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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>M1</u>	Commission Regulation (EU) No 810/2010 of 15 September 2010	L 243	16	16.9.2010
► <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
► <u>M7</u>	Commission Implementing Regulation (EU) No 546/2012 of 25 June 2012	L 165	25	26.6.2012
► <u>M8</u>	Commission Implementing Regulation (EU) No 644/2012 of 16 July 2012	L 187	18	17.7.2012
► <u>M9</u>	Commission Implementing Regulation (EU) No 1036/2012 of 7 November 2012	L 308	13	8.11.2012
► <u>M10</u>	Commission Implementing Regulation (EU) No 1160/2012 of 7 December 2012	L 336	9	8.12.2012
► <u>M11</u>	Commission Implementing Regulation (EU) No 71/2013 of 25 January 2013	L 26	7	26.1.2013
► <u>M12</u>	Commission Implementing Regulation (EU) No 102/2013 of 4 February 2013	L 34	4	5.2.2013
► <u>M13</u>	Commission Implementing Regulation (EU) No 191/2013 of 5 March 2013	L 62	22	6.3.2013
► <u>M14</u>	Commission Implementing Regulation (EU) No 196/2013 of 7 March 2013	L 65	13	8.3.2013

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- ▶<u>C1</u> Corrigendum, OJ L 146, 11.6.2010, p. 1 (206/2010)
- ► C2 Corrigendum, OJ L 49, 24.2.2011, p. 53 (144/2011)
- ► C3 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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⁽¹⁾ OJ L 268, 14.9.1992, p. 54.

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

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

⁽⁵⁾ 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 (¹), 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 (²), 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 (³) 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 (⁴) 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.

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

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

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

^{(&}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 (¹), 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 (²) 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

⁽¹⁾ OJ L 157, 30.4.2004, p. 33.

⁽²⁾ 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 (¹), 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 (²).
- (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.

⁽¹⁾ OJ L 125, 23.5.1996, p. 10.

⁽²⁾ 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 (¹) and Council Regulation (EC) No 1/2005 of 22 December 2004 on the protection of animals during transport and related operations (²) 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 (³) 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.

⁽¹⁾ OJ L 340, 31.12.1993, p. 21.

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

⁽³⁾ 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 (¹);
- (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.

^{(&}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.

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Article 12a

Derogation for the transit of certain consignments of live bovine animals for breeding and production through Lithuania

1. The transit by road through Lithuania of consignments of live bovine animals for breeding and production coming from the Russian region of Kaliningrad and consigned to a destination outside the Union shall be authorised subject to compliance with the following conditions:

- (a) the animals enter Lithuania at the border inspection post at Kybartai road and exit Lithuania at the border inspection post of Medininkai;
- (b) the animals are transported in containers on road vehicles sealed with a serially numbered seal at the border inspection post of introduction into the Union at Kybartai road, by the veterinary services of the competent authority of Lithuania;
- (c) the documents referred to in the third indent of Article 7 (1) of Council Directive 91/496/EEC, including the duly completed veterinary certificate in accordance with the model veterinary certificate 'BOV-X-TRANSIT-RU' set out in Part 2 of Annex I to this Regulation, accompanying the animals from the border inspection post Kybartai road to the border inspection post Medininkai are stamped 'ONLY FOR TRANSIT FROM THE RUSSIAN REGION OF KALININGRAD VIA LITHUANIA' on each page by the official veterinarian of the competent authority responsible for the border inspection post at Kybartai road;

- (d) the requirements provided for in Article 9 of Council Directive 91/496/EEC are complied with;
- (e) the consignment is certified as acceptable for transit through Lithuania on the common veterinary entry document referred to in Article 1(1) of Commission Regulation (EC) No 282/2004 (¹) and signed by the official veterinarian of the border inspection post at Kybartai road;
- (f) the animals are accompanied by a health certificate that allows unhindered entry into Belarus and a veterinary certificate issued for the place of destination of the animals in Russia.

2. The consignment shall not be unloaded in the Union and shall be moved directly to the border inspection post of exit of Medininkai.

The official veterinarian at the border inspection post of Medininkai shall complete part 3 of the Common Veterinary Entry Document after the exit controls on the consignment have verified that it is the same consignment that entered Lithuania at the border inspection post at 'Kybartai road'.

3. In case of any irregularity or emergency during the transit, the Member State of transit shall apply the measures provided for in the second indent of Article 8(1) (b) of Directive 90/425/EEC (²) as appropriate.

4. The competent authority of Lithuania shall verify regularly that the number of consignments entering and leaving the Union territory matches.

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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.

^{(&}lt;sup>1</sup>) OJ L 49, 19.2.2004, p. 11.

⁽²⁾ OJ L 224, 18.8.1990, p. 29.

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 (¹), 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.

^{(&}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 (¹), 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 (²), 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.

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

⁽²⁾ 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

▼M1

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.

▼<u>C1</u>

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

▼<u>M8</u>

PART 1

List of third countries, territories or parts thereof $\left(*\right)$

	1	1	1		-
ISO code and name of	Code of	Description of third country, territory or part	Veterinary certificat	Specific condi-	
third country	Territory	thereof	Model(s)	SG	tions
1	2	3	4	5	6
	CA-0	Whole country	POR-X		
CA – Canada	CA-1	 Whole country, except the Okanagan Valley region of British Columbia described as follows: From a point on the Canada/United States border 120°15' longitude, 49° latitude Northerly to a point 119°35' longitude, 50°30' latitude 	BOV-X, OVI-X, OVI-Y RUM (**)		IVb IX V
		 North-easterly to a point 119° longitude, 50°45′ latitude 			
		 Southerly to a point on the Canada/ United States border 118°15' longitude, 49° latitude 			
CH - Switzerland	CH-0	Whole country	(***)		
CL CL:1	CL 0		BOV-X,OVI-X, RUM		
CL – Chile	CL-0	Whole country	POR-X, SUI	В	
GL – Greenland	GL-0	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
NZ – New Zealand	NZ-0	Whole country	BOV-X, BOV-Y, RUM, POR-X, POR-Y OVI-X, OVI-Y		III 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
RS – Serbia (*****)	RS-0	Whole country			I

	ISO code and name of	Code of	Description of third country, territory or part	Veterinary certificat	e	Specific
	third country	Territory	thereof	Model(s)	SG	tions
	1	2	3	4	5	6
		RU-0	Whole country			
	RU – Russia	RU-1	Whole country except the region of Kaliningrad			condi- tions
		RU-2	Region of Kaliningrad	BOV-X-TRANSIT-RU		X
▼ <u>M12</u>						
	US - United States	US-0	Whole country	POR-X	D	

▼M8

- (*) Without prejudice to specific certification requirements provided for by any relevant agreement between the Union and third countries.
- (**) 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 under UNSCR 1244/99.

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 (¹) for live bovine animals for slaughter and fattening and in accordance with Model I of Annex E to Directive 91/68/EEC (²) 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.

(**) Serbia, not including Kosovo under UNSCR 1244/99.

- **'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.
- **'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.

^(*) Delete country as applicable.

⁽¹⁾ OJ 121, 29.7.1964, p. 1977/64.

⁽²⁾ OJ L 46, 19.2.1991, p. 19.

- **'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.
- **'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.
- **'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.
- **'X':** Only for transit through Lithuania of bovine animals for breeding and/or production from the Kaliningrad region to other regions of Russia.

PART 2

Models of Veterinary Certificates

	Models	
	'BOV-X':	Model of veterinary certificate for domestic bovine animals (including Bubalus and Bison 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.
	'BOV-X-TRANSIT-RU':	Model of veterinary certificate for domestic bovine animals (including Bubalus and Bison species and their cross-breeds) intended for transit from the region of Kaliningrad to other regions of Russia via the territory of Lithuania.
	'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.
▼ <u>M12</u>	'POR-X':	Model of veterinary certificate for domestic porcine animals (Sus scrofa) intended for breeding and/or production after importation or intended for transit through the Union from one third country to another third country.
▼ <u>M8</u>	'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, Tayassuidae 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 guara	intees)
	'A':	guarantees regarding Bluetongue and Epizootic-haem- orrhagic-disease tests on animals certified according to the model of veterinary certificates BOV-X (point II.2.8 B), OVI-X (point II.2.6 D) and RUM (point II.2.6).
	'В':	guarantees regarding Swine-vesicular-disease and Classical-swine-fever tests on animals certified according to the model of veterinary certificates 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 veterinary certificates POR-X (point II.2.4 C) and SUI (point II.2.4 C).
▼ <u>M12</u>	'D':	guarantees regarding vesicular stomatitis test on animals certified according to the model of veterinary certificate POR-X (point II.2.1(b)).

Model BOV-X

col	INTRY	(Veterinary ce	tificate to EU
	l.1.	Consignor Name			e reference No	l.2.a.	
		Address		I.3. Central of	competent authority		
ŧ		Tel.		I.4. Local co	mpetent authority		
mer	1.5.	Consignee		I.6.			
sign		Name					
S		Address					
hed		Postal code		_			
Datc		Tel.					
of dispatched consignment	1.7.	Country of origin ISO code I.	.8. Region of origin Code	I.9. Country destinatio	of ISO code on	I.10. Region of destination	Code
Part I: Details of	l.11.	Place of origin	I	l.12.			
÷		Name	Approval number				
Par		Address		_			
	1.13.	Place of loading		I.14. Date of a	departure		
		Address	Approval number				
	l.15.	Means of transport		I.16. Entry BI	P in EU		
		Aeroplane 🗌 Ship 🗌	Railway wagon 🗖				
		Road vehicle Other		l.17.			
		Identification Documentary references		1.17.			
		· ·		*****	110 Commodity		
	1.18.	Description of commodity			I.19. Commodity c 01.02		
						20. Quantity	
	1.21.					22. Number of package	es
	1.23.	Seal/Container No			١.	24.	
	1.25	Commodities certified for:					
		Breeding		-attening			
			,				
	1.26.			I.27. For impo	ort or admission into	EU 🗌	
	1.28.	Identification of the commodities					
		Species (scientific name)	Breed Identificatio system	n l	dentification number	Age	Sex

С

οu	JNTRY						Model BOV-X
	П.	Health	n information			II.a. Certificate reference number	II.b.
	II.1.	Public	c Health Attesta	ition			
	ĺ	I, the	undersigned offic	cial v	eterinarian, hereby certify, that th	he animals described in this certificate	:
Part II: Certification		II.1.1.	brucellosis, for t	the p		official prohibition on health grounds, t and for the past six months in the ca tisfy these conditions;	
บ้ ≓		II.1.2.	have not receiv	ed:			
Pa			— any stilbene	or t	hyrostatic substances,		
					ogenic, gestagenic or β- agonist irective 96/22/EC);	substances for purposes other than t	herapeutic or zootechnic treatment
		II.1.3.	with regard to b	oovin	e spongiform encephalopathy (B	ISE):	
			(¹) (²) <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	ous cases in the country concerned, th ading of ruminants with meat-and-bon enforced or after the date of birth o ban.]	e meal and greaves derived from
			(¹) (³) or	[(a)		a permanent identification system en nd are not exposed bovine animals a Regulation (EC) No 999/2001;	
				(b)	meal and greaves derived from	date from which the ban on the feedin ruminants had been effectively enforc rn after the date of the feed ban.]	
			(¹) (⁴) or	[(a)		a permanent identification system en nd 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 th greaves derived from ruminants had be digenous case if born after the date of	een effectively enforced or after the
	II.2.	Anima	al Health attesta	ation	::		
		I, the	undersigned offic	cial v	reterinarian, hereby certify, that th	he animals described above meet the	following requirements:
		II.2.1.	they come from	ı the	territory with code:	(5) which, at the date of	of issuing this certificate:
			(¹) either	[(a)	has been free for 24 months fro	om foot-and-mouth disease]	
			(¹) or	[(a)	having had cases/outbreaks af	foot-and-mouth disease since fter that date, and authorised to exp No/, of	ort these animals by Commission
				(b)		m rinderpest, Rift valley fever, contagio morrhagic disease, and for six months	
				(c)		s, no vaccination against the diseases f domestic cloven-hoofed animals vaca	
			(¹) either	[(d)	has been free for 24 months fro	om bluetongue;]	
			(¹) (⁹) or	[(d)	test for the detection of antibody occasions on samples of blood	om bluetongue, and the animals have y for bluetongue and epizootic haemoi taken at the beginning of the isolation (dd/mm/yyyy) and on ithin 10 days before export;]	rrhagic 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.		
		(¹) or	inactivated vaccine, at least 60 serotype/s (inse demonstrated through a surve	ths from bluetongue, and the anima to days before the date of dispatch to <i>rt serotype/s</i>) which are those pres illance programme (¹²) in an area w under box reference I.11, and the an e specifications of the vaccine;]	the Union, against all bluetongue sent in the source population as vith a 150 km radius around the		
	II.2.2.		ined in the territory described under p without contact with imported cloven-	oint II.2.1 since birth, or for at least the hoofed animals for the last 30 days;	e last six months before dispatch to		
	II.2.3.	they have rem reference I.11.:		ays before dispatch in the holding(s	s) of origin described under box		
			nd which, in an area with a 150 km ra previous 60 days,	dius, there has been no case/outbreak	of epizootic haemorrhagic disease		
		rinderpest,		n radius, there has been no case/ou us bovine pleuropneumonia, lumpy sk			
	II.2.4 .	II.2.4. they are not animals to be killed under a national programme for the eradication of diseases, nor have they been vaccina against the diseases referred to under point II.2.1,(a) and (b);					
	II.2.5.		n herds that are not restricted unde enzootic bovine leukosis;	er the national legislation pertaining	to the eradication of tuberculosis,		
	II.2.6 .	they come from	herds recognised as officially tubero	culosis-free (⁶);			
	and	(¹) (⁷) <i>either</i>	[come from a region which is recog	nised as officially tuberculosis-free (6);	1		
		(¹) or	[have been subjected to an intrade 30 days before dispatch to the Unic	rmal tuberculin test (⁸) carried out wi on;]	th negative results within the past		
		(¹) or	[are less than six weeks old;]				
	II.2.7 .	they have not b	peen vaccinated against brucellosis a	nd come from herds recognised as of	ficially brucellosis-free (⁶);		
	and	(¹) (⁷) <i>either</i>	[come from a region which is recog	nised as officially brucellosis-free (6);]			
		(¹) or	[have been subjected to at least one 30 days before dispatch to the Unic	test for bovine brucellosis (⁸) carried o on,]	ut on samples taken within the past		
		(¹) or	[are less than 12 months old,]				
		(¹) or	[are castrated males of any age,]				
(¹) either	[11.2.8.			or the control of enzootic bovine leukos r test of this disease during the past t			
(1) or	[I I.2.8.	they come from	herds recognised as officially enzoo	tic-bovine-leukosis-free (⁶) (^{6a}),]			
	and	(¹) (⁷) <i>either</i>	[come from a region which is recog	nised as officially enzootic-bovine-leuk	osis-free (⁶);]		
		(¹) or	[have been subjected to an individual samples taken within the past 30 dates and the second s	al test for enzootic bovine leukosis (⁸) ays before dispatch to the Union;]	carried out with negative result on		
		(¹) or	[are less than 12 months old;]				
	II.2.9.	they are/were (1) dispatched from their holding(s) of	origin, without passing through any m	arket:		
		(¹) either	[directly to the Union,]				
		(¹) or	[to the officially authorised assembly described under point II.2.1,]	y centre described under box referen	ce I.13 situated within the territory		

COUNTRY				Model BOV-X
II.	Health	information	II.a. Certificate reference number	II.b.
		and, until dispatched to the Union:	1	
		 (a) they did not come in contact with other cloven-h this certificate, 	noofed 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 t		previous 30 days there has been a
	II.2.10	any transport vehicles or containers in which they v authorised disinfectant;	vere loaded were cleaned and disinfe	ected before loading with an officially
	II.2.11	they were examined by an official veterinarian with	in 24 hours of loading and showed n	o clinical sign of disease;
	II.2.12.	they have been loaded for dispatch to the Union o under box reference I.15 above that were cleaned a so constructed that faeces, urine, litter or fodder c	and disinfected before loading with an	officially authorised disinfectant and
II.3.	Anima	I transport attestation		
	loading	undersigned official veterinarian, hereby certify, that t g in accordance with the relevant provisions of Regu re fit for the intended transport.		
(¹) (¹¹) [II.4	Specif	ïc requirements		
	II.4.1.	According to official information, no clinical or parecorded in the holding(s) of origin referred to in but		
	II.4.2.	the animals referred to in box reference I.28 .:		
		 (a) have been isolated in accommodation approve dispatch for export, 	ed by the competent authority for the	e last 30 days immediately prior to
		(b) have been subjected to a serological test for IE results, and all animals in isolation have also gi		ter entry into isolation, with negative
		(c) have not been vaccinated against IBR.]		
Notes				
This certific production.		eant for domestic bovine animals (including Bubalus	and <i>Bison</i> species and their cross-b	preeds) intended for breeding and/or
		animals must be conveyed without delay to the holdi ment outside the holding, except in the case of a di		nain for a minimum period of 30 days
Part I:				
— Box ref	erence I.	8.: Provide the code of territory as appearing in Part	t 1 of Annex I to Regulation (EU) No	206/2010.
— Box refe No 206		13.: The assembly centre, if any, must fulfil the condi	tions for its approval, as laid down in	Part 5 of Annex I to Regulation (EU)
		15.: Registration number (railway wagons or contair ding and reloading, the consignor must inform the B		t) or name (ship) is to be provided.
— Box ref	erence I.	23.: For containers or boxes, the container number a	and the seal number (if applicable) sl	hould be included.
— Box ref	erence I.	28.: Identification system: The animals must bear:		
	ndividual sponder)	number which permits tracing of their premises of a	origin. Specify the identification system	m (such as tag, tattoos, brand, chip,
An .	eartad t	hat includes the ISO code of the exporting country	. The individual number must permit	t tracing of their premises of origin.

COUNTRY			Model BOV-X					
II. Health information		II.a. Certificate reference number	II.b.					
Species: Select amongst "Bo	Species: Select amongst "Bos", "Bison" and "Bubalus" as appropriate.							
Age: Date of birth (dd/mm/yy)).							
Sex (M = male, F = female,	C = castrated).							
Breed: select purebred, cross	sbreed.							
Part II:								
(¹) Keep as appropriate.								
	rn and continuously reared in a country r region posing a negligible BSE risk a							
	of origin is categorised in accordance and is listed as such in Decision 2007		lo 999/2001 as a country or region					
	of origin has not been categorised in a region with undetermined BSE risk and							
(⁵) Code of the territory as it ap	opears in Part 1 of Annex I to Regulation	on (EU) No 206/2010.						
) Officially tuberculosis/brucellosis-free regions and herds as laid down in Annex A to Directive 64/432/EEC; and enzootic-bovine-leukosis-free regions and herds as laid down in Chapter I of Annex D to Directive 64/432/EEC.							
Directive 64/432/EEC for the	ovine-leukosis-free herds recognised as a purpose of exports to the EU of live a < I to Regulation (EU) No 206/2010, ap	nimals according to the model certifica	ate BOV-X from the territory that, in					
	olumn 6 of Part 1 of Annex I to Regulation and/or " IVa " as regards enzootic boving		e entry "II", as regards tuberculosis,					
(⁸) Tests carried out in accorda No 206/2010.	ance with the protocols that, for the dis	sease concerned, are described in Pa	rt 6 of Annex I to Regulation (EU)					
(⁹) Supplementary guarantees to entry "A ".	o be provided when required in colum	n 5 "SG" of Part 1 of Annex I to Reg	ulation (EU) No 206/2010, with the					
Tests for bluetongue and for	r epizootic haemorrhagic disease in acc	cordance with Part 6 of Annex I to Re	gulation (EU) No 206/2010.					
exportation to the Union of	these animals shall not be allowed wh the third country, territory or part there ad by the Union against imports of thes	of referred to in Boxes I.7 and I.8, o	r during a period where restrictive					
	ember State of destination or Switzerlar mmunity and the Swiss Confederation c							
(12) 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.).					
Official veterinarian								
Name (in capital letters):		Qualification and title:						
Date:		Signature:						
Stamp:								

		Model B	OV-Y
co	UNTR	Y	Veterinary certificate to EL
	1.1.	Consignor Name	I.2. Certificate reference No I.2.a.
		Address	I.3. Central competent authority
		Tel.	
			I.4. Local competent authority
dispatched consignment	1.5.	Consignee Name Address	1.6.
tched cor		Postal code Tel.	
ď	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
tails	111	Place of origin	1.12.
Part I: Details		Name Approval number Address	
	I.13.	Place of loading	I.14. Date of departure
		Address Approval number	
	I.15.	Means of transport	I.16. Entry BIP in EU
		Aeroplane 🗌 Ship 🗌 Railway wagon 🗌	
		Road vehicle D Other	
		Identification	1.17.
		Documentary references	
	I.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	
		Species Breed Identification system (scientific name)	Identification number Age Sex

11.	Health	information			II.a. Certificate reference number	II.b.				
	riodian									
11.1.		Public Health Attestation								
	I, the u	indersigned offici	ial veteri	narian, hereby certify, that the	animals described in this certificate:					
	II.1.1. come from holdings which have been free from any official prohibition on health grounds, for the last 42 days in the case 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 in conta with animals from holdings which did not satisfy these conditions;									
	II.1.2.	.1.2. have not received:								
	— any stilbene or thyrostatic substances,									
	 — cestrogenic, androgenic, gestagenic or β- agonist substances for purposes other than therapeutic or zootechnic tr defined in Directive 96/22/EC). 									
II.1.3. with regard to bovine spongiform encephalopathy (BSE):										
	nabling them to be traced back to th cribed in Chapter C, part I, point (4) (
			(b)	from which the ban on the fe	eeding of ruminants with meat-and- y enforced or after the date of birt	d, the animals were born after the da bone meal and greaves derived fro h of the last BSE indigenous case				
		(¹) (³) or	[(a)		re not exposed bovine animals as de	nabling them to be traced back to t escribed in Chapter C, Part II, point (
			(b)	and-bone meal and greaves of		the feeding of ruminants with mea fectively enforced or after the date feed ban.]				
		(¹) (⁴) or	[(a)		e not exposed bovine animals as de	nabling them to be traced back to t ascribed in Chapter C, Part II, point (
			(b)	with meat-and-bone meal and		h the ban on the feeding of ruminar I been effectively enforced or after t e of the feed ban.]				
II.2 .	Anima	l Health Attesta	tion							
	l, the u	ndersigned offici	ial veteri	narian, hereby certify, that the	animals described above meet the t	ollowing requirements:				
	II.2.1 <i>.</i>	they come fron	n the ter	ritory with code:	(⁵) which	, at the date of issuing this certificat				
		(¹) either	[(a)	has been free for 24 months f	rom foot-and-mouth disease]					
		(¹) or	[(a)	had cases/outbreaks after t						
			(b)		om rinderpest, Rift valley fever, conta emorrhagic disease, and for six mon	agious bovine pleuropneumonia, lum ths from vesicular stomatitis,				
			(c)		ns, no vaccination against the diseas of domestic cloven-hoofed animals v					

