of the consignment for introduction into the Union. The competent authorities of the exporting third country shall ensure that rules of certification equivalent to those laid down in Directive 96/93/EC (<sup>1</sup>) are followed.

The colour of the signature shall be different from that of the printing. This requirement also applies to stamps other than those embossed or water-marked.

(i) The certificate reference number referred to in boxes I.2 and II.a. must be issued by the competent authority.

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(1) OJ L 13, 16.1.1997, p. 28.