С

COUNT	ſRY					Model BOV-Y		
II.	Health	information			II.a. Certificate reference number	II.b.		
		(¹) or	inactivated vac serotype/s demonstrated th of origin descril	cine, at least 60 hrough a surveilla	nths from bluetongue, and the anima b) days before the date of dispatch to . (<i>insert serotype/s</i>) which are those p ance programme (⁹) in an area with a 1 eference I.11, and the animals are still s of the vaccine;]	the Union, against all bluetongue present in the source population as 50 km radius around the holding(s)		
	II.2.2.			nt II.2.1 since birth, or for at least the la ofed animals for the last 30 days;	ast three months before dispatch to			
	II.2.3 .	they have rem	ained since birth or at le	ast 40 days befo	ore dispatch in the holding(s) describe	d under box reference I.11:		
			ound which, in an area w previous 60 days, and	rith a 150 km rad	dius, there has been no case/outbreak	of epizootic haemorrhagic disease		
	(b) in and around which, in an area with a 10 km radius, there has been no case/outbreak of foot-and-mouth disease, rinc Rift valley fever, bluetongue, contagious bovine pleuropneumonia, lumpy skin disease and, vesicular stomatitis dur previous 40 days;							
	II.2.4.		animals to be killed unde seases referred to in poir		gramme for the eradication of disease o);	es, nor have they been vaccinated		
	II.2.5.	they come fro	m herds:					
		(a) included in	a) included in an official system for the control of enzootic bovine leukosis, and					
		(b) that are n	ot restricted under the na	ational legislation	regarding eradication of tuberculosis a	and brucellosis, and		
		(c) recognise	as officially tuberculosis	s free; (⁶)				
	II.2.6.	they have not	been vaccinated against	t brucellosis and	they:			
		(¹) either	[come from herds which	n are recognised	as officially brucellosis free;] (⁶)			
		(¹) or	[are castrated males of	any age;]				
	II.2 .7.	they are indiv immediate sla		ast two places o	on their hindquarters as to show that	they are exclusively intended for		
	II.2.8 .	they are/were	(1) dispatched from their	holding(s) of orig	gin, without passing through any mark	et:		
		(¹) either	[directly to the Union,]					
		(¹) or	[to the officially authori described under point II		centre described under box reference	e I.13 situated within the territory		
		and, until dispatched to the Union:						
		(a) they did n certificate,		ther cloven-hoofe	ed animals not complying with the healt	h requirements as described in this		
			not at any place where, eak of any of the diseas		h within a 10 km radius, during the p point II.2.1;	revious 30 days there has been a		
	II.2.9.	any transport authorised dis		ו which they wer	re loaded were cleaned and disinfecte	ed before loading with an officially		
	II.2.10 .	they were exa	mined by an official vete	rinarian within 24	4 hours of loading and showed no clin	ical sign of disease;		
	II.2.11.	under box ref	erence I.15 above that we	ere cleaned and	disinfected before loading with an offic not flow or fall out of the vehicle o	cially authorised disinfectant and so		

COUN	TRY		Model BOV-Y				
II.	Health information	II.a. Certificate reference number	II.b.				
II.3.	Animal transport attestation						
	I, the undersigned official veterinarian, hereby certify, that in accordance with the relevant provisions of Regulation (I the intended transport.						
Note	3						
This d	certificate is meant for live bovine animals (including Bubalu	is and Bison species and their cross-breed	s) intended for immediate slaughter.				
After	importation the animals must be conveyed without delay to	the slaughterhouse of destination to be s	laughtered within five working days.				
Part	:						
— Во	ox reference I.8: Provide the code of territory as appearing i	n Part 1 of Annex I to Regulation (EU) No	206/2010.				
	ix reference I.13: The assembly centre, if any, must fulfil the 206/2010.	conditions for its approval, as laid down in	Part 5 of Annex I to Regulation (EU)				
	ix reference I.15: Registration number (railway wagons or α se of unloading and reloading, the consignor must inform the		or name (ship) is to be provided. In				
— Во	ox reference I.23: For containers or boxes, the container nur	mber and the seal number (if applicable) sh	nould be included.				
— Во	ox reference I.28: Identification system: the animals must be	ar:					
	 An individual number which permits tracing of their premises of origin. Specify the identification system (such as tag, tattoos, brand, chip, transponder). 						
	An ear tag that includes the ISO code of the exporting c	country. The individual number must permit	t tracing of their premises of origin.				
Sp	pecies: Select amongst "Bos", "Bison" and "Bubalus" as app	ropriate.					
Ag	ye: Date of birth (dd/mm/yy).						
Se	ex (M = male, F = female, C = castrated).						
Part	I:						
(¹) Ke	eep as appropriate.						
	nly if the animals were born and continuously reared in a c o 999/2001 as a country or region posing a negligible BSE						
	nly if the country or region of origin is categorised in accorr ssing a controlled BSE risk and is listed as such in Decision		No 999/2001 as a country or region				
	nly if the country or region of origin has not been categorise tegorised as a country or region with undetermined BSE ris						
(⁵) C	ode of the territory as it appears in Part 1 of Annex I to Reg	gulation (EU) No 206/2010.					
(⁶) O	fficially tuberculosis/brucellosis free regions and herds as lai	id down in Annex A to Directive 64/432/EE	С.				
	nis mark shall take the form of "L" having 13 cm in the left s pplied using the technique known as "freeze-branding".	side and 7 cm in the bottom side with 1 cm	n of strength in both lines. It shall be				

OUNTRY Model BOV-Y								
II. Health information	II.a. Certificate reference number II.b.							
(8) 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.								
(⁹) Surveillance programme as laid down in Annex	(I to Commission regulation (EC) No 1266/2007 (OJ L 283, 27.10.2007, p. 37.).							
Official veterinarian								
Name (in capital letters):	Qualification and title:							
Date:	Date: Signature:							
Stamp:								

▼<u>M10</u>

Model BOV-X-TRANSIT-RU

col	INTR	(Veterinary certificate to El					
	I.1.	Consignor Name	I.2. Certificate reference No I.2.a.					
		Address	I.3. Central competent authority					
		Tel.	I.4. Local competent authority					
nent	1.5.	Consignee	I.6. Person responsible for the load in EU					
ignn		Name	Name					
suo:		Address	Address Postal code Tel.					
dispatched consignment		Postal code Tel.						
of	1.7.	Country of ISO code I.8. Region of Code origin Russia Kaliningrad	I.9. Country of ISO code I.10. Region of Code destination Russia					
tails	l.11.	Place of origin	1.12.					
rt I: Details		Name Address						
Part		Postal code						
	l.13.	Place of loading	I.14. Date of departure					
		Address						
		Approval number						
	I.15.	Means of transport	I.16. Entry BIP in EU					
		Aeroplane Ship Railway wagon Road vehicle Other	Kybartai road — Lithuania					
		Identification Documentary references						
			1.17.					
	I.18.	Description of commodity	I.19. Commodity code (HS code) 01.02					
			I.20. Quantity					
	1.21.		I.22. Number of packages					
	1.23.	Seal/Container No	1.24.					
	1.25.	Commodities certified for:						
		Breeding 🗌 Fattening 🗋						
	1.26.	For transit through EU to third country	1.27.					
	Third country Russian Federation ISO code RU							
	I.28. Identification of the commodities							
		Species Breed Identification (scientific name)	n system Identification number Age Sex					

▼<u>M10</u>

	COUNTRY				Model BOV-X-TRANSIT-RU				
	II. He	ealth inf	formation	II.a. Certificate reference No	II.b.				
		II.1.	Animal Health attestation:						
	I, the undersigned official veterinarian, hereby certify, that the animals described in Part I meet the following requirements:								
		II.1.1.	they come from the territory with code: RU-2 $(^{2})$ wh	ich, at the date of issuing this certifica	te:				
ficatior			(1) either [(a) has been free for 24 months from for	oot-and-mouth disease;]					
Part II: Certification			without having had cases/outbreaks	as been considered free from foot-and-mouth disease since					
å	 (b) has been free for 12 months from rinderpest, Rift valley fever, contagious bovine pleuropneumonia, lumpy s disease and epizootic haemorrhagic disease, and for 6 months from vesicular stomatitis; 								
	-		(c) where, during the last 12 months, no carried out and imports of domestic of	vaccination against the diseases referr loven-hoofed animals vaccinated again					
			(1) either [(d) has been free for 24 months from b	luetongue;]					
	(1) or [(d) has not been free for 24 months from bluetongue, and the animals have been vaccinated with an inac vaccine, at least 60 days before the date of the movement, against all bluetongue serotype/s								
	(¹) either	[11.1.2.	they are of European Union origin and they were on (dd/mm/yyyy) and, since that date, origin are kept;]						
	(¹) or	[1.1.2.	they have remained in the territory with code RU-2 s the European Union and without contact with import						
		II.1.3.	they have remained [since birth or at least 40 days box reference I.11.:	before the date of dispatch (5) in the l	holding(s) of origin described under				
			 (a) in and around which, in an area with a 150 km r during the previous 60 days; 	adius, there has been no case/outbreal	 of epizootic haemorrhagic disease 				
	(b) in and around which, in an area with a 10 km radius, there has been no case/outbreak of foot-and-mouth diseas rinderpest, Rift valley fever, bluetongue, contagious bovine pleuropneumonia, lumpy skin disease and vesicular stomati during the previous 40 days;								
		II.1.4 <i>.</i>	they are not animals to be killed under a national p against the diseases referred to under point II.1.1.		ses, nor have they been vaccinated				
	 (a) they did not come in contact with other cloven-hoofed animals not complying with the health requirements as described this certificate; 								
	 (b) they were not at any place where, or around which, within a 10 km radius, during the previous 30 days there has be case/outbreak of any of the diseases referred to in point II.1.1.; II.1.5. any transport vehicles or containers in which they were loaded were cleaned and disinfected before loading with an of authorised disinfectant; 								
		II.1.6.	they were examined by an official veterinarian with	n 24 hours of loading and showed no	clinical sign of disease;				
		II.1.7 <i>.</i>	they have been loaded for dispatch to Russia via the fransport described under box reference 1.15. a authorised disinfectant and so constructed that faec during transportation;	bove that were cleaned and disinfect	ed before loading with an officially				
		ll.1.8.	the consignment is intended to leave the Europea	n Union at the designated Border Ins	pection Post Medininkai, Lithuania.				

▼<u>M10</u>

COUN	TRY			Model BOV-X-TRANSIT-RU				
11.	Health in	formation	II.a. Certificate reference No	ll.b.				
	II.2.	Animal transport attestation						
	I, the undersigned official veterinarian, hereby certify, that the animals described in Part I have been treated before and at the time of loading in accordance with the relevant provisions of Council Regulation (EC) No 1/2005, in particular as regards watering and feeding, and they are fit for the intended transport.							
Notes	Notes:							
	This certificate is meant for transit through the European Union of domestic bovine animais (including Bubalus and Bison species and their cross- breeds) intended for breeding and/or production coming from the region of Kaliningrad and destined to other parts of Russia.							
Part I	:							
— Во	ox reference	I.8.: Provide the code of territory as appearing in Part	1 of Annex I to Commission Regulat	on (EU) No 206/2010.				
		I.13.: The assembly centre, if any, must fulfil the cond) No 206/2010.	litions for its approval, as laid down i	n Part 5 of Annex I to Commission				
		I.15.: Registration number of road vehicle is to be pro- ion Post of entry into the Union.	vided. In case an emergency, the con	signor must immediately inform the				
— Во	ox reference	I.23.: For containers or boxes, the container number a	and the seal number (if applicable) m	ust be included.				
— Во	ox reference	I.28.: Identification system: the animals must bear:						
-	- An individua transponder	al number which permits tracing of their premises of o).	rigin. Specify the identification system	ı (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.				
— Во	ox reference	I.28.: Species: select amongst "Bos", "Bison" and "Bul	palus" as appropriate.					
— Во	ox reference	I.28.: Age: date of birth (dd/mm/yy).						
— Вс	ox reference	I.28.: Sex (M = male, F = female, C = castrated).						
— Во	ox reference	I.28.: Breed: select purebred, cross-breed.						
Part I	l:							
(1) K	eep as appro	ppriate.						
(²) C	ode of the te	rritory as it appears in Part 1 of Annex I to Commissi	on Regulation (EU) No 206/2010.					
R m	³) Date of loading. Transit of these animals shall not be allowed when the animals were loaded either prior to the date of authorisation for transit to Russia via the European Union from this third country, territory or part thereof referred to in Boxes I.7., or during a period where restrictive measures have been adopted by the European Union against transit of these animals from this third country, territory or part thereof via the European Union.							
(4) S	urveillance pr	rogramme as laid down in Annex I to Commission Re	gulation (EC) No 1266/2007.					
(⁵) D	elete the text	t in square brackets if the second option for point II.1.	2. is deleted.					
Officia	al veterinariar	n/Official inspector						
N	lame (in capil	tal letters):	Qualifica	tion and title:				
D	ate:		Signatur	e:				
S	tamp:							

1.1.	Y	T	Veterinary certificate to		
	Consignor Name	I.2. Certificate reference No	l.2.a.		
	Address	I.3. Central competent authori	ty		
	Tel.	I.4. Local competent authority			
1.5.	Consignee	1.6.			
	Name Address				
	Postal code				
	Tel.				
I.5.	Country of origin ISO code I.8. Region of origin Code	I.9. Country of ISO cod destination	le I.10. Region of Code destination		
1.11.	Place of origin	I.12.			
1 -	Name Approval number				
i	Address				
	Disco Charles				
1.13.	Place of loading	I.14. Date of departure			
	Address Approval number				
I.15.	Means of transport	I.16. Entry BIP in EU			
	Aeroplane 🗌 Ship 🗌 Railway wagon 🗌				
	Road vehicle Other I Identification	l.17.			
	Documentary references				
I.18.	Description of commodity	I.19. Commodity	/ code (HS code)		
			I.20. Quantity		
I.21.			I.22. Number of packages		
1.23.	Seal/Container No		1.24.		
1.25.	Commodities certified for:				
	Breeding	Fattening			
1.26.		I.27. For import or admission into EU			
	Identification of the commodities				
1.28.					

Model OVI-X

П.		Health in	nformation			II.a. Certificate reference number	II.b.			
".										
.1	•	Public Health Attestation								
		I, the ur	Idersigned	offici	al veterinarian, hereby certify, that the	e animals described in this certificate:				
		II.1.1.	brucellos	sis, fo		any official prohibition on health grounds, for the last 42 days in the case of inthrax, for the last six months in the case of rables, and, have not been in t satisfy these conditions;				
		II.1.2.	have not	t rece	ved:					
-			— any	stilbei	ne or thyrostatic substances,					
					c, androgenic, gestagenic or β- agoni d in Directive 96/22/EC).	ist substances for purposes other than	therapeutic or zootechnic treatmer			
.2		Animal	Health at	testat	ion					
		l, the ur	Idersigned	offici	al veterinarian, hereby certify, that the	e animals described above meet the t	following requirements:			
		II.2.1.	they con	ne fro	m the territory with code:	(¹) which, at the date of issu	ing this certificate:			
			(²) either	[(a)	has been free for 24 months from fo	pot-and-mouth disease]				
			(²) or	[(a)	having had cases/outbreaks after th	and-mouth disease since nat date, and authorised to export th of	ese animals by Commission Imple			
				(b)		nderpest, Rift valley fever, peste des p umonia, and epizootic haemorrhagic				
				(c)		vaccination against the diseases ment tic cloven-hoofed animals vaccinate				
			(²) either	[(d)	has been free for 24 months from b	luetongue;]				
			(²) (⁹) or	[(d)	the detection of antibody for bluetone samples of blood taken at the be	uetongue, and the animals have reacte gue and epizootic haemorrhagic disea ginning of the isolation/quarantine p /) and on	se, carried out on two occasions o period and at least 28 days late			
			(²) or	[(d)	vaccine, at least 60 days before the serotype/s) which are those prese programme $(^{11})$ in an area with a	m bluetongue, and the animals have date of dispatch to the Union, against ent in the source population as de 150 km radius around the holding still within the immunity period of time	t all bluetongue serotype/s … (inse monstrated through a surveillanc (s) of origin described under bo			
		II.2.2.				point II.2.1 since birth, or for at least th 1-hoofed animals for the last 30 days;				
		11.2.3.	they hav	re ren	nained since birth or at least 40 day	vs in the holding(s) described under	box reference I.11 before dispatch			
					ound which, in an area with a 150 luring the previous 60 days, and	km radius, there has been no case/	outbreak of epizootic haemorrhagi			
			rind	erpes		m radius, there has been no case/o te des petits ruminants, sheep pox				

COUNTR	RY			Model OVI-X		
П.	Health i	nformation		II.a. Certificate reference number	II.b.	
	I I.2. 4.	according to	o my knowledge and to the written dec	aration made by the owner, the anima	als:	
			come from holdings, and have not been inically detected:	in contact with animals of a holding, ir	n which the following diseases have	
			tagious agalactia of sheep or goats (<i>My</i> coides large colony), within the last six		ricolum, Mycoplasma mycoides var.	
		(ii) para	is, within the last 12 months,			
		(iv) Mae				
		(²) eithe	er [within the last three years,]			
		(²) or		all the infected animals were slaugh wo tests carried out at least six month		
		(b) are inclu	uded in an official system for notificatio	n of these diseases, and		
		(c) have be	een free from clinical or other evidenc	e of tuberculosis and brucellosis duri	ng the three years prior to export;	
	II.2.5.		t animals to be killed under a national p diseases referred to in point II.2.1(a) a		ses, nor have they been vaccinated	
	11.2.6.	they origina	ite:			
		(²) (³) either	r [from the territory described under bo	x reference I.8, which has been reco	gnised as officially brucellosis-free;]	
		(²) or	[from the holding(s) described under	box reference I.11, where, in respect	of brucellosis (Brucella melitensis):	
			(a) all susceptible animals have bee	n free from clinical or any signs of th	his disease for the last 12 months,	
			(b) a representative number of the do each year to a serological test, (⁴	mestic ovine and caprine animals over)	an age of six months are submitted	
		(²) (⁵) either	 (c) all domestic ovine or caprine anin with Rev. 1 vaccine more than tw 		this disease, save those vaccinated	
				y an interval of at least six months, ca (dd/mm/yyyy) on all domestic ults, and]		
		(²) or	[(c) domestic ovine or caprine animals Rev. 1 vaccine;	s under the age of seven months are v	vaccinated against this disease with	
			(d) the last two tests (6), separated b	y an interval of at least six months, ca	arried out:	
				m/yyyy) and on	dd/mm/yyyy) on all non-vaccinated	
			— on (dd/m domestic ovine and caprine a	m/yyyy) and on	dd/mm/yyyy) on all vaccinated	
			gave negative results, and]			
			(e) there are only domestic ovine and	I caprine animals that fulfil at least the a	above conditions and requirements;]	

cou	NTRY				Model OVI-X
II.		Health in	formation	II.a. Certificate reference number	II.b.
	(2)	[11.2.7.	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
		II.2.8.	In respect of scrapie		
	(²) (⁷) [II.2.8.1.	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, the animals comply wit	h the guarantees provided for in the
► ⁽¹) (²) eithe	r [II.2.8.2.	are animals intended for production born in and conti agnosed;] ◀	inuously reared on holdings in which a	case of scrapie has never been di-
	(²) (⁸) or	[II.2.8.2.	they shall have been kept continuously since birth or following requirements for at least three years:	for the last three years on a holding o	or holdings which have satisfied the
			- they are subject to regular official veterinary che	cks,	
			- the animals are identified in conformity with Unic	on legislation,	
			- no case of scrapie has been confirmed;		
			 — 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; 	slaughtered for human consumption)	have been examined for scrapie in
			 domestic ovine and caprine animals, with the exc have been introduced into the holding only if t 		
	(²) or	[11.2.8.2.	they are domestic ovine animals of the ARR/ARR p	rion protein genotype, as defined in A	Annex I to Decision 2002/1003/EC;]
		II.2.9.	$\blacktriangleright^{(2)}$ they are/were(²) dispatched from their holding(s) of origin, without passing through a	ny market, ◀
			(²) <i>either</i> [directly to the Union,]		
			(²) or [to the officially authorised assembly of described under point II.2.1.]	entre described under box reference	e I.13 situated within the territory
			and, until dispatched to the Union:		
			 (a) they did not come in contact with other cloven-he this certificate, and 	oofed animals not complying with the l	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		previous 30 days there has been a
		II.2.10.	any transport vehicles or containers in which they w authorised disinfectant;	ere loaded were cleaned and disinfed	ted before loading with an officially
		II.2.11.	they were examined by an official veterinarian within	n 24 hours of loading and showed no	clinical sign of disease;
		II.2.12.	they have been loaded for dispatch to the Union or described under box reference I.15 above that we disinfectant and so constructed that faeces, urine, during transportation.	ere cleaned and disinfected before lo	bading with an officially authorised

OUNTR			Model OVI->
II.	Health information	II.a. Certificate reference number	II.b.
11.3.	Animal transport attestation		
	 the undersigned official veterinarian, hereby certify, that t loading in accordance with the relevant provisions of Regula are fit for the intended transport. 		
Notes			
This cer productio	tificate is meant for live domestic ovine animals (<i>Ovis aries</i> on.) and domestic caprine animals (<i>Cap</i>	ra hircus) intended for breeding or
	portation the animals must be conveyed without delay to the hol urther movement outside the holding, except in the case of a		ain for a minimum period of 30 days
Part I:			
— Box r	reference I.8: Provide the code of territory as appearing in Pa	rt 1 of Annex I to Regulation (EU) No 3	206/2010.
	reference I.13: The assembly centre, if any, must fulfil the conc 06/2010.	litions for its approval, as laid down in I	Part 5 of Annex I to Regulation (EU)
	reference I.15: Registration number (railway wagons or contair of unloading and reloading, the consignor must inform the BII		or name (ship) is to be provided. In
— Box r	reference I.19: Use the appropriate HS code: 01.04.10 or 01.0	04.20.	
— Box r	reference I.23: For containers or boxes, the container number	and the seal number (if applicable) sh	ould be included.
— Box r	reference I.28; Identification system: The animals must bear:		
	n individual number which permits tracing of their premises of ansponder) and the anatomic place used in the animal.	origin. Specify the identification system	n (such as tag, tattoos, brand, chip,
— A	n ear tag that includes the ISO code of the exporting count	ry. The individual number must permit	tracing of their premises of origin.
Spec	ies: Select amongst "Ovis aries" and "Capra hircus" as approp	priate.	
Age:	(months).		
Sex ((M = male, F = female, C = castrated).		
Part II:			
(¹) Cod	e of the territory as it appears in Part 1 of Annex I to Regula	tion (EU) No 206/2010.	
(²) Kee	p as appropriate.		
(³) Only	\prime for a territory appearing with the entry "V" in column 6 of Pa	rt 1 of Annex I to Regulation (EU) No	206/2010.
(⁴) The	representative number of animals to be tested for brucellosis	must, for each holding, consist of:	
— a	all non-castrated male animals, which have not been vaccinate	ed against brucellosis, over six months	old,
— a	all non-castrated male animals, which have been vaccinated a	gainst brucellosis, over 18 months old,	
— a	all animals brought onto the holding since the previous tests, a	and	
— 2	25% of females which are sexually mature, within a minimum	of 50 females.	
	must be completed when the destination is a Member State 2/EEC.	or part of a Member State laid down	in one of the Annexes of Decision

cou	NTRY		Model OVI-X
П.	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 mu	st 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.	
(⁹)	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, o	or during a period where restrictive
(11)	Surveillance programme as laid down in Annex I to Commission F	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:		
1			

Model OVI-Y

cou	INTR	(Veterinary certificate to EU					
	1.1.	Consignor Name	I.2. Certificate reference No I.2.a.					
		Address	I.3. Central competent authority					
		Tel.						
ient			I.4. Local competent authority					
ignm	1.5.	Consignee	1.6.					
Part I: Details of dispatched consignment		Name Address						
ede								
atch		Postal code Tel.						
disp	17	Country of ISO code I.8. Region of Code	I.9. Country of ISO code I.10. Region of Code					
s of		origin origin	destination destination					
etail								
ŏ ∺	1.11.	Place of origin	1.12.					
Part		Name Approval number						
		Address						
	113	Place of loading	I.14. Date of departure					
	1.10.	hade of loading						
		Address Approval number						
	I.15.	Means of transport	I.16. Entry BIP in EU					
		Aeroplane 🗌 Ship 🗌 Railway wagon 🗌						
		Road vehicle Other	I.17.					
		Identification						
		Documentary references						
	I.18.	Description of commodity	I.19. Commodity code (HS code)					
			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						
		Species Breed Identification (scientific name) system	Identification number Age Sex					

cou	NTRY									Μ	lodel OVI-Y
	II.	Health	n information	I			II.a. Certifica	ate reference number		II.b.	
	II.1.	Public	c Health At	testat	tion						
		I, the	undersigned	l offic	ial veterinarian,	hereby certify, that th	ne animals des	cribed in this certifica	ate:		
Part II: Certification		II.1.1.	brucellosis,	for th	ne last 30 days i		for the last six			or the last 42 days in t bies, and, have not beer	
l: Ce		Ⅱ.1.2.	have not re	ceive	id:						
Part			— any stilt	bene	or thyrostatic su	ibstances,					
					androgenic, ges rective 96/22/E0		substances for	purposes other than t	thera	apeutic or zootechnic tre	atment (as
	II.2.	Anima	al Health at	testa	tion						
		I, the	undersigned	l offic	ial veterinarian,	hereby certify, that th	ne animals des	cribed above meet th	ne fo	bllowing requirements:	
		II.2.1.	they come this certifica		the territory with	h code:				(¹) which, at the date) of issuing
			(²) either	[(a)	has been free f	or 24 months from fo	oot-and-mouth o	disease]			
			(²) or		without having	had cases/outbreaks	after that date	e, and authorised to	expo	ort these animals by C (dd/mm/yyyy)	Commission
				• •						iits ruminants, sheep po se, and for 6 months fro	
										ned in points (a) and (b t these diseases are no	
			(²) either	[(d)	has been free f	or 24 months from b	iuetongue;]				
			(²) or	,	vaccine, at leas (<i>insert serotype</i> programme (⁵) i	st 60 days before the √s) which are those in an area with a 150	date of dispate present in the km radius aro	ch to the Union, again source population as bund the holding(s) of	inst a s der f orig	een vaccinated with an all bluetongue serotype monstrated through a s gin described under bo n the specifications of th	e/s surveillance x reference
		II.2.2.						birth, or for at least th for the last 30 days;		ast three months before	dispatch to
		II.2.3.	they have	rema	ined since birth	or at least 40 days	s before dispat	tch in the holding(s)	des	scribed under box refe	rence I.11:
					nd which in an a revious 60 days		adius there has	s been no case/outbre	əak (of epizootic haemorrha	gic disease
			rinderp	est, F	Rift valley fever,		es petits rumina	ants, sheep pox and g		break of foot-and-mout t pox; contagious caprir	
		II.2.4.				d under a national pr in point II.2.1(a) and		he eradication of dise	ease	es, nor have they been	vaccinated
		II.2.5.	they are/we	əre (²)	dispatched from	m their holding(s) of	origin, without p	passing through any r	marł	≺et,	
			(²) either	[dire	ectly to the Unio	n]					

COUNTRY Model OVI-1						
II.	Health ir	nformation	II.a. Certificate reference number	II.b.		
		(²) or [to the officially authorised assembly centre of under point II.2.1,]	lescribed under box reference I.13 si	tuated within the territory described		
		and, until dispatched to the Union:				
		 (a) they did not come in contact with other cloven-ho this certificate, and 	ofed animals not complying with the l	health requirements as described in		
		(b) they were not at any place where, or around which 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 benefit or (c) of Chapter A(I) of Annex VIII to Regulation (EC the programmes referred to in those points, as laid of) No 999/2001, the animals comply	with the guarantees provided for in		
(²) either	[11.2.6.2.	were born in and continuously reared on holdings in	which a case of scrapie has never	been diagnosed;]		
(²) or	[11.2.6.2.	are domestic ovine animals of the ARR/ARR prion profrom a holding where no case of scrapie has been r		to Decision 2002/1003/EC, coming		
	II.2.7.	any transport vehicles or containers in which they we authorised disinfectant;	re loaded were cleaned and disinfed	ted before loading with an officially		
	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 described under box reference I.15 above that wer disinfectant and so constructed that faeces, urine, during transportation.	e cleaned and disinfected before lo	bading 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 Regulation or the intended transport.				
Notes						
This certific after impor		eant for live domestic ovine animals (Ovis aries) and do	omestic caprine animals (Capra hircu	<i>s</i>) intended for immediate slaughter		
After impor	tation the	e animals must be conveyed without delay to the sla	ughterhouse of destination to be sla	aughtered within five working days.		
Part I:						
- Box ref	erence I.8	3: Provide the code of territory as appearing in Part 1	of Annex I to Regulation (EU) No 2	06/2010.		
— Box ref No 206		3: The assembly centre, if any, must fulfil the conditio	ns for its approval, as laid down in P	art 5 of Annex I to Regulation (EU)		
		15: Registration number (railway wagons or container g and reloading, the consignor must inform the BIP of		or name (ship) is to be provided. In		
- Box ref	erence I.1	19: Use the appropriate HS code: 01.04.10 or 01.04.2	0.			
- Box ref	erence 1.2	23: For containers or boxes, the container number and	the seal number (if applicable) sho	ould be included.		

co	DUNTRY		Model OVI-Y						
II.	Health information	II.a. Certificate reference number	II.b.						
_	- Box reference I.28: Identification system: The animals must bear:								
	— An individual number which permits tracing of their premises of origin. Specify the identification system (such as tag, tattoos, brand, chip, transponder) and the anatomic place used in the animal.								
	- 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 appropri	iate.							
	Age: months.								
	Sex (M = male, F = female, C = castrated).								
Pa	art II:								
(1)) Code of the territory as it appears in Part 1 of Annex I to Regulation	ו (EU) No 206/2010.							
(2)) Keep as appropriate.								
(3)) Guarantees in relation to a programme of control of scrapie, as require 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 whe 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 Reg	gulation (EC) No 1266/2007 (OJ L 28	3, 27.10.2007, p. 37.).						
Of	fficial veterinarian								
	Name (in capital letters):	Qualification and title:							
	Date:	Signature:							
	Stamp:								

Model POR-X

cou	INTR	(Veterinary ce	ertificate to EU
	l.1.	Consignor Name Address		I.2. Certific	ate reference	ə No	l.2.a.	
		Tel.		I.3. Centra	l competent :	authority		
ment				I.4. Local o	competent au	Ithority		
consign	1.5.	Consignee Name		I.6.				
of dispatched consignment		Address Postal code Tel.						
Part I: Details	1.7.	Country ISO I.8. Region of origin code of origin	Code	I.9. Countr of dest		ISO code	I.10. Region of destination	Code
Part I	l.11.	Place of origin Name Approval numb Address	oer	1.12.				
	I.13.	Place of loading Address Approval numb	per	I.14. Date of departure				
	I.15.	Means of transport Aeroplane A Ship Railway wagon Road vehicle Other I Identification		I.16. Entry BIP in EU				
		Documentary references	l.17.					
	l.18.	Description of commodity			I.19. Com		ode (HS code) 01.03	
						1.2	20. Quantity	
	I.21.					1.2	I.22. Number of packages	
	1.23.	Identification of container/seal number				1.2	24.	
	1.25.	Commodities certified for: Breeding						
	1.26.			I.27. For im	port or admis	ssion into	EU 🗌	
	I.28. Identification of the commodities							
		Species Identification system (scientific name)	Identifi	ication numbe	r		Age	Sex

	COUNTRY						Model POR-X					
	П.	Health	informatio	on		II.a. Certificate reference number	II.b.					
	II.1.	Public	Health A	ttestation								
		l, the	I, the undersigned official veterinarian, hereby certify, that the animals described in this certificate:									
tion		II.1. 1 .	II.1.1. come from holdings which have been free from any official prohibition on health grounds, for the last 42 days in the case of brucellosis, for the last 30 days in the case of anthrax and for the past six months in the case of rabies and, the animals have not been in contact with animals from holdings which did not satisfy these conditions;									
Part II: Certification		II.1.2. have not received:										
ŭ.			— any st	ilbene or thyrost	atic substances,							
Part				genic, androgeni d in Directive 96		substances for purposes other than the	erapeutic or zootechnic treatment (as					
	II.2.	Anima	al Health a	attestation								
		l, the	undersigne	ed official veterin	arian, hereby certify, that t	he animals described above meet the	e following requirements:					
		II.2.1.	they com	e from the territo	bry with code:	(¹) which, a	at the date of issuing this certificate:					
		(²) eith	er [(a)			nd-mouth disease, for 12 months fr ase and vesicular exanthema, and]	om rinderpest, African swine fever,					
		(²) or	[(a)			oot-and-mouth disease] (²), for 12 mo swine fever] (²) and [swine vesicular o						
 (ii) has been considered free from [foot- disease] (²), sinceauthorised to export these animals by and] 						(dd/mm/yyyy), without having had	cases/outbreaks from that date, and					
(²) <i>either</i> [(b) for 6 months from vesicular stomatitis, and]												
		(²) (⁹) (<i>or</i> [(b)	export quarantin during the pre- vector insects v test for vesicula	ne in a holding in which no export quarantine of not les where they were subjected	rs, or since birth if younger than 21 da case of vesicular stomatitis was offic ss than 30 days prior to shipment in with negative results at a serum dilutic eferred to in Part 6 of Annex I to Regu int of the quarantine; and]	ially reported during that period and a quarantine station protected from on of 1 in 32 to a virus neutralisation					
			(c)			ation against these diseases has beer these diseases are not permitted;	a carried out and imports of domestic					
		II.2.2.				oint II.2.1 since birth, or for at least th hoofed animals for the last 30 days;	e last six months before dispatch to					
		II.2.3.	and, durir	ng this period, in		er box reference I.11 since birth, or fo ea with a 10 km radius around the ho 2.1;						
	11.	2.4. A			e killed under a national pr rred to in point II.2.1;	ogramme for the eradication of disea	ses, nor have they been vaccinated					
(²) (³) [II.2.4. B they have been subjected within the past 30 days to a test for swine vesicular disease antibodies and fever antibodies with negative results in both cases;]							bodies and a test for classical swine					
	(²) (⁴) [II	.2.4. C	they have results;]	e been subjecte	d within the past 30 days	to a buffered Brucella antigen test f	or porcine brucellosis with negative					
		II.2.5	they com	e from herds wh	nich are not restricted unde	r the national brucellosis eradication	programme;					
		II.2.6	they are/\	were (²) dispatch	ed from their holding(s) of	origin, without passing through any n	narket,					
	(²)	either	[directly to	o the Union,]								
	(2)	or	[to the of point II.2.		d assembly centre describe	ed under box reference I.13 situated	within the territory described under					

COUNTRY				Model POR-X
П.	Healt	n information	II.a. Certificate reference number	II.b.
		and, until dispatched to the Union:		
		 (a) they did not come in contact with other cloven-he this certificate, and 	pofed animals not complying with the l	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 40 days there has been a
		 (c) in the case the country has not been free for 6 m protected from vector insects; 	onths of vesicular stomatitis, they were	e transported to the place of loading
	II.2.7.	any transport vehicles or containers in which they w authorised disinfectant;	ere loaded were cleaned and disinfed	eted before loading with an officially
	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 described under box reference I.15 that were cleane and so constructed that faeces, urine, litter or fodder	ed and disinfected before loading with	an officially authorised disinfectant
II.3.	Anim	al transport attestation		
	loadir	undersigned official veterinarian, hereby certify, that t g in accordance with the relevant provisions of Regul are fit for the intended transport.		
(²) (⁶) [II.4.	Spec	ific requirements		
	II.4.1.	Aujeszky's disease is notifiable in the country referre	ed to in box reference I.7;	
	II.4.2.	according to official information, no clinical, patholog the last 12 months in the holding(s) of origin referre within 5 km;		
	II.4.3.	the animals referred to in box reference I.28:		
		 (a) prior to dispatch for exportation, have remained si have remained in this(ese) holdings(s) for the last 		
		(b) have been isolated in accommodation approved dispatch for export, without direct or indirect con		last 30 days immediately prior to
		(c) have been subjected to an ELISA test for the pre- negative results; and, all animals in isolation hav		
		(d) have not been vaccinated against Aujeszky's dise origin has not been vaccinated during the previo		vaccinated animals and the herd of
(2) (8)	[.4.4.			• • •
Notes				
This certific	ate is i	meant for live domestic porcine animals (Sus scrofa)	intended for breeding or production.	
before furth	er mov	ne animals must be conveyed without delay to the hold rement outside the holding, except in the case of ani ird country to another third country.		
Part I:				
- Box refe	rence	I.8: Provide the code of territory as appearing in Part	1 of Annex I to Regulation (EU) No 2	206/2010.

- Box reference I.13: The assembly centre, if any, must fulfil the conditions for its approval, as laid down in Part 5 of Annex I to Regulation (EU) No 206/2010.

со	UNTRY		Model POR-X
11.	Health information	II.a. Certificate reference number	II.b.
_	Box reference I.15: Registration number (railway wagons or containe case of unloading and reloading, the consignor must inform the BIP		or name (ship) is to be provided. In
-	Box reference I.23: For containers or boxes, the container number a	nd the seal number (if applicable) sl	nould be included.
-	Box reference I.28.: Identification system: the animals must bear:		
	 An individual number which permits tracing of their premises of o transponder). 	rigin. Specify the identification syste	m (such as tag, tattoos, brand, chip,
	- An ear tag that includes the ISO code of the exporting country.	. The individual number must permi	t tracing of their premises of origin.
-	Box reference I.28: Age: months.		
-	Box reference I.28.: 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	(EU) No 206/2010.	
(2)	Keep as appropriate.		
(3)	Supplementary guarantees to be provided when required in column entry ${}^{\prime}\textbf{B}{}^{\prime}.$	5 'SG' of Part 1 of Annex I to Re	gulation (EU) No 206/2010, with the
(4)	Supplementary guarantees to be provided when required in column entry 'C'.	5 'SG' of Part 1 of Annex I to Rep	gulation (EU) No 206/2010, with the
(5)	Date of loading. Imports of these animals shall not be allowed whe 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 during a period where restrictive
(6)	When required by the EU Member State of destination or Switzerland the Community and the Swiss Confederation on trade in agricultural p in column 6 'Specific conditions' of Part 1 of Annex I to Regulation (roducts (OJ L 114, 30.4.2002, p. 132	
(7)	To be carried out according to the standards laid down in Annex III to used shall be the whole virus ELISA.	Decision 2008/185/EC. In the case	of pigs aged over 4 months, the test
(8)	Further requirements requested by Finland in respect of transmissible	e gastro-enteritis.	
(9)	Supplementary guarantees to be provided when required in column entry 'D'.	5 'SG' of Part 1 of Annex I to Re	gulation (EU) No 206/2010, with the
Of	ficial veterinarian		
	Name (in capital letters):	Qualific	ation and title:
	Date:	Signatu	ire:
	Stamp:		

	co	UNTRY		mouer	POR-Y			Veterinary cer	tificate to EU
		Consignor			I.2. Certifica	ate reference i	number	I.2.a.	
		Name				<u> </u>			
		Address			I.3. Central	Competent A	uthority		
		Tel. No			I.4. Local Co	ompetent Aut	hority		
ţ	I.5.	Consignee			I.6.				
mer		Name							
sign		Address							
con		Postal code							
hed		Tel. No							
patc	I.7.	Country ISO I.8	3. Region	Code	I.9. Country	of Is	so	I.10. Region of	Code
dis		of origin code	of origin		destinat	tion co	ode	destination	
Part I: Details of dispatched consignment	I.11.	Place of origin	I		l.12.				
etai			pproval number						
t I: D		Address							
Par		Name A Address	pproval number						
		Name A	pproval number						
		Address							
	I.13.	Place of loading			I.14. Date of	departure	tir	me of departure	
			pproval number						
	I.15.	Means of transport Aeroplane Ship	Railway wago	n 🗆	I.16. Entry BI	P in EU			
				" 🗀					
		Road vehicle Other			l.17.				
		Identification: Documentary references:							
	I.18	Description of commodity				I.19. Commo	odity coo	de (HS code)	01.03
					L		I.20. Q	uantity	
	I.21						I.22. N	umber of package	es
	1.23	. Identification of container/seal r	number				I.24.		
	1.25	. Commodities certified for: Slaughter							
	1.26				I.27. For impo	ort or admissi	on into E	EU	
	I.28. Identification of the commodities								
		Species (Scientific name)	Identification system		Identification number		Age	e	Sex

Model POR-Y

	COUNTRY Mode			Model POR-Y		
	Ш.	Health	information		II.a. Certificate reference number	II.b.
ation	II.1.	Public	Health Attest	ation		
		l, the u	ndersigned off	icial veterina	arian, hereby certify, that the animals described	I in this certificate:
		II.1.1	case of bruce	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,	
rtifica		II.1.2	have not rece	eived:		
Part II: Certification			— any stilbe	ene or thyros	static substances,	
Part					enic, gestagenic or β- agonist substances for pu I in Directive 96/22/EC).	rposes other than therapeutic or zootechnic
	II.2.	Anima	I Health attes	tation		
		I, the u	ndersigned 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]	
			(²) or		has been free [for 24 months from foot-and-mout African swine fever, vesicular exanthema, [cla disease] (²), and for 6 months from vesicular sto	ssical swine fever] (2) and [swine vesicular
				[has been considered free from [foot-and-moutl swine vesicular disease] (²), since cases/outbreaks from that date, and authorise Regulation (EU) No/, of	
				and	re during the last 12 months, no vaccination ag 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	l, during this	e holding(s) described under box reference I.1 period, in the holding(s) and in an area with a 1 putbreak of the diseases referred to in point II.2	10 km radius around the holding(s) of origin,
		II.2.4			be 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 disj	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.	
l						

I.	Health	information	II.a. Certificate reference number	II.b.	
	II.2.6	any transport vehicles or officially authorised disin	containers in which they were loaded were cle fectant;	aned and disinfected before loading with a	
	II.2.7	they were examined by a	n official veterinarian within 24 hours of loading	g and showed no clinical sign of disease;	
	II.2.8 they have been loaded for dispatch to the Union on				
.3.	Anima	I 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.				
²) (⁴) [II	.4. Specif	ic requirements			
	II.4.1	Aujeszky's disease is not	ifiable in the country referred to in box reference	ce I.7;	
	II.4.2		mation, no clinical, pathological or serologica) of origin referred to in box reference I.11, for i		
	II.4.3	the animals referred to in	box reference I.28:		
	(a) have remained in the holding(s) of origin referred to in box reference I.11 since birth or for the last 60 days p to dispatch for exportation, and				
		(b) have not been vaccir	nated against Aujeszky's disease.]		
lotes					
his ce	rtificate is	meant for live domestic po	prcine animals (Sus scrofa) intended for immed	liate slaughter after importation.	
fter im ays.	portation	the animals must be conve	yed without delay to the slaughterhouse of des	tination to be slaughtered within five workin	
Part I:					
– Box	c reference	e I.8: Provide the code of te	erritory as appearing in Part 1 of Annex I to Reg	gulation (EU) No 206/2010.	
		e I.13: The assembly cent EU) No 206/2010.	re, if any, must fulfil the conditions for its app	proval, as laid down in Part 5 of Annex I t	
			r (railway wagons or container and lorries), flig ading, the consignor must inform the BIP of ent		
– Box	k reference	e I.23: For containers or bo	xes, the container number and the seal numbe	er (if applicable) should be included.	
– Box	k reference	e I.28: Identification system	n: The animals must bear:		
_			s tracing of their premises of origin. Specify the natomic place used in the animal.	e identification system (such as tag, tattoo	
_	An ear ta origin.	g that includes the ISO co	de of the exporting country. The individual num	nber must permit tracing of their premises of	
– Box	<pre>creference</pre>	e I.28: <i>Age</i> : months.			

COUNTRY Model POR-Y						
II. Health information	II.a. Certificate reference number	II.b.				
Part II:						
(1) Code of the territory as it appears in Pa	art 1 of Annex I to Regulation (EU) No 206/2010).				
(2) Keep as appropriate.						
for exportation to the Union of the third	Is shall not be allowed when the animals were lo d country, territory or part thereof referred to in ed by the Union against imports of these anim	boxes I.7 and I.8, or during a period where				
(4) When required by the EU Member Stat	e of destination, in accordance with Decision 2	2008/185/EC.				
Official veterinarian	Official veterinarian					
Name (in capital letters):	Qualification	n and title:				
Date:	Signature:					
Stamp:						

Model RUM

cou	NTR	1	Veterinary certificate to EL			
	l.1.	Consignor Name	I.2. Certificate reference No I.2.a.			
		Address	I.3. Central competent authority			
÷		Tel.	I.4. Local competent authority			
signmer	I.5.	Consignee Name	1.6.			
Š		Address				
atched		Postal code Tel.				
Part I: Details 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			
Deta	I.11.	Place of origin	1.12.			
Part I:		Name Approval number Address				
	I.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			
	I.21.		I.22. Number of packages			
	I.23.	Seal/Container No	1.24.			
	1.25.	Commodities certified for:				
		Breeding Fattening	Slaughter			
	I.26.		I.27. For import or admission into EU			
	1.28.	Identification of the commodities	1			
		Species Identification system Identifi (scientific name)	cation number Age Sex			

cou	NTRY					Model RUM		
	II.	Health	information		II.a. Certificate reference number	II.b.		
	II.1.	Public	Health Attest	ation				
ation		l, the u	ne undersigned official veterinarian, hereby certify, that the animals described in this certificate:					
		II.1.1.	brucellosis an	ome from a holding which has been free from any official prohibition on health grounds, for the last 42 days in the case of ucellosis and tuberculosis, for the last 30 days in the case of anthrax, for the last six months in the case of rabies, and, have ot been in contact with animals from holdings which did not satisfy these conditions;				
Part II: Certification		∥.1.2.	have not rece	vived:				
ŏ ≓			— any stilber	ne or thyrostatic substances,				
Part				c, androgenic, gestagenic or β- agonist d in Directive 96/22/EC).	t substances for purposes other than	therapeutic or zootechnic treatment		
	II.2.	Anima	Health Attes	tation				
		I, the u	ndersigned off	icial veterinarian, hereby certify, that the	e animals described above meet the	following requirements:		
		II.2.1.	they come fro	om the territory with code:	(1) which, at the d	late of issuing this certificate:		
			contagiou	free for 24 months from foot-and-mouth s bovine pleuropneumonia, lumpy skin o leuropneumonia and epizootic haemorrh	disease, peste des petits ruminants, s	heep pox and goat pox, contagious		
	(b) where during the last 12 months, no vaccination against foot-and-mouth disease, rinderpest, Rift valley fever, con bovine pleuropneumonia, lumpy skin disease, peste des petits ruminants, sheep pox and goat pox, contagious pleuropneumonia and epizootic haemorrhagic disease and during the last 24 months no vaccination against bluetong been carried out and imports of cloven-hoofed animals vaccinated against these diseases are not permitted;				and goat pox, contagious caprine vaccination against bluetongue has			
		II.2.2.	they have ren	nained				
			(²) either	[in the territory described under point I Union and without contact with clove				
(²) or [in the country of dispatch for at least 60 days since entry, if they are animals of Part 7 of Annex I to Regulation (EU) No 206/2010 and they were imported directly for each species in Part 7 of Annex I to Regulation (EU) No 206/2010 from a third than six months prior to embarkation to the Union and in any case they have been not of the same health status after being released in the exporting country Union (³)		rectly under the conditions specified third country during a period of less been separated from other animals						
		II.2.3.	they have rer reference I.11	nained since birth or at least 40 days and I.13:	before dispatch in the holding/estab	lishment (²) described under boxes		
				ound 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 ng the previous 40 days;	there has been no case/outbreak of th	e other diseases referred to in point		
		II.2.4.		animals to be killed under a national pro f the diseases referred to in point II.2.1		ses, nor have they been vaccinated		
			(²) (⁴) <i>either</i>	[come from a herd which is recognise	ed as officially tuberculosis free, and]			
			(²) (⁵) or	[have been subjected to an intraderr	mal tuberculin test within the past 3	0 days with negative results, and]		
			they have not	been vaccinated against brucellosis an	nd they:			
			(²) (⁴) <i>either</i>	[come from a herd which is recognise	ed as officially brucellosis free;]			
			(²) (⁵) or	[have been subjected to a serum agaglutination per ml, within the past 3		ucella count of less than 30 IU of		
			(²) or	[are castrated males of any age;]				

cou	NTRY				Model RUM			
11.		Health	information	II.a. Certificate reference number	II.b.			
		II.2.5.	according to my knowledge and to the written declar	ation made by the owner, the animals				
			(a) do not come from holdings/establishments (²), ar which the following diseases have been clinically		nals of a holding/establishment, in			
			 (i) contagious agalactia of sheep or goats (Myco mycoides 'large colony'), within the last six m 		icolum, Mycoplasma mycoides var.			
			(ii) paratuberculosis and caseous lymphadenitis,	within the last 12 months,				
			(iii) pulmonary adenomatosis, within the last three years, and					
			(iv) Maedi/Visna or caprine viral arthritis/encepha	litis,				
			(²) <i>either</i> [within the last three years,]					
				the infected animals were slaughtered ests carried out at least six months a				
		(b) are included in an official system for notification of these diseases, and						
			(c) have been free from clinical or other evidence of	tuberculosis and brucellosis during th	e three years prior to export;			
	(²) (⁶)	(²) (⁶) [II.2.6. the animals have reacted negatively to a serological test for the detection of antibody for bluetongue and epizootic rhagic-disease, carried out on two occasions on samples of blood taken at the beginning of the isolation/quarantine p at least 28 days later on			the isolation/quarantine period and			
		II.2.7. they are dispatched from the holding/establishment described under boxes reference I.11 and I.13 directly to the Union and dispatched to the Union:						
			 (a) they did not come in contact with other cloven-hc this certificate, and 	ofed animals not complying with the h	ealth requirements as described in			
			(b) they were not at any place where, or around whi case/outbreak of any of the diseases referred to		previous 30 days there has been a			
		II.2.8.	any transport vehicles or containers in which they we authorised disinfectant;	ere loaded were cleaned and disinfec	ted before loading with an officially			
		II.2.9.	they were examined by an official veterinarian within	24 hours of loading and showed no o	linical sign of disease;			
		ll.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 r	d disinfected before loading with an offi	cially authorised disinfectant and so			
II.3.		Anima	I transport attestation					
		I, the undersigned official veterinarian, hereby certify, that the animals described above have been treated before and at the time o 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.						
(2) (⁸) [II.4.	Specif	ic requirements					
		II.4.1.	According to official information, no clinical or patholog in the holding/establishment (²) of origin referred to in					
		II.4.2.	the animals referred to in box reference l.28.:					
			 (a) have been isolated in accommodation approved by for export, and 	y the competent authority for the last 30) days immediately prior to dispatch			
			(b) have been subjected to a serological test for IBF results, and all animals in isolation have also give		r entry into isolation, with negative			

COUNTRY			Model RUM
II. Health ir	nformation	II.a. Certificate reference number	II.b.
(c)) have not been vaccinated against IBR.;		
(²) [II.4.3	(further requirement	ts and/or tests)]]
Notes			
	eant for live animals of the order Artiodactyla (excludi <i>Capra hircus</i> , Suidae and Tayassuidae), and of the fa		
	animals must be conveyed without delay to the holdi ment outside the holding, except in the case of a di		in for a minimum period of 30 days
Part I:			
- Box reference I.8	3.: Provide the code of territory as appearing in Part	1 of Annex I to Regulation (EU) No 2	206/2010.
- Box reference I.1 No 206/2010.	3.: The assembly centre, if any, must fulfil the condit	ions for its approval, as laid down in P	art 5 of Annex I to Regulation (EU)
	5.: Registration number (railway wagons or containe g and reloading, the consignor must inform the BIP		or name (ship) is to be provided. In
- Box reference I.1	9.: Use the appropriate HS code: 01.02, 01.04.10,	01.04.20 or 01.06.19.	
- Box reference I.2	23.: For containers or boxes, the container number a	and the seal number (if applicable) sho	ould be included.
	8.: Identification system: Specify the identification sys orting country. The individual number must permit tra		nder). The ear tag includes the ISO
Age: months.			
Sex (M = male, I	F = female, C = castrated).		
Species: Select t	the species amongst those listed for the following fa	milies:	
Antilocapridae:	Antilocapra spp.;		
Bovidae:	Addax spp., Aepyceros spp., Alcelaphus spp., Am laphus spp., Budorcas spp., Capra spp. (excluding (including Beatragus), Dorcatragus spp., Gazella s Madoqua spp., Naemorhedus spp. (including Nem spp., Oryx spp., Ourebia spp., Ovibos spp., Ovis Pseudois spp., Pseudoryx spp., Raphicerus spp., F Sylvicapra spp., Syncerus spp., Taurotragus spp.,	I Capra hircus), Cephalophus spp., Co spp., Hemitragus spp., Hippotragus s orhaedus and Capricornis), Neotragus spp. (excluding Ovis aries), Pantholop Redurica 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.,
Camelidae:	Camelus spp., Lama spp., Vicugna spp.		
Cervidae:	Alces spp., Axis-Hyelaphus spp., Blastocerus spp Hippocamelus spp., Hydropotes spp., Mazama sp spp., Pudu spp., Rangifer spp.		
Giraffidae:	Giraffa spp., Okapia spp.		
Hippopotamidae:	Hexaprotodon-Choeropsis spp., Hippopotamus spp).,	
Moschidae:	Moschus spp.		
Tragulidae:	Hyemoschus spp., Tragulus-Moschiola spp.,		
Rhinocerotidae:	Ceratotherium spp., Dicerorhinus spp., Diceros spp	o., <i>Rhinoceros</i> spp.	
Elephantidae:	Elephas spp., Loxodonta spp., as appropriate.		

OUNTRY Model RUM							
II. Health information	II.a. Certificate reference number	II.b.					
Part II:							
(1) Code of the territory as it appears in Part 1 of Annex I to Regulation	on (EU) No 206/2010.						
(²) Keep as appropriate.							
(³) In this case the health certificate has to be accompanied by the offici- I to Regulation (EU) No 206/2010 (model "CAM").	al document on quarantine and test cor	nditions laid down in Part 2 of Annex					
(4) Officially tuberculosis/brucellosis free regions or herds recognised 64/432/EEC and which appear in column 6 of Part 1 of Annex I to I "VIII", as regards brucellosis.							
(5) Tests carried out in accordance with the protocols that, for the dise 206/2010. However for the tuberculin test a result of an increase in exudation, necrosis, pain and/or inflammation shall be deemed to b	skin fold thickness of 2mm or more, o						
(⁶) Supplementary guarantees to be provided when required in column "A". Tests for Bluetongue and for Epizootic-haemorrhagic-disease							
exportation to the Union of the third country, territory or part there	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.							
Official veterinarian	Official veterinarian						
Name (in capital letters):	Qualification and	title:					
Date:	Signature:						
Stamp:	Stamp:						

		WOO	el SUI				
			10 Cartifia		number	Veterinary cer	tificate to EU
	I.1. Consignor		I.2. Certifica	ate reference	number	l.2.a.	
	Name		I.3. Central Competent Authority				
	Address			ompetent Aut	hority		
	Tel. No		I.4. Local Competent Authority				
ant	I.5. Consignee		I.6.			_	
шu	Name						
nsig	Address						
l co	Postal code						
chec	Tel. No						
Part I: Details of dispatched consignment	I.7. Country ISO I.8. Reg of origin code of o	gion Code rigin	I.9. Country destinat		SO I ode	I.10. Region of destination	Code
ils o	I.11. Place of origin		l.12.				
l: Deta	Name Approva Address	l number					
Part	Name Approva Address						
	Name Approva Address						
	I.13. Place of loading Address Approva	-			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 🗌 Other 🗌						
	Identification: Documentary references:		I.17. No(s) of CITES				
	I.18. Description of commodity		I.19. Commodity code (HS code)				
					l.20. Q	uantity	
	l.21.				1.22. N	umber of package	es
	I.23. Identification of container/seal numbe			l.24.			
	I.25. Commodities certified for:						
	Breeding			Slauç	ghter		
	l.26.	I.27. For imp	ort or admissi	ion into E	U		
	I.28. Identification of the commodities	1					
		fication stem	Identification number	I	Age	e	Sex

	COUNTRY	Model SU		
	II. Healt	h information II.a. Certificate reference number II.b.		
Part II: Certification	ll.1. Publi	c Health Attestation		
	I, the	undersigned official veterinarian, hereby certify, that the animals described in this certificate:		
	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, for the last 30 days in the case of anthrax and for the past six months in the case of rabies and, the animals have not been in contact with animals from holdings which did not satisfy these conditions;		
rtifica	ll.1.2	have not received:		
II: Ce		 any stilbene or thyrostatic substances, 		
Part		 oestrogenic, androgenic, gestagenic or β - agonist substances for purposes other than therapeutic or zootechnic treatment (as defined in Directive 96/22/EC). 		
	II.2. Anim	al Health attestation		
	I, the	undersigned official veterinarian, hereby certify, that the animals described above meet the following requirements:		
	II.2.1	they come from the territory with code:		
		(a) has been free for 24 months from foot-and-mouth disease, for 12 months from rinderpest, African swine fever, classical swine fever, swine vesicular disease and vesicular exanthema, and for 6 months from vesicular stomatitis, and		
		(b) where during the last 12 months, no vaccination against these diseases has been carried out and imports of cloven-hoofed animals vaccinated against these diseases are not permitted;		
	II.2.2	they have remained in the territory described under point II.2.1 since birth, or for at least the last six months before dispatch to the Union and without contact with cloven-hoofed animals imported into this territory less than six months ago;		
	II.2.3	they have remained in the holding described under boxes reference I.11 and I.13 since birth, or for 40 days prior to dispatch, and, during this period, in the holding(s) and in an area with a 10 km radius around the holding(s) of origin, there has been no case/outbreak of the diseases referred to in point II.2.1;		
	II.2.4 A	they are not animals to be killed under a national programme for the eradication of diseases, nor they have been vaccinated against the diseases referred to in point II.2.1 and they have been subjected within the past 30 days to a buffered Brucella antigen test for porcine brucellosis with negative results;		
	(²) (³) [II.2.4 B	they have been subjected within the past 30 days to a test for swine vesicular disease antibodies and a test for classical swine fever antibodies with negative results in both cases]		
	(²) (⁴) [II.2.4 C	they have been subjected within the past 30 days to a buffered Brucella antigen test for porcine brucellosis with negative results]		
	II.2.5	they come from holdings which:		
		(a) are not restricted under a national control and eradication programme for brucellosis, porcine enteroviral encephalomyelitis (Teschen disease), and		
	(b) are included in an official system for notification of these diseases;			
	II.2.6	they are dispatched from the holding described under boxes reference I.11 and I.13 directly to the Union and, until dispatched to the Union:		
		(a) they did not come in contact with other cloven-hoofed animals not complying with the health requirements as described in this certificate, and		
		(b) they were not at any place where, or around which within a 10 km radius, during the previous 40 days there has been a case/outbreak of any of the diseases referred to in point II.2.1;		

COUNTR	COUNTRY Model SUI					
II.	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 disinfe officially authorised disinfectant;				aned and disinfected before loading with an		
II.2.8 they were examined by a			n official veterinarian within 24 hours of loading	g 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 o		arian, hereby certify, that the animals described th the relevant provisions of Regulation (EC) N he intended transport.			
(²) (⁶) [II.4	I. Specif	ic requirements				
	II.4.1	Aujeszky's disease is not	tifiable in the country referred to in box reference	ce I.7;		
II.4.2 According to official information, no clinical, pathological or serological evidence of Aujeszky's disease recorded for the last 12 months in the holding(s) of origin referred to in boxes reference I.11 and I.13, and with a 5 km radius around 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 competen export, without direct or indirect contact with ot			
			d to an ELISA test for the presence of gl antib vith negative results; and, all animals in isolation			
			nated against Aujeszky's disease and have not s not been vaccinated during the previous 12 m			
(²) (⁸) [II.4.4			(further requirements and/or tests)		
Natas						
This cert	Notes This certificate is meant for live non-domestic Suidae (Babyrousa spp., Hylochoerus spp., Phacochoerus spp., Potamochoerus spp.,					
			p., <i>Pecari</i> spp., <i>Tayassu</i> spp.) and Tapiridae (<i>Ta</i> j			
			veyed without delay to the holding of destinat outside the holding, except in the case of a dis			

СС	COUNTRY Model SUI								
II.	Health information	II.a. Certificate reference number	II.b.						
Pa	Part I:								
_	Box reference I.8: Provide the code of the	erritory as appearing in Part 1 of Annex I to	Regulation (EU) No 206/2010.						
_	Box reference I.13: The assembly cent Regulation (EU) No 206/2010.	re, if any, must fulfil the conditions for its	approval, as laid down in Part 5 of Annex I to						
_	9	r (railway wagons or container and lorries) ading, the consignor must inform the BIP of	, flight number (aircraft) or name (ship) is to be entry into the Union.						
_	Box reference I.19: Use the appropriate								
_		exes, the container number and the seal number	mber (if applicable) should be included.						
_	Box reference I.28: Identification system								
	brand, chip, transponder) and the a	natomic place used in the animal.	the identification system (such as tag, tattoos,						
	origin.	de of the exporting country. The individual	number must permit tracing of their premises of						
_	Box reference I.28: Age: months.								
_	Box reference I.28: Sex (M = male, F = 1	female, C = castrated).							
_	Box reference I.28: <i>Species</i> .								
Ра	rt II:								
(1)		rt 1 of Annex I to Regulation (EU) No 206/2	010.						
(²)	Keep as appropriate.								
(3)	Supplementary guarantees to be provise with the entry 'B'.	ded when required in column 5 'SG' of Par	t 1 of Annex I to Regulation (EU) No 206/2010,						
(4)	Supplementary guarantees to be provise with the entry 'C'.	ded when required in column 5 'SG' of Par	t 1 of Annex I to Regulation (EU) No 206/2010,						
(5)	for exportation to the Union of the third	country, territory or part thereof referred to	re loaded either prior to the date of authorisation o in boxes I.7 and I.8, or during a period where animals from this third country, territory or part						
(⁶)	When required by the EU Member State	e of destination, in accordance with Decisio	on 2008/185/EC.						
(7)	To be carried out according to the star 4 months, the test used shall be the wh		2008/185/EC. In the case of animals aged over						
(⁸)	Further requirements requested by Finla	and in respect of transmissible gastro-enter	ritis.						
Of	ficial veterinarian								
	Name (in capital letters):	Qualifica	tion and title:						
	Date: Signature:								
	Stamp:								

	со	UNTRY						Veterinary cer	tificate to EU	
	I.1.	Consignor			I.2. Certifica	ate reference nu	ımber	I.2.a.		
		Name			I.3. Central	Competent Auth	hority			
		Address								
		Tel. No			I.4. Local C	ompetent Autho	brity			
ent	I.5.	Consignee			I.6.					
hme		Name								
nsig		Address								
d co		Postal code								
tche		Tel. No								
Part I: Details of dispatched consignment	I.7.	Country ISO I. of origin code	.8. Region of origin	Code	I.9. Country destina			10. Region of destination	Code	
ils o	I.11	. Place of origin			l.12.					
I: Deta		Name Address	Approval number							
Part		Name Address	Approval number							
		Name Address	Approval number							
	I.13	. Place of loading			I.14. Date of departure time of departure					
		Address A	Approval number							
	l.15	. Means of transport Aeroplane	Railway wagor	י 🗆	I.16. Entry B	IP in EU				
		Road vehicle Other			I.17. No(s) of (CITES				
		Identification: Documentary references:								
	l.18	. Description of commodity				I.19. Commod	lity code	e (HS code)	01.06.19	
							I.20. Qu	antity		
	I.21						I.22. Nu	mber of package	es	
	1.23	B. Identification of container/seal	number				1.24.			
	1.25	6. Commodities certified for:								
		Breeding	Fi	attening			Slaugh	nter		
	I.26). 			I.27. For imp	ort or admission	n into EL	J		
	1.28	B. Identification of the commoditi	es							
		Species (Scientific name)	Identification 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

COUNT	ſRY					Model CAI					
П.	Health	information		II.a. Certificate re	ference number	II.b.					
II.1.	Quarar	ntine conditio	ns attesta	tion							
			dersigned official veterinarian, hereby certify, that the animals described in the animal health certificate (1) number 								
	(date (d Part 7 d Union a	dd/mm/yyyy) c of Annex I to Re	of entry (²)) egulation (period the	in the quarantine s EU) No 206/2010 for ey have been subjec	tation of St. Pierre and M a period of: day	Aiquelon under the conditions provided for in ys before being released for exportation to the , carried out in an approved laboratory within					
	II.1.1.	Brucellosis:									
		(a) <i>B. abortus</i> least 42 c		gglutination Test (SA	T) and Rose Bengal Test	(RBT) within two days after arrival and after at					
		(b) <i>B. ovis</i> : C	omplemen	t Fixation Test (CFT)	within two days after arri	val and after at least 42 days					
		(c) <i>B. meliter</i>	<i>isis</i> : SAT a	nd RBT within two d	ays after arrival and after	at least 42 days					
	II.1.2.	Bluetongue a	nd Epizool	ic haemorrhagic dis	ease						
		(⁵) either	[two tes 21 days	• •	e competitive Elisa test v	vithin two days after arrival and after at least					
		(⁵) or		ed free of Bluetongu		and during this period the quarantine station and no evidence of clinical disease has been					
	II.1.3.	Tuberculosis									
					o annex B to Directive 6 fter at least 42 days from	34/432/EC using bovine and avian tuberculin the first test					
	II.1.4.	Foot-and-mou after arrival ar			detection of antibodies	and a virus neutralizaton test within two days					
	II.1.5.	Rinderpest: c	ompetitive	ELISA test within tw	o days after arrival and a	fter at least 42 days					
	II.1.6.	Vesicular stor	natitis: ELI	SA or virus- neutrali	sation test within two days	s after arrival and after at least 42 days					
	II.1.7.	Rift valley feve	er: an ELIS	A test or a virus neu	tralisation test within two	days after arrival and after at least 42 days					
	II.1.8.	Lumpy skin d	umpy skin disease: ELISA or virus neutralisation test within two days after arrival and after at least 42 days								
 II.1.9. Crimean Congo haemorrhagic fever: ELISA or virus neutralisation test within two days after arrival and after 42 days II.1.10. Surra: blood microscopy within two days after arrival and after at least 42 days 											
									II.1.11.	Malignant cat	arrhal feve
II.2.	Supple	mentary gua	rantees								
	II.2.1	Bovine leukos Member State			vo days after arrival and a	fter at least 42 days (When required by the EU					

I.	Health	information		II.a. Certificate reference number	II.b.			
.3.	Treatm	nents						
	They h	ave been subj	ected to:					
	II.3.1.	an internal a	nd external a	ntiparasitic treatment during the quarantir	ne period			
	II.3.2.							
		(⁵) either	[a treatm	ent with streptomycin 25ma/kg]				
		(⁵) or	[an antib		ira spp. (specify			
	(⁵) [II.3.3.			es (if requested) on and with the test result	(dd/mm/yyyy) using vaccine]			
lote		meant for live	animals of th	ne family Camelidae.				
Part								
		- L O: Descride t						
				erritory as appearing in Part 1 of Annex I to				
		EU) No 206/20		re, ir any, must ruini the conditions for its	s approval, as laid down in Part 5 of Annex I t			
				r (railway wagons or container and lorries iding, the consignor must inform the BIP c), flight number (aircraft) or name (ship) is to b f entry into the Union.			
- E	Box reference	e I.23: For con	tainers or bo	xes, the container number and the seal nu	umber (if applicable) should be included.			
- E	Box reference	e I.28: Identific	ation system	r: The animals must bear:				
_				its tracing of their premises of origin. S and the anatomic place used in the an	pecify the identification system (such as tag imal.			
-	 An ear ta origin. 	g that include:	s the ISO coo	de of the exporting country. The individual	number must permit tracing of their premises			
- E	Box reference	e I.28: <i>Age</i> : mo	onths.					
– E	Box reference	e I.28: <i>Sex</i> (M	= male, F = f	emale, $C = castrated$).				
- E	Box reference	e I.28: Species	: Select amo	ongst <i>'Camelus</i> spp.', <i>'Lama</i> spp.', 'Vicugna	a spp.' as appropriate.			
art	11:							
		n certificate for Regulation (El			o the Union (model 'RUM') as laid down in Part			
²) C	Date in which	te in which the last animal in a group entered the quarantine facility.						
³) T	ests perform	sts performed in accordance with the methods described in Chapter 2 of Part 7 of Annex I to Regulation (EU) No 206/2010.						
¹) F	Results of the	e tests perform	ed must be a	attached in original to this health attestatio	on.			
⁵) k	(eep as appr	opriate.						
	Sampling and excessive ha				respecting the minimum time intervals to avoi			

COUNT	RY		Model CAM
П.	Health information	II.a. Certificate reference number	II.b.
Official v	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

I, the undersigned, master of ship (name have rem attached veterinary certificate No have rem from in (<i>exporting country</i>) to at any place outside (<i>exporting country</i>) en re (<i>Ports of call en route</i>). Moreover, during the jou with other animals on board of a lower health sta	mained on board the ship during the voyage in the Union and that the ship did not call bute 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 H₂ O₂ 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)

	Controls		Controls Test Sera									
	1	2	3	4	5	6	7	8	9	10	11	12
А	Cc	C-	1	2	3	4	5	6	7	8	9	10
В	Cc	C-	1	2	3	4	5	6	7	8	9	10

	Controls						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)

	Controls		ntrols Test Sera									
	1	2	3	4	5	6	7	8	9	10	11	12
Α	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
antibody control and contain BTV antigen, monoclonal
antibody and conjugate. These wells represent maximum
colour. The mean of the optical density readings from
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 μ 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 μ l sera + 40 μ 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 μ 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 μl 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 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 μ 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 μ 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 μ 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.

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).
,	Foot-and-mouth disease (FMD)
1	oot and mouth disease (1 mb)

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 CO₂ or liquid

^{(&}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 1.0×10^6 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 μ 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 μl 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 H₂SO₄.

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 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 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

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 (¹).

Classical swine fever (CSF)

Tests for classical swine fever (CSF) shall be carried out according to Decision 2002/106/EC $(^2\!).$

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.

▼<u>M12</u>

Vesicular stomatitis (VS)

The virus neutralisation (VN) test shall be carried out in accordance with the testing protocols for vesicular stomatitis set out in Chapter 2.1.19 of the OIE Manual of Diagnostic Tests and Vaccines for Terrestrial Animals.

Sera that prevent cytopathic effect (CPE) at dilutions of 1 in 32 or greater shall be considered to contain antibodies to the vesicular stomatitis virus.

▼<u>C1</u>

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.					

(¹) OJ L 167, 7.7.2000, p. 22.

^{(&}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 (¹), 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;

⁽¹⁾ 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>C1</u>

▼<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	de of Territory Description of third country, territory or part thereof	Veterinary certificate		Specific	Closing date (2)	Opening date (³)
third country	Code of Terntory		Model(s)	SG	conditions	Closing date (2)	Opening date (*)
1	2	3	4	5	6	7	8
AL – Albania	AL-0	Whole country	_				
AR – Argentina	AR-0	Whole country	EQU				
	AR-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, La Rioja, 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, Santa Fe, Tucuman, Cordoba, La Pampa, Santiago del Estero, Chaco, Formosa, Jujuy and Salta, excluding the buffer area of 25 Km from the border with Bolivia and Paraguay that extends from the Santa Catalina District in the Province of Jujuy, to the Laishi District in the Province of Formosa	BOV	А	1		18 March 2005	
		La Rioja, 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	А	1		1 December 2007
		Tucuman, Cordoba, La Pampa, Santiago del Estero, Chaco, Formosa, Jujuy and Salta, excluding the buffer area of 25 Km from the border with Bolivia and Paraguay that extends from the Santa Catalina District in the Province of	RUW	А	1		1 August 2010

VI	M2
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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	HK-0	Whole country	_				
HN – Honduras	HN-0	Whole country	BOV, EQU				
HR — Croatia	HR-0	Whole country	BOV, OVI, EQU, RUF, RUW				
			POR				8 Novemb 2012
IL – Israel	IL-0	Whole country	_				

V	12

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				
4							
JP — Japan	JP	Whole country	BOV				28 March 2013
2							
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 (⁴)	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
PY – Paraguay	PY-0	Whole country	EQU				
	PY-1	Whole country except the designated high surveillance zone of 15 km from the external borders	BOV	А	1	18 September 2011	1 August 2008
RS – Serbia (⁵)	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	_				

▼ <u>M2</u>								
	1	2	3	4	5	6	7	8
	US – United States	US-0	Whole country	BOV, OVI, POR, EQU,SUF, SUW, RUF, RUW	G			
▼ <u>M11</u>								
	UY – Uruguay	UY-0	Whole country	EQU				
				BOV	A and J	1		1 November 2001
				OVI	А	1		
▼ <u>M3</u>								
	ZA - South Africa	ZA-0	Whole country	EQU, EQW				
		ZA-1	 The whole country except: 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 the district of Camperdown, in the province of KwaZulu-Natal. 	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):

- [']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).
- ^(RUF) 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).
- ^cC[']: 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).
- ^(D): 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).
- ^cH[']: 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.
- ^cJ²: 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

ou	NTRY					Veterinary	certificate to EU
	l.1.	Consignor		I.2. Certificat	e reference No	l.2.a.	
		Name		I.3. Central competent authority			
		Address			-		
ŧ		Tel.		I.4. Local cor	mpetent authority		
nme	1.5.	Consignee		1.6.			
dispatched consignment		Name					
200		Address					
atche		Postal code					
dispa		Tel.			-		
s of	1.7.	Country of origin ISO code I.8. Region of o	rigin Code	I.9. Country	of ISO code	I.10. Region of	Code
etail				destinatio	n	destination	1
Part I: Details of	I.11.	Place of origin		I.12.			
Part		Name Approval number					
		Name Approval number Address					
	l.13.	Place of loading		I.14. Date of d	eparture		
	l.15.	Means of transport		I.16. Entry BIP in EU			
		Aeroplane 🗌 Ship 🗌 Railwa	y wagon 🔲				
		Road vehicle Other		I.17.			
		Identification Documentary references					
	l.18.	Description of commodity			I.19. Commodity	code (HS code)	
						I.20. Quantity	
	I.21.	Temperature of product				I.22. Number of pack	ages
		Ambient Chilled		Frozen 🗌			
	1.23.	Seal/Container No				I.24. Type of packagi	ng
	1.25.	Commodities certified for:					
		Human consumption 🗌					
	1.26.			I.27. For impo	rt or admission in	to EU []
	1.28.	Identification of the commodities					
		Species Nature of Treat (scientific name) commodity ty			of establishment	s Number of packages d store	Net weight
				Ū			

	COUNT	ſŖŶ	Model BOV								
	11.	Health information	II.a. Certificate reference number	II.b.							
	11.1.	Public Health Attestation									
		I, the undersigned official veterinarian declare that I am aware of the relevant requirements of Regulations (EC) No 178/2002, (EC) No 852/2004, (EC) No 853/2004, (EC) No 854/2004 and (EC) No 999/2001 and certify that the meat of domestic bovine animals described in Part I was produced in accordance with those requirements, in particular that:									
Part II: Certification	II.1.1.	the [meat] [minced meat] (1) comes from (an) establishment(s) implementing a programme based on the HACCP principles in accordance with Regulation (EC) No 852/2004;									
rt II: Cei	II.1.2.	the meat has been obtained in compliance with Section I of Annex III to Regulation (EC) No 853/2004;									
Ран		(¹) II.1.3. [the minced meat has been produced in compliance with Section V of Annex III to Regulation (EC) No 853/2004 and frozen t Internal temperature of not more than - 18 °C;]									
		II.1.4. the meat has been found fit for human consumption follo Chapter II of Section I and Chapters I and IX of Section IV									
		II.1.5. (¹) <i>either</i> [the carcass or parts of the carcass have been Annex I to Regulation (EC) No 854/2004;]	marked with a health mark in accorda	ance with Chapter III of Section I of							
		(¹) or [the packages of [meat] [minced meat] (¹) have Annex II to Regulation (EC) No 853/2004;]	been marked with an identification m	ark in accordance with Section I of							
	2005 on microbiological criteria for										
II.1.7. the guarantees covering live animals and products thereof provided by the residue plans submitted in accordance 96/23/EC, and in particular Article 29 thereof, are fulfilled;											
	II.1.8. the [meat] [minced meat] (¹) has been stored and transported in accordance with the relevant requirements of Section respectively of Annex III to Regulation (EC) No 853/2004;										
		II.1.9. with regard to bovine spongiform encephalopathy (BSE):									
		(¹) <i>either</i> [II.1.9.1. for imports from a country or a r 2007/453/EC:	region with a negligible BSE risk	and listed as such in Decision							
		 (a) the country or region is classified country or region posing a neglig 		Regulation (EC) No 999/2001 as a							
		(b) the animals from which the bovin slaughtered in a country with a n		vere born, continuously reared and							
		$(^1)$ [(c) if in the country or region there h	ave been BSE indigenous cases:								
			after the date from which the ban on l eaves derived from ruminants had be								
			ced meat does not contain and is not / to Regulation (EC) No 999/2001, bovine animals.]]]								
		(¹) or [II.1.9.2. for imports from a country or a r 2007/453/EC:	region with a controlled BSE risk	and listed as such in Decision							
		 (a) the country or region is classified country or region posing a contro 		Regulation (EC) No 999/2001 as a							

	Health info	rmation			II.a. Certificate reference number	II.b.
			(b)	stunning by means of gas inje	vine meat or minced meat was deriv cted into the cranial cavity or killed by entral nervous tissue by means of a vity;	the same method or slaughtered b
			(¹) <i>either</i> [(c)		meat does not contain and is not der lation (EC) No 999/2001, or mechani	
			(¹) <i>or</i> [(c)	quarters contain no specifie ganglia. The carcasses or	es or half carcasses cut into no mo d risk material other than the verte wholesale cuts of carcasses of b ed by a blue stripe on the labe	ebral column, including dorsal ro ovine animals containing vertebr
	(¹) or [1.1.9.3.		2001 or has been categorised	h has not been categorised in accorc as a country or region with undeterm	
					gorised in accordance with Article 5(2) egion with undetermined BSE risk;) of Regulation (EC) No 999/2001
				als from which the bovine mea derived from ruminants;	t or minced meat was derived have n	not been fed meat-and-bone meal
			means of	gas injected into the cranial	or minced meat was derived have no cavity or killed by the same methoc neans of an elongated rod-shaped in	d or slaughtered by laceration aft
	((¹) either	· [(d) the bovi	ne meat or minced meat was i	not derived from:	
			(i) spec	ified risk material as defined ir	Annex V to Regulation (EC) No 999	9/2001;
			(ii) nervi	ous and lymphatic tissues exp	osed during the deboning process;	
			(iii) mecl	nanically separated meat obtai	ned from bones of bovine animals.]	
	((¹) or	no spec wholesal	ified risk material other than	arcasses cut into no more than three v the vertebral column, including dors e animals containing vertebral colun ation (EC) No 1760/2000. (³)]]	sal root ganglia. The carcasses
	(⁴) [II.1.10.	Parli	ament and of		1688/2005 implementing Regulation ial guarantees concerning Salmonelli	
.2.	Animal He	alth atte	estation			
	I, the unde	ersigned	official veterin	arian, hereby certify, that the fi	resh meat described in Part I:	
	II.2.1.	has bee	en obtained in	the territory/ies with code:	(²) which, a	at the date of issuing this certificat
			s been free fo ice, and	^r 12 months from rinderpest, a	nd during the same period no vaccin	ation against this disease has take
	(¹) either		s been free for s taken place;]	12 months from foot-and-mout	h disease, and during the same period	d no vaccination against this diseas
	(¹) or			red free from foot-and-mouth uthorised to export this meat t	disease since (dd/mm/yyyy),	without having had cases/outbreak

COUNTRY M							
II.	Health inf	ormati	on II.a. Certificate reference number II.b.				
	(¹) (⁵) or		vaccination programmes against foot-and-mouth disease are being officially carried out and controlled in domestic bovine animals;]				
	(¹)(⁶) or		has a systematic vaccination programme against foot and mouth disease and from herds where the efficacy of this vaccination programme is controlled by the competent veterinary authority through a regular serological surveillance indicating adequate antibody levels and which also demonstrates the absence of foot and mouth virus circulation;]				
	(¹) (⁶) or		has been free for 12 months from foot-and-mouth disease, and during the same period no vaccination against this disease has taken place and is controlled by the competent veterinary authority through a regular surveillance demonstrating the absence of foot and mouth infection;]				
	I I.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 last three months before slaughter;]				
		(1)	or [have been introduced on				
		(1)	or [have been introduced on				
	II.2.3.	has	been obtained from animals coming from holdings in which:				
		(a)	None of the animals present therein have been vaccinated against [foot-and-mouth disease or] $(^7)$ rinderpest, and				
	(¹) <i>either</i>	[(b)	in these holdings, and in the holdings situated in their vicinity within 10 km, there has been no case/outbreak of foot-and- mouth disease or rinderpest during the previous 30 days,]				
	(¹) (⁸) or	[(b	there is no official restriction for animal health reasons and where, in these holdings and in the holdings situated in their vicinity within 25 km, there has been no case/outbreak of foot-and-mouth disease or rinderpest during the previous 60 days, and,				
		(c)	they have remained for at least 40 days before direct dispatch to the slaughterhouse;]				
	(¹) (¹⁴) or	[(c)	they have remained for at least 40 days before passing through one assembly centre approved by the competent veterinary authority without coming into contact with animals of a different health status prior to subsequently going directly to a slaughterhouse;]				
	(¹) (⁹) or	[(b)	there is no official restriction for animal health reasons and where, in these holdings and in the holdings situated in their vicinity within 10 km, there has been no case/outbreak of foot-and-mouth disease or rinderpest during the previous 12 months, and				
		(c)	they have remained for at least 40 days before direct dispatch to the slaughterhouse;]				
	(¹) (⁶)	[(d)	animals have not been introduced during the last 3 months from areas not approved by the EU;				
		(e)	animals are identified and registered in the national System of Identification and Certification of Origin for bovine animals;				
		(f)	the holdings in question are listed as approved holdings, following a favourable competent authorities' inspection and official report, in TRACES (¹⁰) and inspections are regularly carried out by the competent authorities to ensure that the relevant requirements provided for in Regulation (EU) No 206/2010 are respected.]				
	II.2.4. has	s beer	obtained from animals which:				
	(a)		been transported from their holdings in vehicles, cleaned and disinfected before loading, to an approved slaughterhouse ut contact with other animals which did not comply with the conditions referred to in point II.2.1, II.2.2 and II.2.3,				

I.	Heal	Ith inform	matic	n		II.a. Certificate reference number	II.b.
					slaughterhouse, have passed ante-morter no evidence of the diseases referred to	m health inspection during the 24 hours bef in point II.2.1,	iore slaughter and, in particular, ha
			(c)		een slaughtered on m/yyyy) (¹¹);	(dd/mm/yyyy) or between	(dd/mm/yyyy) and
		(1) (12)	[(d)	have r	reacted negatively to an official intra-derr	mal tuberculosis test carried out within 3 m	nonths before slaughter;]
		(1) (6)	[(e)	at the the Ur		laughter completely separate from animals	the meat of which is not intended
			refei impo	red to ortation	in point II.2.1 during the previous 30 da	ch, within a radius of 10 km, there has ber ays or, in the event of a case/outbreak of after slaughter of all animals present, remov trol of an official veterinarian;	disease, the preparation of meat
		II.2.6.					
			(¹) є	ither	[has been obtained and prepared witho certificate.]	out contact with other meats not complying	g with the conditions required in t
			(¹) (⁸) or	from carcasses in which the main acce maturation at a temperature above + 2	meat] (¹), obtained only from de-boned me essible lymphatic glands have been remo °C for at least 24 hours before the bones en tested electronically in the middle of	ved, which have been submitted were removed and in which the
						eat not conforming to the requirements re d storage until it has been packed in box	
			(¹) (9) or	from carcasses in which the main acce	meat] $^{(1)}$, obtained only from de-boned messible lymphatic glands have been remo $^\circ C$ for at least 24 hours before the bones	ved, which have been submitted
						eat not conforming to the requirements re d storage until it has been packed in box	
(1)	II.3.	Anima	l we	lfare a	attestation		
		been h	andl	ed in th	ne slaughterhouse before and at the time	e fresh meat described in Part I of this certifi of slaughter or killing in accordance with the laid down in Chapters II and III of Council f	e relevant provisions of Union legis
	Notes						
	This ce cross-b		is r	neant ·	for fresh meat, including minced meat,	of domestic bovine animals (including Bi	son and Bubalus species and th
	Fresh meat means all animal parts fit for human consumption whether fresh, chilled or frozen.						
	Part I						
	— Box	referen	ce I.	8: Prov	vide the code of territory as appearing in	Part 1 of Annex II to Regulation (EU) No	206/2010.
	— Box	referen	ce I.	11: Pla	ace of origin: name and address of the d	lispatch establishment.	
					gistration number (railway wagons or co reloading, the consignor must inform the	ntainer and lorries), flight number (aircraft) e BIP of entry into the Union.	or name (ship) is to be provided.
					e the appropriate HS code: 02.01, 02.02, of Part 1 of Annex II to Regulation (E	02.06 or 05.04. In addition, for those territo	

►⁽¹⁾ <u>M13</u>

cou	COUNTRY Model BOV							
11.	I	Health information	II.a. Certificate reference number	II.b.				
	_	Box reference I.20: Indicate total gross weight and total net weight.						
	_	ox 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", "cuts", "offal" 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.						
	_	Box reference I.28: Treatment type: If appropriate, indicate "debone	d"; "bone in"; "matured"					
	Par	rt II:						
	(1)	Keep as appropriate.						
	(²)	Code of the territory as it appears in Part 1 of Annex II to Regulati	on (EU) No 206/2010.					
	(³)	The number of bovine carcasses or wholesale cuts of carcasses, number where removal of the vertebral column is not required must 2 (1) of Regulation (EC) No 136/2004.						
	(4)	Delete if the consignment is not intended for introduction into Finla	nd or Sweden.					
	(5)	Only matured de-boned meat fulfilling the supplementary guarantee	es referred to in footnote (⁸).					
	(6)	Supplementary guarantees regarding import of matured de-boned m to Regulation (EU) No 206/2010 with the entry "H".	eat to be provided when required in c	olumn 5 "SG" of Part 1 of Annex II				
	(7)	Delete when the exporting country carries out vaccination against allowed to import into the Union matured de-boned meat which fulf						
	(⁸)	(⁸) Supplementary guarantees regarding meats from matured de-boned meat to be provided when required in column 5 "SG" of Part 1 of A II to Regulation (EU) No 206/2010, with the entry "A".						
	(⁹) Supplementary guarantees regarding meats from matured de-boned meat to be provided when required in column 5 "SG" of Part 1 of Ar II to Regulation (EU) No 206/2010, with the entry "F". The matured de-boned meat shall not be allowed for importation into the Union unt days after the date of slaughter of the animals.							
	(¹⁰) The list of approved holdings provided by the competent authority is reviewed on a regular basis and kept up to date by the comp authority. The Commission will ensure that this list of approved holdings is made publicly available for information purposes throug integrated computerised veterinary system (TRACES).							
	 (¹¹) Date or dates of slaughter. Imports of this meat shall not be allowed when obtained from animals slaughtered either prior to the d 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 where restrictive measures have been adopted by the Union against imports of this meat from this third country, territory or part the (¹²) Supplementary guarantees concerning tuberculosis test, to be provided when required in column 5 "SG" of Part 1 of Annex II to Regi (EU) No 206/2010, with the entry "E". Intra-dermal tuberculosis test to be carried out in accordance with the provisions of Annex B to Dir 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 [·]	I of Annex II to Regulation (EU)				
▶ ⁽¹⁾	(15)	OJ L 303, 18.11.2009, p. 1. ◀						
	Offi	icial veterinarian						
		Name (in capital letters):	Qualifica	tion and title:				
		Date:	Signatur	e:				
		Stamp:						

►⁽¹⁾ <u>M13</u>

Model OVI

coul	NTRY		Veterinary certificate to EL			
	1.1.	Consignor	I.2. Certificate reference No I.2.a.			
		Name				
		Address	I.3. Central competent authority			
ţ		Tel.	I.4. Local competent authority			
dispatched consignment	I.5.	Consignee	1.6.			
nsiç		Name				
о р		Address				
tche		Postal code				
lispa		Tel.				
đ	l.7.	Country of origin ISO code I.8. Region of origin Code	I.9. Country of ISO code I.10. Region of destination Code destination			
Part I: Details	l.11.	Place of origin	1.12.			
Par		Name Approval number Address				
	l.13.	Place of loading	I.14. Date of departure			
	l.15.	Means of transport	I.16. Entry BIP in EU			
		Aeroplane 🗌 Ship 🗌 Railway wagon 🗌				
		Road vehicle Other	1.17.			
		Identification Documentary references				
	l.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:	1			
		Human consumption 🔲				
	1.26.		I.27. For import or admission into EU			
	1.28.	Identification of the commodities	1			
		Species Nature of Treatment (scientific name) commodity type Abatte	Approval number of establishments Number of Net packages weight vir Cutting plant Cold store			

	COUNTRY					Model OVI		
	II. Hea	lth informatior	ı		II.a. Certificate reference number	ll.b.		
	ll.1. Publi	c Health Atte	estation					
	I, the undersigned official veterinarian, declare that I am aware of the relevant requirements of Regulations (EC) No 178/20 (EC) No 852/2004, (EC) No 853/2004, (EC) No 854/2004 and (EC) No 999/2001 and certify that the meat of domestic ovine a caprine animals described in Part I was produced in accordance with those requirements, in particular that:							
Part II: Certification	II.1.1.			at] (¹) comes from (an) establishm ation (EC) No 852/2004;	nent(s) implementing a programme b	ased on the HACCP principles in		
t II: Cei	(¹) .1.2	. the meat ha	as been obl	ained in compliance with the cond	litions set out in Section I of Annex II	I to Regulation (EC) No 853/2004;		
Par	(¹) II.1.3. [the minced meat has been produced in compliance with Section V of Annex III to Regulation (EC) No 853/2004 and frozen to a internal temperature of not more than - 18 °C;]							
	II.1.4.				wing ante and post-mortem inspectic V of Annex I to Regulation (EC) No 8			
	II.1.5.			or parts of the carcass have been egulation (EC) No 854/2004;]	marked with a health mark in accorda	ince with Chapter III of Section I of		
				s of [meat] [minced meat] (¹) have Regulation (EC) No 853/2004;]	been marked with an identification man	ark in accordance with Section I of		
	II.1.6.	the [meat] [i foodstuffs;	minced mea	at] (¹) satisfies the relevant criteria	set out in Regulation (EC) No 2073/	2005 on microbiological criteria for		
	ll.1.7.			g live animals and products thereo ular Article 29 thereof, are fulfilled;	f provided by the residue plans subr	nitted in accordance with Directive		
	II.1.8. the [meat] [minced meat] (¹) has been stored and transported in accordance with the relevant requirements of Sections I and respectively of Annex III to Regulation (EC) No 853/2004;					requirements of Sections I and V		
	II.1.9.	with regard f	to bovine sp	oongiform encephalopathy (BSE):				
	(¹) either	[ll.1.9.1. for in	nports from	a country or a region with a neglig	ible BSE risk and listed as such in D	ecision 2007/453/EC:		
		(a)		y or region is classified in accordan negligible BSE risk;	ce with Article 5(2) of Regulation (EC)	No 999/2001 as a country or region		
		(b)		Is from which the meat or minced ith negligible BSE risk; (²)	meat was derived were born, continu	iously reared and slaughtered in a		
		(¹) [(c)	if in the co	ountry or region there have been B	SE indigenous cases:			
			(¹) either	[the animals were born after the d meal and greaves derived from ru	late from which the ban on the feedin minants had been enforced.]	g of ruminants with meat-and-bone		
			(¹) or		not contain and is not derived from s 9/2001, or mechanically separated me			
	(¹) or	[II.1.9.2. for	imports fro	m a country or a region with a cor	ntrolled BSE risk and listed as such in	Decision 2007/453/EC:		
		(a)		y or region is classified in accordance controlled BSE risk;	ce with Article 5(2) of Regulation (EC)	No 999/2001 as a country or region		
		(b)	injected in	to the cranial cavity or killed by t	t was derived have not been slaughter the same method or slaughtered by d-shaped instrument introduced into th	laceration after stunning of central		

COUN	TRY		Model OVI			
П.	Health	information	II.a. Certificate reference number II.b			
		(¹) either	[(c) the meat or minced meat does not contain and is not derived from specified risk material as defined in Annex V to Regulation (EC) No 999/2001, or mechanically separated meat obtained from bones of domestic ovine or caprine animals.]			
		(¹) or	[(c) the carcasses, half carcasses or half carcasses cut into no more than three wholesale cuts, and quarters contain no specified risk material other than the vertebral column, including dorsal root ganglia.]]			
	(¹) or [II.1.9.3. for imports from a country or a region which has not been categorised in accordance with Article 5(2) of Reg (EC) No 999/2001 or has been categorised as a country or region with undetermined BSE risk and listed as a Decision 2007/453/EC:					
			 (a) the country or region has not been categorised in accordance with Article 5(2) of Regulation (EC) No 999/2001 or has been categorised as a country or region with undetermined BSE risk; 			
			 (b) the animals from which the meat or minced meat was derived have not been fed meat-and-bone meal or greaves derived from ruminants; 			
			(c) the animals from which the meat or minced meat was derived have not been slaughtered after stunning by means of gas injected into the cranial cavity or killed by the same method or slaughtered by laceration after stunning of central nervous tissue by means of an elongated rod-shaped instrument introduced into the cranial cavity;			
		(¹) either	[(d) the meat or minced meat was not derived from:			
			(i) specified risk material as defined in Annex V to Regulation (EC) No 999/2001;			
			(ii) nervous and lymphatic tissues exposed during the deboning process;			
			(iii) mechanically separated meat obtained from bones of domestic ovine or caprine animals.]			
		(¹) or	[(d) the carcasses, half carcasses or half carcasses cut into no more than three wholesale cuts, and quarters contain no specified risk material other than the vertebral column, including dorsal root ganglia.]]			
II.2.	Animal	Health atte	estation			
	l, the u	ndersigned	official veterinarian, hereby certify, that the fresh meat described in Part I:			
	II.2.1.	has been	obtained in the territory/ies with code:			
		(a) has b and	een free for 12 months from rinderpest, and during the same period no vaccination against this disease has taken place,			
	(¹) eithei		een free for 12 months from foot-and-mouth disease, and during the same period no vaccination against this disease aken place;]			
	(¹) or	break	peen considered free from foot-and-mouth disease since			
	(¹) (⁴) or	r [(b) vaccii anima	nation programmes against foot-and-mouth disease are being officially carried out and controlled in domestic bovine als;]			
	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 last three months before slaughter;]			
		(¹) or	[have been introduced on (dd/mm/yyyy) into the territory described under point II.2.1, from the territory with code (³) that at that date was authorised to import this fresh meat into the Union;];			
		(¹) or	[have been introduced on			

OUNTR	4			Model OV		
I. ⊢	lealth infor	nation	II.a. Certificate reference number	II.b.		
	II.2.3.	has been obtained from animals coming from holdings:				
		(a) in which none of the animals present therein have be	en vaccinated against [foot-and-mouth	n disease or] (⁵) rinderpest,		
		(b) not subject to prohibition as a result of an outbreak of	f ovine or caprine brucellosis during t	he previous six weeks, and		
	(¹) 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		
	(1) (4) or [(c) where there is no official restriction for health reasons and in and around which, in area of 50 km radius, there has been n case/outbreak of foot-and-mouth disease or rinderpest during the previous 90 days, and,					
		(d) where they have remained for at least 40 days before	e direct dispatch to the slaughterhous	ə;]		
	(¹)(⁸) or	[(d) where they have remained for at least 40 days bet veterinary authority without coming into contact with a slaughterhouse;]				
	II.2.4.	has been obtained from animals which:				
		(a) have been transported from their holdings in vehicles without contact with other animals which did not com				
		(b) at the slaughterhouse, have passed ante-mortem heal shown no evidence of the diseases referred to in point		re slaughter and, in particular, have		
		(c) have been slaughtered on (dd/mm/yyyy)	or between (dd/mm/yyyy) and(dd/mm/yyyy) (⁶);		
	II.2.5.	has been obtained in an establishment around which, will 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 of	in the event of a case/outbreak of d slaughter of all animals present, remove	isease, the preparation of meat for		
	II.2.6.					
	(¹) either	[has been obtained and prepared without contact with o	ther meats not complying with the co	nditions required in this certificate.]		
	(¹)(⁴) or	[contains [boneless meat] [and] [minced meat] $(^{1})$, 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 longi	ds have been removed, which have e bones were removed and in which th	been submitted to maturation at a ne 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				
	(¹)(⁷) or	[contains [boneless meat] [and] [minced meat] (¹), obta carcasses in which the main accessible lymphatic glar temperature above + 2 °C for at least 24 hours before t	ds have been removed, which have			
		has been kept strictly separate from meat not conform production, de-boning and storage until it has been pac				
▶ ⁽¹⁾ Ⅱ.3	Animal	welfare attestation				
	been ha	dersigned official veterinarian, hereby certify, that the fresh ndled in the slaughterhouse before and at the time of slaugh e met requirements at least equivalent to those laid down in	nter or killing in accordance with the rele	evant provisions of Union legislation		

►⁽¹⁾ <u>M13</u>

COUN	TR	γ		Model OVI			
П.		Health information	II.a. Certificate reference number	II.b.			
	No	tes					
	This certificate is meant for fresh meat, including minced meat, of domestic ovine animals (Ovis aries) and caprine animals (Capra hircu Fresh meat means all animal parts fit for human consumption whether fresh, chilled or frozen.						
	Pa	rt 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.				
	_	Box reference I.15: Registration number (railway wagons or container case of unloading and reloading, the consignor must inform the BIP		or name (ship) is to be provided. In			
	_	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					
	_	Box reference I.20: Indicate total gross weight and total net weight.					
	_	Box reference I.23: For containers or boxes, the container number an	nd the seal number (if applicable) sho	ould be included.			
		Box reference I.28: <i>Nature of commodity:</i> Indicate "carcass-whole", "or meat is de-boned meat that has been minced into fragments and that adjoining fatty tissues) except heart muscle.					
		Box reference I.28: <i>Treatment type</i> : If appropriate, indicate "de-bone freezing (mm/yy) of the cuts/pieces.	ed"; 'bone in"; "matured" and/or "minc	ed". If frozen, indicate the date of			
	Ра	rt II:					
	(1)	Keep as appropriate.					
	(²)	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.				
	(4)	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	column 5 "SG" of Part 1 of Annex II			
	(⁵)	Delete when the exporting country carries out vaccination against authorised to import into the Union matured de-boned meat which fu					
	(6)	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			
	(7)	Supplementary guarantees regarding meats from matured de-boned n to Regulation (EU) No 206/2010, with the entry "F". The matured de-t days after the date of slaughter of the animals.					
	(⁸)	Alternative guarantee may be provided when allowed for by the (EU) No 206/2010.	ə entry "J " in column 5 "SG" of F	Part 1 of Annex II to Regulation			
▶ ⁽¹⁾	(⁹)	OJ L 303, 18.11.2009, p. 1. ◀					
	Official veterinarian						
		Name (in capital letters):	Qualification and title				
		Date:	Signature:				
		Stamp:					

►(1) <u>M13</u>

	col	Mode UNTRY	el POR Veterinary certificate to EU			
			I.2. Certificate reference number I.2.a.			
	1.1.	Consignor Name				
		Address	I.3. Central Competent Authority			
÷		Tel. No	I.4. Local Competent Authority			
men	15	Consignee	1.6.			
ign	1.5.	Name	1.0.			
con		Address				
ber						
atcl		Postal code				
disp		Tel. No				
Part I: Details of dispatched consignment	1.7.	Country ISO I.8. Region Code of origin code of origin	I.9. Country of ISO I.10. Region of Code destination code destination			
Det	l.11.	Place of origin	l.12.			
it l		Name Approval number Address				
å		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 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			
	1.23	. Identification of container/seal number	I.24. Type of packaging			
	1.25	. Commodities certified for: Human consumption				
	1.26		I.27. For import or admission into EU			
	1.28	. Identification of the commodities				
	(5	Species Nature of Treatment App Scientific name) commodity type	roval number establishments Number Net of packages weight			
		Abattoi	r Cutting plant Cold store			

cc	DUNTRY				Model F		
П.	Health	information		II.a. Certificate reference number	II.b.		
11.1	1. Public	Health Attest	ation				
	(EC) N	lo 852/2004, (E	C) No 853/	arian, declare that I am aware of the relevant re 2004 and (EC) No 854/2004 and hereby cert ance with those requirements, in particular th	tify that the meat of domestic swine describe		
	II.1.1] (1) comes from (an) establishment(s) imple with Regulation (EC) No 852/2004;	ementing a programme based on the HACC		
	II.1.2	II.1.2 the meat has been obtained in compliance with the conditions set out in Section I of Annex III to Regula No 853/2004;					
II.1.3 the meat fulfils the requirements of Regulation (EC) No 2075/2005 laying down specific rules on official <i>Trichinella</i> in meat, and in particular:							
		(1) either	[has bee	en subjected to an examination by a digestic	on method with negative results]		
		(1) or	[has bee No 2075	en subjected to a freezing treatment in ad //2005;]	ccordance with Annex II to Regulation (Ed		
_		(1) or	holding	ase of meat from domestic swine kept sole or category of holdings that has been officia n Trichinella in accordance with Annex IV to F	ally recognized by the competent authority a		
(1) II.1.4 [the minced meat has been produced in accordance with Section V of Annex III to Regulation (EC) frozen to an internal temperature of not more than -18 °C;]							
	II.1.5 the meat has been found fit for human consumption following ante and post-mortem inspections carried out accordance with Chapter II of Section I and Chapters IV and IX of Section IV of Annex I to Regulation (E No 854/2004;						
	II.1.6 (1) either		cass or parts of the carcass have been ma III of Section I of Annex I to Regulation (EC)			
		(1) or		kages of [meat] [minced meat] (1) have b nce with Section I of Annex II to Regulation (B			
	II.1.7	the [meat] [mi criteria for foc		(1) satisfies the relevant criteria set out in Reg	gulation (EC) No 2073/2005 on microbiologic		
	II.1.8	0		live animals and products thereof provided hand in particular Article 29, are fulfilled.	by the residue plans submitted in accordance		
	II.1.9			t] (1) has been stored and transported in a ively of Annex III to Regulation (EC) No 853/2			
	(²) [II.1.10			of Regulation (EC) No 1688/2005 implemen erning Salmonella for consignments to Finlar			
11.2	II.2. Animal Health attestation						
	l, the u	undersigned off	icial veterina	arian, hereby certify, that the fresh meat desc	cribed in Part I :		
	II.2.1	has been obt	ained in the	territory/ies with code:	(³) which, at the date of issuing this certificat		
		(1) either		been free for 12 months from foot-and-mo sical swine fever, swine vesicular disease, and			
		(1) <i>or</i>		has been free for 12 months from rinderpest, Ai [classical swine fever] (¹) and [swine vesicular			

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l. He	ealth inform	ation		II.a. Certificate reference number	II.b.
				has been considered free from [foot-and-mot [swine vesicular disease] (1), since had cases/outbreaks afterwards, and author Regulation (EC) No, of	(dd/mm/yyyy), without havin orised to export this meat by Commissio
			imp	ing the last 12 months no vaccination agains orts of domestic animals vaccinated agains itory;	
	II.2.2	has been ob	tained from	animals that:	
		(1) either	-	mained in the territory described under point before slaughter;]	II.2.1 since birth, or for at least the last thre
		(1) or	point II.2	een introduced on(dc 2.1, from the territory with code his fresh meat into the Union;]	
		(1) or	-	een introduced on(dc 2.1, from the EU Member State	
	II.2.3	has been ob	tained from	animals coming from holdings:	
		(a) in which point II.2		he animals present therein have been vaco	cinated against the diseases referred to i
				ı, in an area of 10 km radius, there has been n e previous 40 days,	o case/outbreak of the diseases referred to
		(c) that are weeks;	not subject	t to prohibition as a result of an outbreak of	f porcine brucellosis during the previous s
	(1) (4)			ng has been received that pigs are not fed with the list established by the competent authority	
	II.2.4	has been ob	tained from	animals that:	
		(a) have rem	nained sepa	arate since birth from wild cloven-hoofed anima	als,
			rhouse with	ted from their holdings in vehicles, cleaned ar out contact with other animals which did not co	- · · · ·
				se, have passed ante-mortem health inspectio wn no evidence of the diseases referred to in p	
				red on(dd/mm/yyyy) or (dd/mm/yyyy). (⁵);	between (dd/mm/yyyy
	II.2.5	of the diseas	ses referred of meat for i	n establishment around which, within a radius d to in point II.2.1 during the previous 40 day importation into the Union has been authorise d the total cleaning and disinfection of the e	ys or, in the event of a case of disease, th ed only after slaughter of all animals presen
	II.2.6	has been ob certificate.	ained and p	prepared without contact with other meats not	complying with the conditions required in th
▶ ⁽¹⁾ Ⅱ.3.	Anima	I welfare atte	station		
				arian, hereby certify, that the fresh meat descrit n the slaughterhouse before and at the time of	

COUNTRY Model Pe							
II.		Health information	II.a. Certificate reference number	II.b.			
	No	tes	I				
	This certificate is meant for fresh meat, including minced meat, of domestic swine (Sus scrofa).						
	Fre	esh meat means all animal parts fit for hu	man consumption whether fresh, chilled or fro	ozen.			
	Pa	rt I:					
		Box reference I.8: Provide the code of the	erritory as appearing in Part 1 of Annex II to Re	egulation (EU) No 206/2010.			
	_	Box reference I.11: Place of origin: nam	e and address of the dispatch establishment.				
			er (railway wagons or container and lorries), fli ading, the consignor must inform the BIP of er				
		Box reference I.19: Use the appropriate	HS code: 02.03, 02.06, 02.09, 05.04 or 15.01	1.			
		Box reference I.20: Indicate total gross	weight and total net weight.				
			oxes, the container number and the seal numb				
		Box reference I.28: Nature of commodia	ty: Indicate 'carcass-whole', 'carcass-side', 'ca	rcass-quarters', 'cuts' or 'minced meat'.			
		Minced meat is deboned meat that has muscle (including the adjoining fatty tis	s been minced into fragments and that must h sues) except heart muscle.	nave been prepared exclusively from striated			
	_	Box reference I.28: Treatment type: If ap of freezing (mm/yy) of the cuts/pieces.	propriate, indicate 'deboned'; 'bone in'; 'mature	ed' and/or 'minced'. If frozen, indicate the date			
	Pa	rt II:					
	(1)	Keep as appropriate.					
	(²)	Delete if the consignment is not intende	ed for import into Finland or Sweden.				
	(3)	Code of the territory as it appears in Pa	rt 1 of Annex II to Regulation (EU) No 206/201	0.			
	(4)	Supplementary guarantees to be provise with the entry 'D'.	ded when required in column 5 'SG' of Part 1	of Annex II to Regulation (EU) No 206/2010,			
			od intended for human consumption from resta ins of the farmer or persons tending pigs.	urants, catering facilities or kitchens, including			
	(5)	of authorisation for importation into the	is meat shall not be allowed when obtained from Union of the third country, territory or part there been adopted by the Union against imports of	eof referred to in boxes I.7 and I.8, or during a			
► ⁽¹⁾	(6)	OJ L 303, 18.11.2009, p. 1. ◀					
	Off	icial veterinarian					
	011						
		Name (in capital letters):	Qualification	n and title:			
		Date:	Signature:				
		Stamp:					

			el EQU			
		UNTRY	Veterinary certificate to EU			
	l.1.	Consignor	I.2. Certificate reference number I.2.a.			
		Name	I.3. Central Competent Authority			
		Address	I.4. Local Competent Authority			
lent		Tel. No				
gnm	I.5.	Consignee	1.6.			
onsi		Name				
sd ce		Address				
tche		Postal code				
ispa		Tel. No				
Part I: Details of dispatched consignment	1.7.	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	l.11.	. Place of origin	1.12.			
rt I:		Name Approval number				
Ра		Address				
	I.13.	. Place of loading	I.14. Date of departure			
		-				
	I.15.	. Means of transport Aeroplane Ship Railway wagon	I.16. Entry BIP in EU			
		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			
	1.23	B. Identification of container/seal number	I.24. Type of packaging			
	I.25	i. Commodities certified for: Human consumption				
	1.26		I.27. For import or admission into EU			
	1.28	B. Identification of the commodities	1			
			umber establishments Number Net of packages weight			
		Abattoir C	Cutting plant Cold store			

	COUNT	RY						Model EQU		
	II.	Health	information		II.a. Certificate reference	number	II.b.			
Part II: Certification	II.1.	Public Health Attestation								
		I, the undersigned official veterinarian, declare that I am aware of the relevant requirements of Regulations (EC) No 178/2002 (EC) No 852/2004, (EC) No 853/2004 and (EC) No 854/2004 and hereby certify that the meat of domestic solipeds described in Part I was produced in accordance with those requirements, in particular that:								
		II.1.1			an) establishment(s) imple ion (EC) No 852/2004;	ementing a progra	amme based on t	he HACCP principles in		
t II: Cer		II.1.2	the meat has b No 853/2004;	s been obtained in compliance with the conditions set out in Section I of Annex III to Regulation (EC) ;						
Par		II.1.3 the meat fulfils the requirements of Regulation (EC) No 2075/2005 laying down specific rules on official contr for Trichinella in meat, and in particular, has been subject to an examination by a digestion method with negat results;								
		II.1.4			d fit for human consumptioner II of Section I and Cha					
		II.1.5	(1) either		cass or parts of the carcas III of Section I of Annex I to			mark in accordance with		
			(1) or		ages of meat have been ma to Regulation (EC) No 853/		ification mark in acc	cordance 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	II.1.7 the guarantees covering live animals and products thereof provided by the residue plans submitted in accordance with Directive 96/23/EC, and in particular Article 29 thereof, are fulfilled;							
		II.1.8	the meat has b Regulation (EC		d and transported in accord 2004.	lance with the relev	vant requirements o	of Section I of Annex III to		
	II.2. Animal Health attestation									
		I, the u	ndersigned offici	ial veterina	arian, hereby certify, that the	e fresh meat descri	ibed 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 o	domestic solipeds, which:					
			(1) either		nained in the territory desc before slaughter;]	ribed under point l	I.2.1 since birth, or	for at least the last three		
			(¹) or	point II.2	en introduced on 1, from the territory with co this fresh meat to the Union	ode:				
			(1) <i>or</i>		en introduced on .1, from the EU Member Sta			territory described under		
		II.2.3	which, within a previous 40 da has been auth	radius of ys or, in th orised onl	animals which were slau dd/mm/yyyy) and 10 km, there has been no c e event of a case of such d y after slaughter of all anir shment under the control of	ase/outbreak of Af iseases, the prepa nals present, remo	d/mm/yyyy) (³) in a frican horse sickne tration of meat for it oval of all meat, ar	slaughterhouse around ss or glanders during the mportation into the Union		

	Hea	th inform	nation	II.a. Certificate reference nu	ımber	II.b.	
		II.2.4	has been obtained an certificate.	d prepared without contact with c	other meats not	complying with the conditions required	d in thi
(1)	II.3.	Anima	I welfare attestation				
		which h sions of	ave been handled in the	slaughterhouse before and at the	time of slaughte	d in Part I of this certificate derives from r or killing in accordance with the relevar id down in Chapters II and III of Council I	nt prov
	Notes						
	This cert breeds).	ficate is	meant for fresh meat, e	excluding minced meat, of domes	stic solipeds (E	quus caballus, Equus asinus and their	r cros
	Fresh me	at mean	is all animal parts fit for	human consumption whether free	sh, chilled or fr	ozen.	
	Part I:						
	— Box	eference	e I.8: Provide the code of	of territory as appearing in Part 1	of Annex II to F	legulation (EU) No 206/2010.	
	— Box	eference	e I.11: Place of origin: n	ame and address of the dispatch	establishment		
	prov	ded. In c	ase of unloading and re	eloading, the consignor must info	rm the BIP of e	light number (aircraft) or name (ship) i ntry into the Union.	is to t
				ate HS code: 02.05, 02.06 or 05.0	04.		
			-	ss weight and total net weight.	d the cool num	acr (if applicable) should be included	
				boxes, the container number and boxes, the container number an		per (if applicable) should be included.	
	— Box	reference				d/or 'matured'. If frozen, indicate the	date
	Part II:		,,,,,,,,,,,,,,,,,,,,,,,,,,,,,,,,,,,,,,				
			apriata				
	(¹) Keep	• •		Part 1 of Annex II to Regulation (I	ELI) No 206/20	10	
	(³) Date for in	s: import iportatio	s of this meat shall not l n into the Union of the	be authorised when obtained from third country, territory or part ther	n animals slaug reof referred to	htered either prior to the date of autho in boxes I.7 and I.8, or during a perioc om this third country, territory or part th	d whe
(2)			1.2009, p. 1 . ৰ	.			
	m <u></u>	. .					
	Official v	eterinaria	an				
		Name	(in capital letters):		Qualificatio	on and title:	
		Date:			Signature:		
		Stamp	:				

	<u> </u>	Mode UNTRY	el RUF Veterinary certificate to EU			
		Consignor	I.2. Certificate reference number I.2.a.			
	1.1.	Name	1.2. Certificate reference number 1.2.a.			
		Address	I.3. Central Competent Authority			
		Tel. No	I.4. Local Competent Authority			
nent	1.5					
ign	1.5.	Consignee	1.6.			
suos		Name				
ped (Address				
atch		Postal code				
disp		Tel. No				
Part I: Details of dispatched consignment	1.7.	Country ISO I.8. Region Code of origin code of origin	I.9. Country of ISO I.10. Region of Code destination code destination			
Det	l.11.	. Place of origin	1.12.			
art I:		Name Approval number Address				
Pa		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 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			
	1.23	B. Identification of container/seal number	I.24. Type of packaging			
	1.25	i. Commodities certified for: Human consumption				
	I.26		I.27. For import or admission into EU			
	1.28	B. Identification of the commodities	1			
			roval number establishments Number Net			
	(5	Scientific name) commodity type	of packages weight			
		Abattoi	r Cutting plant Cold store			

	COUNT	RY			Model RUF	
	Ш.	Health	information	II.a. Certificate reference number	II.b.	
tion	II.1.	Public	Health Attestati	on		
		No 178 the me and the	3/2002, (EC) No a at of farmed anir eir cross-breeds)	cial veterinarian, declare that I am aware of the re 852/2004, (EC) No 853/2004, (EC) No 854/2004 and nals of the order Artiodactyla (excluding bovine anim <i>o, Ovis aries, Capra hircus,</i> Suidae and Tayassuidae I in Part I was produced in accordance with those rec	(EC) No 999/2001 and hereby certify that hals (including <i>Bison</i> and <i>Bubalus</i> species), and of the families Rhinocerotidae and	
Part II: Certification		II.1.1		s from (an) establishment(s) implementing a progra Regulation (EC) No 852/2004;	mme based on the HACCP principles in	
Part II		II.1.2	in Section III of Annex III to Regulation (EC)			
		II.1.3		een found fit for human consumption following ante an Chapter II of Section I and Chapters VII and IX of		
		II.1.4	.,	[the carcass or parts of the carcass have been mark Chapter III of Section I of Annex I to Regulation (EC) No		
				[the packages of meat have been marked with a Section I of Annex II to Regulation (EC) No 853/2004		
		II.1.5	the meat satisfi foodstuffs;	es the relevant criteria set out in Regulation (EC) N	o 2073/2005 on microbiological criteria for	
		II.1.6		covering live animals and products thereof provided by /23/EC, and in particular Article 29 thereof, are fulfilled.		
	(¹) (²) [II.1.7 with regard to Chronic Wasting Disease (CWD):					
			animals which hother diagnostic	ntains or is derived exclusively from meat, excludin have been examined for Chronic Wasting Disease by the method recognised by the competent authority with from a herd where Chronic Wasting Disease has been	histopathology, immunohistochemistry or n negative results and is not derived from	
		ll.1.8	the meat has be Regulation (EC)	en 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 attestat	on		
		I, the u	ndersigned officia	l veterinarian, hereby certify, that the fresh meat descri	bed in Part I:	
		II.2.1		ed in the territory/ies with code:	-	
			(a) has been fre has taken pl	ee for 12 months from rinderpest, and during the same ace, and	period no vaccination against this disease	
		(1) either		e for 12 months from foot-and-mouth disease, and dun has taken place;]	ring the same period no vaccination against	
		(1) or	having had o	onsidered free from foot-and-mouth disease since ases/outbreaks afterwards, and authorised to export th f		
		(1) (4) <i>or</i>		programmes against foot-and-mouth disease are be vine animals;]	ing officially carried out and controlled in	

co	COUNTRY Model RU						
Π.	Health	information	II.a. Certificate reference number	II.b.			
	II.2.2	has been obtained from a	nimals that:				
		-	nained in the territory described under point I efore 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	nimals coming from holdings:				
		 (a) in which none of the or] (⁵) rinderpest, 	ne animals present therein have been va	ccinated against [foot-and-mouth disease			
			ary inspections are carried out to diagnose d re not subject to prohibition as a result of an or				
	(1) either	[(c) in and around which in rinderpest during the	n an area of 10 km radius, there has been no previous 30 days,]	case/outbreak of foot-and-mouth disease or			
	(1) (4) or		ial restriction for health reasons and in and ar brack of foot-and-mouth disease or rinderpes				
		(d) where the animals ha	ve remained for at least 40 days before direct	dispatch to the slaughterhouse;]			
	II.2.4	has been obtained from a	nimals:				
	(1) either		nsported from their holdings in vehicles, cle buse, without contact with other animals which				
			rhouse, have passed ante-mortem health ins e shown no evidence of the diseases referred				
			ughtered on(dd/mm/yyyy) (6);]	ı/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 f the animals to an slaughterhouse,	he welfare of the animals or to their handlers			
		 the holding had animals, 	been inspected and authorised by the com	petent authority for the slaughter of game			
			passed the ante-mortem health inspection d shown no evidence of the diseases referred				
		 the animals were (dd/mm/yyyy), (⁶) 	slaughtered between	(dd/mm/yyyy) and			
		 the bleeding of the 	e animals was performed correctly, and				
		 the slaughtered a 	nimals were eviscerated within three hours of	the time of slaughter, and			
		where more than one	ch have been transported to the approved sla hour elapsed since the time of slaughter, a to ival of the vehicle used for the transport;]				
	(¹) (²) II.2.5	[has been obtained from a hoofed animals;]	animals that have remained since birth or for t	the last 3 months separate from wild cloven-			

	Health	informa	ation		II.a. Certificate reference number	II.b.
		II.2.6	of the diseas	ses referred of meat for ir Ill meat, and	to in point II.2.1 during the previous 30 damportation into the Union has been authoris	us of 10 km, there has been no case/outbrea ays or, in the event of a case of disease, th sed only after slaughter of all animals preser establishment under the control of an offici
		II.2.7				
			(1) either	[has bee required		h other meats not complying with the condition
			(¹) (⁴) or	carcasse submitte removed	es in which the main accessible lymphatic d to maturation at a temperature above + 2	ed meat other than offal that was obtained fro glands have been removed, which have bee °C for at least 24 hours before the bones we as below 6.0 when tested electronically in th tion and before de-boning, and
				certificat		nforming to the requirements set out in th oning and storage until it has been packed as.]
			(1) (8) or	carcasse	es in which the main accessible lymphatic d to maturation at a temperature above + 2	ed meat other than offal that was obtained fro glands have been removed, which have bee °C for at least 24 hours before the bones we
				certificat		onforming to the requirements set out in th oning and storage until it has been packed as.]
(1)	(1) II.3.	Anima	al welfare attes	station		
		terhous time of	se, I, the under f slaughter or k	signed officia cilling in acco	al veterinarian, hereby certify, that they were I	which have been slaughtered or killed in a slaug handled in the slaughterhouse before and at th legislation and have met requirements at lea o 1099/2009 ([®]). ◀
	Natas					
	animals (i	includin	ig <i>Bison</i> and B	<i>ubalus</i> speci		ra hircus, Suidae and Tayassuidae), and of th
	This certi animals (i families F	includin Ihinocei	ng <i>Bison</i> and <i>B</i> irotidae and Ele	<i>ubalus</i> speci ephantidae, t	ies and their cross-breeds), Ovis aries, Cap	<i>ra hircus</i> , Suidae and Tayassuidae), and of th n or for the last three months in farms.
	This certi animals (i families F	includin Ihinocei	ng <i>Bison</i> and <i>B</i> irotidae and Ele	<i>ubalus</i> speci ephantidae, t	ies and their cross-breeds), <i>Ovis aries, Cap</i> that are domestically kept or bred since birth	
	This certii animals (i families R Fresh me Part I:	includin Rhinocei at mear	ng <i>Bison</i> and <i>B</i> rotidae and Ele ns all animal pa	<i>ubalus</i> speci ephantidae, t arts fit for hur	ies and their cross-breeds), <i>Ovis aries, Cap</i> that are domestically kept or bred since birth	ra hircus, Suidae and Tayassuidae), and of th n or for the last three months in farms. rozen.
	This certi animals (i families F Fresh me Part I: — Box re	includin Rhinocei at mear eferenc	ng <i>Bison</i> and <i>B</i> rotidae and Ele ns all animal pa ne I.8: Provide t	<i>ubalus</i> speci ephantidae, t arts fit for hur he code of te	ies and their cross-breeds), <i>Ovis aries, Cap</i> that are domestically kept or bred since birth man consumption whether fresh, chilled or f	<i>ra hircus</i> , Suidae and Tayassuidae), and of th n or for the last three months in farms. rozen. Regulation (EU) No 206/2010.
	This certii animals (i families F Fresh me Part I: — Box re — Box re — Box re	includin Rhinocel at mear eferenc eferenc eferenc	ng <i>Bison</i> and <i>B</i> rotidae and Ele ns all animal pa re I.8: Provide ti re I.11: Place of re I.15: Registra	ubalus speci ephantidae, t arts fit for hur he code of te f origin: name ation numbe	ies and their cross-breeds), <i>Ovis aries, Cap</i> that are domestically kept or bred since birth man consumption whether fresh, chilled or f erritory as appearing in Part 1 of Annex II to e and address of the dispatch establishmen r (railway wagons or container and lorries),	<i>ra hircus</i> , Suidae and Tayassuidae), and of th n or for the last three months in farms. rozen. Regulation (EU) No 206/2010. It. flight number (aircraft) or name (ship) is to b
	This certii animals (i families F Fresh me Part I: — Box re — Box re — Box re	includin Rhinocel at mear eferenc eferenc ded. In c	ng <i>Bison</i> and <i>B</i> rotidae and Ele ns all animal pa e I.8: Provide ti re I.11: Place of ree I.15: Registra case of unload	ubalus speci ephantidae, t arts fit for hur he code of te f origin: name ation numbe ing and reloa	ies and their cross-breeds), Ovis aries, Cap that are domestically kept or bred since birth man consumption whether fresh, chilled or f erritory as appearing in Part 1 of Annex II to e and address of the dispatch establishmen	<i>ra hircus</i> , Suidae and Tayassuidae), and of th n or for the last three months in farms. rozen. Regulation (EU) No 206/2010. It. flight number (aircraft) or name (ship) is to b
	This certii animals (i families F Fresh me Part I: — Box r — Box r — Box r — Box r	includin Rhinocel at mear eference eference ded. In c eference	ng <i>Bison</i> and <i>B</i> rotidae and Ele ns all animal pa re I.8: Provide t re I.11: Place of re I.15: Registra case of unload re I.19: Use the	ubalus speci ephantidae, t arts fit for hur he code of te f origin: name ation numbe ing and reloa appropriate	ies and their cross-breeds), <i>Ovis aries, Cap</i> that are domestically kept or bred since birth man consumption whether fresh, chilled or f erritory as appearing in Part 1 of Annex II to e and address of the dispatch establishmen r (railway wagons or container and lorries), ading, the consignor must inform the BIP of HS code: 02.06, 02.08.90 or 05.04.	<i>ra hircus</i> , Suidae and Tayassuidae), and of th n or for the last three months in farms. rozen. Regulation (EU) No 206/2010. It. flight number (aircraft) or name (ship) is to b
	This certii animals (i families F Fresh me Part I: — Box ru — Box ru — Box ru — Box ru — Box ru	includin Rhinocel at mear eferenc eferenc ded. In c eferenc eferenc	ng <i>Bison</i> and <i>B</i> rotidae and Ele ns all animal pa e I.8: Provide t re I.11: Place of re I.15: Registra case of unload re I.19: Use the re I.20: Indicate	ubalus speci ephantidae, t arts fit for hur he code of te f origin: name ation numbe ing and relos appropriate total gross s	ies and their cross-breeds), <i>Ovis aries, Cap</i> that are domestically kept or bred since birth man consumption whether fresh, chilled or f erritory as appearing in Part 1 of Annex II to e and address of the dispatch establishmen r (railway wagons or container and lorries), ading, the consignor must inform the BIP of HS code: 02.06, 02.08.90 or 05.04. weight and total net weight.	<i>ra hircus</i> , Suidae and Tayassuidae), and of th n or for the last three months in farms. rozen. Regulation (EU) No 206/2010. It. flight number (aircraft) or name (ship) is to t entry into the Union.
	This certii animals (i families F Fresh me Part I: — Box r — Box r — Box r — Box r — Box r — Box r	includin Rhinocer at mear eferenc eferenc ded. In c eferenc eferenc eferenc eferenc	ng <i>Bison</i> and <i>B</i> rotidae and Ele ns all animal pa e I.8: Provide t re I.11: Place of re I.15: Registra case of unload re I.19: Use the re I.20: Indicate re I.23: For con	ubalus speci ephantidae, t arts fit for hur he code of te f origin: name ation numbe ing and reloa appropriate e total gross to tainers or bo	ies and their cross-breeds), <i>Ovis aries, Cap</i> that are domestically kept or bred since birth man consumption whether fresh, chilled or f erritory as appearing in Part 1 of Annex II to e and address of the dispatch establishmen r (railway wagons or container and lorries), ading, the consignor must inform the BIP of HS code: 02.06, 02.08.90 or 05.04.	ra hircus, Suidae and Tayassuidae), and of th n or for the last three months in farms. rozen. Regulation (EU) No 206/2010. It. flight number (aircraft) or name (ship) is to t entry into the Union.

	Health information	II.a. Certificate reference number	II.b.
Pa	rt II:	1	
• •			be provided when required in column 5 'SG' of Pa
(³)	-	I No 206/2010, with the entry 'G' . s in Part 1 of Annex II to Regulation (EU) No 2	206/2010
	Supplementary guarantees reg		to be provided when required in column 5 'SG' of
(5)			-mouth disease with serotypes A, O or C, and thi Ifils the supplementary guarantees described unde
(6)	date of authorisation for importa	tion into the Union of the third country, territo	obtained from animals slaughtered either prior to th ory or part thereof referred to in boxes I.7 and I.8, of against imports of this meat from this third country
		animals kept permanently in Arctic regions.	
(8)	of Annex II to Regulation (EU) No		be provided when required in column 5 'SG' of Part be-boned meat shall not be authorised for importatio
(1) (9)	OJ L 303, 18.11.2009, p. 1. ◀		
Of	ficial veterinarian		
	Name (in capital letters):	Qua	lification and title:
	Date:	Sign	nature:
	Stamp:		

	<u> </u>	Mode UNTRY	el RUW Veterinary certificate to EU			
	1.1.	Consignor Name	I.2. Certificate reference number I.2.a.			
		Address	I.3. Central Competent Authority			
_		Tel. No	I.4. Local Competent Authority			
nen	1.5					
ign	1.5.	Consignee	1.6.			
suo:		Name				
edo		Address				
atch		Postal code				
lisp		Tel. No				
Part I: Details of dispatched consignment	1.7.	Country ISO I.8. Region Code of origin code of origin	I.9. Country of ISO I.10. Region of Code destination code destination			
Deti	l.11.	Place of origin	1.12.			
ï		Name 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 wagon				
		Road vehicle Other				
		Identification:	1.17.			
		Documentary references:				
	1.18	. 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		I.27. For import or admission into EU			
	1.28	. Identification of the commodities	1			
	(9	Species Nature of Treatment App Scientific name) commodity type	roval number establishments Number Net of packages weight			
	(Abattoi				

	COUNTRY	(Model RUW
	II.	Health	information	II.a. Certificate reference number	II.b.
	II.1. I	Public	Health Attestation		
ation		No 178 animals <i>Ovis ai</i>	/2002, (EC) No 852/200 s of the order Artiodactyla <i>ries, Capra hircus,</i> Suida	erinarian, declare that I am aware of the re 4 , (EC) No 853/2004 and (EC) No 854/2004 a a (excluding bovine animals (including <i>Bison</i> an ae and Tayassuidae), and of the families Rhin nce with those requirements, in particular that:	nd hereby certify that the fresh meat of wild d <i>Bubalus</i> species and their cross-breeds), ocerotidae and Elephantidae described in
Part II: Certification	I	II.1.1	the meat comes from accordance with Regula	(an) establishment(s) implementing a progra tion (EC) No 852/2004;	mme based on the HACCP principles in
Part II:	I	II.1.2	the meat has been obt 853/2004, and in particu	ained in compliance with the conditions set c llar:	out in Section IV of Annex III to Regulation
			(i) before skinning, it h	as been stored and handled separately from oth	ner food and not frozen;
			and		
			(ii) after skinning, it has	undergone a final inspection as referred to in p	oint II.1.4;
	(1)	ll.1.3		ble species, the meat fulfils the requirements of controls for Trichinella in meat;]	Regulation (EC) No 2075/2005 laying down
	I	II.1.4		d fit for human consumption following a post-m n I and Chapters VIII and IX of Section IV of An	
	l	II.1.5		ase of large wild game, the carcass or parts of t accordance with Chapter III of Section I of Ann	
				kages of meat have been marked with an identi I to Regulation (EC) No 853/2004;]	fication mark in accordance with Section I of
	I	II.1.6	the meat satisfies the foodstuffs;	relevant criteria set out in Regulation (EC) No	o 2073/2005 on microbiological criteria for
	J	II.1.7		live animals and products thereof provided by and in particular Article 29 thereof, are fulfilled.	
	(¹) (²) [[II.1.8	with regard to Chronic V	Vasting Disease (CWD):	
			have been examined for method recognised by t	is derived exclusively from meat, excluding offal r Chronic Wasting Disease by histopathology, he 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
	I	II.1.9	the meat has been store Regulation (EC) No 853	ed and transported in accordance with the relev /2004.	rant requirements of Section I of Annex III to
	II.2.	Animal	Health attestation		
	l	I, the ur	ndersigned official veterir	narian, hereby certify, that the fresh meat descri	bed in Part I:
	ĺ	II.2.1	has been obtained in the	e territory/ies with code:	which, at the date of issuing this certificate:
			(a) has been free for 12 has taken place, an	2 months from rinderpest, and during the same d	period no vaccination against this disease
	(1) eithe	er	(b) has been free for 12 this disease has tak	? months from foot-and-mouth disease, and due en place;]	ring the same period no vaccination against
L					

. Health	information	II.a. Certificate reference number	II.b.
having had cases			since(dd/mm/yyyy), with to export these animals by Commission Regulat /):]
(1) (4) or		programmes against foot-and-mouth diseas vine animals;]	e are being officially carried out and controlled
II.2.2			en (dd/mm/yyyy) a eferred to in point II.2.1, and the killing took place
		hat exceeds 20 km from the borders of a cour iorting this fresh meat into the Union,	try or part thereof, which is not authorised during t
	(b) in an area w point II.2.1;	here during the last 60 days, there has be	een no restrictions for the diseases referred to
II.2.3	game-handling e diseases referred of meat for impor	stablishment around which, within a radius to in point II.2.1 during the previous 30 days	orted as soon as possible for chilling to an approv of 10 km, there has been no case/outbreak of or, in the event of a case of disease, the preparat vafter removal of all meat, and the total cleaning a veterinarian;
II.2.4			
	-	nas been obtained and prepared without conta aquired above.]	ct with other meats not complying with the condition
	c s r	arcasses in which the main accessible lymp ubmitted to maturation at a temperature abov	e-boned meat other than offal that was obtained fr hatic glands have been removed, which have be re +2 °C for at least 24 hours before the bones w eat was below 6.0 when tested electronically in laturation and before de-boning, and
	c		not conforming to the requirements set out in t de-boning and storage until it has been packed ad areas.]
	c	arcasses in which the main accessible lymp	s-boned meat other than offal that was obtained fr hatic glands have been removed, which have be e +2 °C for at least 24 hours before the bones w
	c		not conforming to the requirements set out in t de-boning and storage until it has been packed ad areas.]
otes			
his certificate is	meant for fresh m	eat, excluding offal and minced meat, of wild	animals of the order Artiodactyla (excluding bov

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

СС	DUNTRY		Model RUW				
II.	Health information	II.a. Certificate reference number	II.b.				
Pa	rt I:						
	 Box reference I.8: Provide the code of territory as appearing in Part 1 of Annex II to Regulation (EU) No 206/2010. Box reference I.11: Place of origin: name and address of the dispatch establishment. 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. Box reference I.19: Use the appropriate HS code: 02.01, 02.02, 02.04, 02.06, 02.08.90 or 05.04. 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) should be included. Box reference I.28: <i>Nature of commodity</i>: Indicate 'carcass-whole', 'carcass-side', 'carcass-quarters' or 'cuts'. 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. Box reference I.28: <i>Abattoir</i> : any abattoi						
De	rt II:	r er game nandning esidbilsinnent.					
(1) (2) (3)	Keep as appropriate Supplementary guarantees regarding f of Annex II to Regulation (EU) No 206 Code of the territory as it appears in Par Supplementary guarantees regarding Part 1 of Annex II to Regulation (EU) N The matured de-boned meat shall not animals. Dates. Imports of this meat shall not be a for importation into the Union of the thir restrictive measures have been adopted Supplementary guarantees regarding m	/2010, with the entry 'G'. rt 1 of Annex II to Regulation (EU) No meat from matured de-boned mea No 206/2010 with the entry 'A'. be authorised for importation into th uthorised when obtained from animals d country, territory or part thereof refe d by the Union against imports of this eats from matured de-boned meat to t IO, with the entry 'F'. The matured de	be provided when required in column 5 'SG' of Part 1 206/2010. It to be provided when required in column 5 'SG' of the Union until 21 days after the date of killing of the skilled or hunted either prior to the date of authorisation erred to in boxes 1.7 and 1.8, or during a period where meat from this third country, territory or part thereof. De 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 letters):	Qu	alification and title:				
	Date:	Sig	nature:				
	Stamp:						

	<u></u>	Mode	el SUF Veterinary certificate to EU			
			I.2. Certificate reference number I.2.a.			
	1.1.	Consignor Name				
		Address	I.3. Central Competent Authority			
Ŧ		Tel. No	I.4. Local Competent Authority			
mer	15	Consignee	1.6.			
sign	1.5.	Name	1.0.			
cou		Address				
ber						
atcl		Postal code				
disp		Tel. No				
Part I: Details of dispatched consignment	1.7.	Country ISO I.8. Region Code of origin code of origin	I.9. Country of ISO I.10. Region of Code destination code destination			
Det	l.11.	Place of origin	I.12.			
Ë		Name Approval number				
e		Address				
	I.13.	Place of loading	I.14. Date of departure			
	145	Manage of August and				
	1.15.	Means of transport Aeroplane Ship Railway wagon	I.16. Entry BIP in EU			
		Road vehicle Other				
		Identification: Documentary references:	1.17.			
	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:				
		Human consumption				
	1.26		I.27. For import or admission into EU			
	1.28	Identification of the commodities				
	(5	Scientific name) commodity type	roval number establishments Number Net of packages weight			
		Abatto	r Cutting plant Cold store			

	COUN	FRY				Model SUF
	Ш.	Health	information		II.a. Certificate reference number	II.b.
Part II: Certification	II.1.	Public	Health Attest	ation		
		(EC) N animal	lo 852/2004, (I	EC) No 853 the Suidae	arian declare that I am aware of the relevant p /2004 and (EC) No 854/2004 and hereby ce , Tayassuidae, or Tapiridae families described that:	rtify that the meat of farmed non-domestic
		II.1.1			an) establishment(s) implementing a progra ion (EC) No 852/2004;	mme based on the HACCP principles in
		II.1.2	the meat has No 853/2004		ned in compliance with the conditions set out	in Section III of Annex III to Regulation (EC)
č		II.1.3			rements of Regulation (EC) No 2075/2005 lay nd in particular, has been subject to an exami	
		II.1.4		with, Chapt	d fit for human consumption following ante a er II of Section I and, Chapters VII and IX of	
		II.1.5	(1) either		cass or parts of the carcass have been mark III of Section I, of Annex I to Regulation (EC) N	
			(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 sat foodstuffs;	isfies the re	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 Regulation (E		d and transported in accordance with the relev 2004.	rant requirements of Section I of Annex III to
	II.2.	Anima	I Health attes	tation		
		I, the u	ndersigned off	icial veterina	arian, hereby certify, that the fresh meat descri	bed in Part I:
		II.2.1	has been obt	ained in the	territory/ies with code:	ch, at 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		nas been free for 12 months from rinderpest, Afric classical swine fever] (') and [swine vesicular d	
				[has been considered free from [foot-and-mout swine vesicular disease] (1), since ad cases/outbreaks afterwards, and author Regulation (EU) No/, of	(dd/mm/yyyy), without having ised to export this meat by Commission
				. ,	ng the last 12 months no vaccination against orts of domestic animals vaccinated against ory;	
		II.2.2	has been obt	ained from a	animals that:	
			(1) either	-	nained in the territory described under point II pefore slaughter;]	.2.1 since birth, or for at least the last three

II.	Hoalth	information		II.a. Certificate reference number	II.b.
	пеаш	mormation		II.a. Certificate reference number	11.0.
		(1) or	point II.:	veen introduced on(dd/ 2.1, from the territory with code his fresh meat into the Union;]	
	II.2.3	has been obta	ned from	animals coming from holdings:	
		(a) in which i point II.2.1		the animals present therein have been vacci	nated against the diseases referred to
				h in an area of 10 km radius, there has been no ne previous 40 days,	case/outbreak of the diseases referred to
			holdings	erinary inspections are carried out to diagnose d s are not subject to prohibition as a result of ar	
	II.2.4	has been obta	ned from	animals which:	
		(1) either	to a	ve been transported from their holdings in vehicl an approved slaughterhouse without contact with nditions mentioned above,	
				he slaughterhouse, have passed ante-mortem h ughter and, in particular, have shown no eviden d	
				/e been slaughtered on(dd //mm/yyyy) and(dd/mm/	
		(1) <i>or</i>	,	re been slaughtered on the holding of origin, follc ponsible for the holding, who has provided a writ	•
			_	in his opinion an unacceptable risk would have to their handlers by the transport of the animals	
			—	the holding had been inspected and authorised of game,	l by the competent authority for the slaught
			_	the animals have passed the ante-mortem he the slaughter and, in particular, have shown point II.2.1,	
			_	the animals were slaughtered between	(dd/mm/yyyy) ar
			_	the bleeding of the animals was performed corr	rectly, and
			-	the slaughtered animals were eviscerated within	in three hours of the time of slaughter, and
			cor tem	ir carcasses have been transported to the a nditions and, where more than one hour of nperature of between 0 °C and + 4 °C has been the transport;]	elapsed since the time of slaughter,
	II.2.5	has been obta	ned from	animals that have remained separate since birt	th from wild cloven-hoofed animals;
	II.2.6	of the disease preparation of	s referre meat for	n establishment around which, within a radius of d to in point II.2.1 during the previous 40 days importation into the Union has been authorised ad the total cleaning and disinfection of the est	s or, in the event of a case of disease, th d only after slaughter of all animals preser
	II.2.7	has been obtai certificate.	ned and	prepared without contact with other meats not co	omplying with the requirements set out in th

C

COUN	COUNTRY Model SUF					
II.	Healt	h information	II.a. Certificate reference number	II.b.		
► ⁽¹⁾	II.3.	which have been handled in the sla	aughterhouse before and at the time of s	escribed in Part I of this certificate derives from animals aughter or killing in accordance with the relevant provi- hose laid down in Chapters II and III of Council Regula-		
	Notes	for the internet for free large to a	al allow a field and a classical and a field and			
		families that are domestically kept		I animals belonging to the Suidae, Tayassuidae, or		
	Fresh me	at means all animal parts fit for hu	man consumption, whether fresh, chille	ed or frozen.		
	Part I:					
			erritory as appearing in Part 1 of Annex			
		•	e and address of the dispatch establish r (railway wagons or container and lor	nment. ries), flight number (aircraft) or name (ship) is to be		
	provid	ded. In case of unloading and reloa	ading, the consignor must inform the B			
		eference I.19: Use the appropriate eference I.20: Indicate total gross	HS code: 02.03, 02.08.90 or 05.04.			
			•	I number (if applicable) should be included.		
	— Box r	eference I.28: Nature of commodit	y: Indicate 'carcass-whole', 'carcass-si	de', 'carcass-quarters' or 'cuts'.		
		eference I.28: <i>Treatment type</i> : If a uts/pieces.	opropriate indicate deboned, or bone-i	n. If frozen, indicate the date of freezing (mm/yy) of		
	Part II:					
	., ,	as appropriate				
	(³) Date of aut perior	or dates of slaughter. Imports of thi horisation for importation into the l	Union of the third country, territory or pa	106/2010. ned from animals slaughtered either prior to the date art thereof referred to in boxes I.7 and I.8, or during a aports of this meat from this third country, territory or		
► ⁽²⁾	(4) OJL3	303, 18.11.2009, p. 1. ◀				
	Official ve	eterinarian				
		Name (in capital letters):	Qua	ification and title:		
		Date:		ature:		
		Stamp:	e.g.			
		otanp.				

	co	Mode UNTRY	I SUW Veterinary certificate to EU			
		Consignor	I.2. Certificate reference number I.2.a.			
	1.1.	Name				
		Address	I.3. Central Competent Authority			
ŧ		Tel. No	I.4. Local Competent Authority			
Inel	I.5.	Consignee	1.6.			
Isigr		Name				
S		Address				
hed		Postal code				
patc		Tel. No				
Part I: Details of dispatched consignment	1.7.	Country ISO I.8. Region Code of origin	I.9. Country of ISO I.10. Region of Code destination code destination			
etai	111	Place of origin	1.12.			
<u>-</u>	1.111	Name Approval number				
Par		Address				
	140	Disco of locality a				
	1.13.	Place of loading	I.14. Date of departure			
	I.15.	Means of transport Aeroplane Ship Railway wagon	I.16. Entry BIP in EU			
		Road vehicle Other				
		Identification: Documentary references:	l.17.			
	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			
	1.23	. 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			
	1.28	. Identification of the commodities				
	(5	Species Nature of Treatment App Scientific name) commodity type	roval number establishments Number Net of packages weight			
		Abattoi	r Cutting plant Cold store			

	COUNT	RY				Model SUW			
	Ш.	Health	information		II.a. Certificate reference number	II.b.			
ion	II.1.	Public	Health Attestation						
		(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 iridae families described in Part I was produced	that the meat of wild animals belonging to			
Part II: Certification		II.1.1			(an) establishment(s) implementing a progra ttion (EC) No 852/2004;	mme based on the HACCP principles in			
rt II: Cel		II.1.2	the meat has bee particular:	n obt	ained in accordance with Section IV of Annex	III to Regulation (EC) No 853/2004, an in			
Ра			(i) before skinning	g, it h	as been stored and handled separately from oth	ner food and not frozen;			
			and						
			(ii) after skinning,	it has	undergone a final inspection as referred to in p	oint II.1.4;			
		II.1.3			uirements of Regulation (EC) No 2075/2005 lay and in particular, has been subject to an exami	5 I			
		II.1.4			d fit for human consumption following a post-m n I and Chapters VIII and IX of Section IV of An				
		II.1.5			rcass or parts of the carcass have been mark r III of Section I of Annex I to Regulation (EC) No				
					kages of meat have been marked with an identi I to Regulation (EC) No 853/2004;]	fication mark in accordance with Section I of			
		II.1.6	the meat satisfies foodstuffs;	the	relevant criteria set out in Regulation (EC) No	o 2073/2005 on microbiological criteria for			
		II.1.7	the guarantees covering live animals and products thereof provided by the residue plans submitted in accordance with Directive 96/23/EC, and in particular Article 29 thereof, are fulfilled.						
		II.1.8	the meat has been Regulation (EC) N		ed and transported in accordance with the relev /2004	rant requirements of Section I of Annex III to			
	II.2.	Anima	I Health attestation	ı					
		I, the u	ndersigned official v	eterir	narian, hereby certify, that the fresh meat descri	bed in Part I:			
		II.2.1	has been obtained	in the	e territory/ies with code:	t the date of issuing this certificate:			
			(1) either [(a	,	been free for 12 months from foot-and-mout sciences, and sciences, swine vesicular disease, and				
			(1) or [(a) (i)	has been free for 12 months from rinderpest, Afric [classical swine fever] (1) and [swine vesicular d				
				(ii)	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			
			(b)	imp	ing the last 12 months no vaccination against orts of domestic animals vaccinated against itory;				

I. Health	information		II.a. Certificate reference number	II.b.
			wild animals that were killed between d/mm/yyyy) (3) inside the territory referred to i	
			eeds 20 km from the borders of a country or pathis fresh meat into the Union,	art thereof, which is not authorised during th
	(b) in an ar point II.2		uring the last 60 days, there has been no	restrictions for the diseases referred to i
II.2.3.A	centre, and i of 10 km, the in the event	mmediately are has been of a case of I of all meat,	animals which after killing were transported afterwards] (1) to an approved game-handling no case/outbreak of the diseases referred to disease, the preparation of meat for importal and the total cleaning and disinfection of the	gestablishment around which, within a radiu in point II.2.1 during the previous 40 days o tion into the Union has been authorised on
(1) (4) [II.2.3.B	has been ob negative res		carcasses on which the following test for class	ical swine fever was carried out and provide
	(1) either	[virus isc	plation from blood (EDTA);]	
	(1) or	[virus isc	plation from samples of	
	(1) or	[immuno	ofluorescence for viral antigen on samples of .	
	()		5	,
II.2.4	has been ob certificate.	tained and p	repared without contact with other meats not	complying with the conditions required in th
		tained and p	repared without contact with other meats not	complying with the conditions required in th
otes his certificate is	certificate.	sh meat, ex	cluding offal and minced meat, of wild anima	
lotes his certificate is apiridae familie:	certificate. s meant for fre s that are killed	sh meat, ex	cluding offal and minced meat, of wild anima	als belonging to the Suidae, Tayassuidae, o
otes his certificate is apiridae families resh meat mear	certificate. s meant for fre s that are killed ns all animal pa	sh meat, ex I or hunted ir arts fit for hui	cluding offal and minced meat, of wild anima n the wild.	als belonging to the Suidae, Tayassuidae, o
otes his certificate is apiridae families resh meat meal fter importation	certificate. s meant for fre s that are killed ns all animal pa	sh meat, ex I or hunted ir arts fit for hui	cluding offal and minced meat, of wild anima n the wild. man consumption whether fresh, chilled or fro	als belonging to the Suidae, Tayassuidae, o
otes his certificate is apiridae families resh meat meal fter importation art I:	certificate. s meant for fre s that are killed ns all animal pa , unskinned ca	ish meat, exi l or hunted ir arts fit for hui	cluding offal and minced meat, of wild anima n the wild. man consumption whether fresh, chilled or fro st be conveyed without delay to the processing	als belonging to the Suidae, Tayassuidae, ozen. g establishment of destination.
otes his certificate is apiridae families resh meat mean fter importation art I: - Box referenc	certificate. s meant for fre s that are killed ns all animal pa , unskinned ca e I.8: Provide t	ish meat, exi l or hunted ir arts fit for hui arcasses mus he code of te	cluding offal and minced meat, of wild anima n the wild. man consumption whether fresh, chilled or fro	als belonging to the Suidae, Tayassuidae, o ozen. g establishment of destination. egulation (EU) No 206/2010.
otes his certificate is apiridae families resh meat mean fter importation art I: - Box referenc - Box referenc - Box referenc	certificate. s meant for fre s that are killed ns all animal pa , unskinned ca e I.8: Provide t e I.11: Place o e I.15: Registra	ish meat, ex l or hunted ir arts fit for hun ircasses mus he code of te f origin: nam- ation numbe	cluding offal and minced meat, of wild anima I the wild. man consumption whether fresh, chilled or fro st be conveyed without delay to the processing erritory as appearing in Part 1 of Annex II to Re	als belonging to the Suidae, Tayassuidae, ozen. g establishment of destination. egulation (EU) No 206/2010. ight number (aircraft) or name (ship) is to t
otes his certificate is apiridae families resh meat mean fter importation art I: - Box referenc - Box referenc provided. In o	certificate. s meant for fre s that are killed ns all animal pa , unskinned ca e I.8: Provide t e I.11: Place o e I.15: Registra case of unload	Ish meat, ex I or hunted ir arts fit for hun arcasses mus he code of te f origin: nam- ation numbe ing and reloa	cluding offal and minced meat, of wild anima n the wild. man consumption whether fresh, chilled or fro st be conveyed without delay to the processing erritory as appearing in Part 1 of Annex II to Re e and address of the dispatch establishment. or (railway wagons or container and lorries), fil	als belonging to the Suidae, Tayassuidae, o ozen. g establishment of destination. egulation (EU) No 206/2010. ight number (aircraft) or name (ship) is to b
lotes his certificate is apiridae families resh meat meat fter importation art I: - Box referenc - Box referenc provided. In o - Box referenc	certificate. s meant for fre s that are killed ns all animal pa , unskinned ca e I.8: Provide t e I.11: Place o e I.15: Registra case of unload e I.19: Use the	Ish meat, ex l or hunted ir arts fit for hun arcasses mus he code of te f origin: nam- ation numbe ing and reloa	cluding offal and minced meat, of wild anima n the wild. man consumption whether fresh, chilled or fro st be conveyed without delay to the processing erritory as appearing in Part 1 of Annex II to Re e and address of the dispatch establishment. or (railway wagons or container and lorries), fil ading, the consignor must inform the BIP of er	als belonging to the Suidae, Tayassuidae, o ozen. g establishment of destination. egulation (EU) No 206/2010. ight number (aircraft) or name (ship) is to b
lotes his certificate is apiridae families resh meat meau fter importation art I: - Box referenc - Box referenc provided. In - Box referenc - Box referenc - Box referenc	certificate. s meant for fre s that are killed ns all animal pa , unskinned ca 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	Ish meat, exi l or hunted in arts fit for hun arcasses mus he code of te f origin: nam- ation numbe ing and reloa appropriate e total gross o	cluding offal and minced meat, of wild anima n the wild. man consumption whether fresh, chilled or fro st be conveyed without delay to the processing erritory as appearing in Part 1 of Annex II to Re e and address of the dispatch establishment. rr (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.	als belonging to the Suidae, Tayassuidae, o ozen. g establishment of destination. egulation (EU) No 206/2010. ight number (aircraft) or name (ship) is to b
lotes his certificate is apiridae families resh meat meau fter importation art I: - Box referenc - Box referenc - Box referenc - Box referenc - Box referenc - Box referenc - Box referenc	certificate. s meant for fre s that are killed ns all animal pa , unskinned ca 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	sh meat, ex l or hunted ir arts fit for hun ircasses mus he code of te f origin: nam ation numbe ing and reloa appropriate total gross v tainers or bo	cluding offal and minced meat, of wild anima n the wild. man consumption whether fresh, chilled or fro st be conveyed without delay to the processing erritory as appearing in Part 1 of Annex II to Re e and address of the dispatch establishment. or (railway wagons or container and lorries), fl ading, the consignor must inform the BIP of er HS code: 02.03, 02.08.90 or 05.04. weight and total net weight.	als belonging to the Suidae, Tayassuidae, o ozen. g establishment of destination. egulation (EU) No 206/2010. ight number (aircraft) or name (ship) is to b ntry into the Union.
otes his certificate is apiridae families resh meat meat fter importation art I: - Box referenc - Box referenc	certificate. s meant for fre s that are killed ns all animal pa , unskinned ca 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 e I.28: <i>Nature</i> o e I.28: <i>Treatme</i>	sh meat, ex l or hunted ir arts fit for hun arcasses mus he code of te f origin: nam- ation numbe ing and reloa e appropriate total gross to tainers or bo of commodit	cluding offal and minced meat, of wild anima n the wild. man consumption whether fresh, chilled or fro st be conveyed without delay to the processing erritory as appearing in Part 1 of Annex II to Re e and address of the dispatch establishment. or (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. ixxes, the container number and the seal numb	als belonging to the Suidae, Tayassuidae, o ozen. g establishment of destination. egulation (EU) No 206/2010. ight number (aircraft) or name (ship) is to b ntry into the Union. per (if applicable) should be included. ircass-quarters' or 'cuts'.

COUNTRY

COUNTR	YY		Model SUW
Ш.	Health information	II.a. Certificate reference number	II.b.

Part II:

- (1) Keep as appropriate.
- (2) Code of the territory as it appears in Part 1 of Annex II to Regulation (EU) No 206/2010.
- (3) Dates. Imports of this meat shall not be authorised when obtained from animals killed or hunted either prior to the date of authorisation for importation into the Union of the third country, territory or part thereof referred to in boxes reference 1.7 and 1.8, 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.
- (4) Supplementary guarantees to be provided when required in column 5 'SG' of Part 1 of Annex II to Regulation (EU) No 206/2010, with the entry 'C'. For such purpose, in tests other than EDTA, the samples to be used are a sample of tonsil and of spleen plus a sample of ileum or kidney and a sample of at least one of the following lymph nodes: retropharyngeal, parotid, mandibular or mesenteric. The samples used shall be indicated.

Official veterinarian

Name (in capital letters):

Date:

Stamp:

Qualification and title:

Signature:

	~~		el EQW			
		UNTRY	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			
nent		Tel. No				
ignr	1.5.	Consignee	1.6.			
suos		Name				
equ		Address				
atch		Postal code				
disp		Tel. No				
Part I: Details of dispatched consignment	1.7.	Country ISO I.8. Region Code of origin code of origin	I.9. Country of ISO I.10. Region of Code destination code destination			
Det	l.11.	Place of origin	I.12.			
i ti		Name Approval number Address				
a		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 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			
	1.23	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			
	1.28	. Identification of the commodities	1			
	(5	Species Nature of Approval nu Scientific name) commodity	Imber establishments Number Net of packages weight			
			utting plant Cold store			

				Model EQ			
II.	Health	information	II.a. Certificate reference number	II.b.			
II.1.	Public Health Attestation						
	I, the undersigned official veterinarian, declare that I am aware of the relevant requirements of Regulations (EC) No 178/2002, (EC) No 852/2004, (EC) No 853/2004 and (EC) No 854/2004 and hereby certify that the meat of wild solipeds belonging to the subgenus <i>Hippotigris</i> (zebra) described in Part I was produced in accordance with those requirements, in particular that:						
	II.1.1		(an) establishment(s) implementing a prog ation (EC) No 852/2004;	ramme based on the HACCP principles in			
	II.1.2	the meat was obtained	in compliance with Section IV of Annex III to Re	egulation (EC) No 853/2004;			
	II.1.3		irements of Regulation (EC) No 2075/2005 lay articular, has been subject to an examination b				
	II.1.4		nd fit for human consumption following a post- on I and Chapters VIII and IX of Section IV of A				
	II.1.5	• •	arcass or parts of the carcass have been ma er III of Section I of Annex I to Regulation (EC) I				
		.,	ckages of meat have been marked with an ider II to Regulation (EC) No 853/2004;]	tification mark in accordance with Section I of			
	II.1.6	the meat satisfies the foodstuffs;	relevant criteria set out in Regulation (EC) I	No 2073/2005 on microbiological criteria for			
	II.1.7	U	g live animals and products thereof provided t b, and in particular Article 29 thereof, are fulfille	, , , , , , , , , , , , , , , , , , , ,			
	II.1.8 the meat has been stored and transported in accordance with the relevant requirements of Section I of Annex Regulation (EC) No 853/2004.						
	II.1.8			evant requirements of Section I of Annex III to			
II.2.				evant requirements of Section I of Annex III to			
II.2.	Anima	Regulation (EC) No 85					
II.2.	Anima	Regulation (EC) No 85 I Health attestation Indersigned official veter has been obtained fro	3/2004.	ribed in Part I: 			
II.2.	Anima I, the u	Regulation (EC) No 85 I Health attestation Indersigned official veter has been obtained fro centre, and immediate of 10 km, there has be the event of a case of s	3/2004. Inarian, hereby certify, that the fresh meat desc m wild animals that were killed between	ribed in Part I: 			
II.2.	Anima I, the u II.2.1	Regulation (EC) No 85 I Health attestation Indersigned official veter has been obtained fro centre, and immediate of 10 km, there has bee the event of a case of s after removal of all mea veterinarian;	3/2004. Inarian, hereby certify, that the fresh meat desc m wild animals that were killed between (dd/mm/yyyy) (²) inside the territory/ies with co n wild animals which after killing were transpor y afterwards] (¹) to an approved game-handling an no case/outbreak of African horse sickness uch diseases, the preparation of meat for expo	ribed in Part I: 			
II.2.	Anima I, the u II.2.1 II.2.2	Regulation (EC) No 85	3/2004. Inarian, hereby certify, that the fresh meat desc m wild animals that were killed between (dd/mm/yyyy) (²) inside the territory/ies with co- n wild animals which after killing were transpor y afterwards] (¹) to an approved game-handling en no case/outbreak of African horse sickness uch diseases, the preparation of meat for expo t, and the total cleaning and disinfection of the	ribed in Part I: 			
II.2 .	Anima I, the u II.2.1 II.2.2	Regulation (EC) No 85	3/2004. Inarian, hereby certify, that the fresh meat desc m wild animals that were killed between (dd/mm/yyyy) (²) inside the territory/ies with co- n wild animals which after killing were transpor y afterwards] (¹) to an approved game-handling en no case/outbreak of African horse sickness uch diseases, the preparation of meat for expo t, and the total cleaning and disinfection of the	ribed in Part I: 			
II.2.	Anima I, the u II.2.1 II.2.2	Regulation (EC) No 85	3/2004. Inarian, hereby certify, that the fresh meat desc m wild animals that were killed between (dd/mm/yyyy) (²) inside the territory/ies with co- n wild animals which after killing were transpor y afterwards] (¹) to an approved game-handling en no case/outbreak of African horse sickness uch diseases, the preparation of meat for expo t, and the total cleaning and disinfection of the	ribed in Part I: 			
Notes	Anima I, the u II.2.1 II.2.2 II.2.3	Regulation (EC) No 85 I Health attestation Indersigned official veter has been obtained fro centre, and immediate of 10 km, there has be the event of a case of s after removal of all mea veterinarian; has been obtained and certificate.	3/2004. Inarian, hereby certify, that the fresh meat desc m wild animals that were killed between (dd/mm/yyyy) (²) inside the territory/ies with co- n wild animals which after killing were transpor y afterwards] (¹) to an approved game-handling en no case/outbreak of African horse sickness uch diseases, the preparation of meat for expo t, and the total cleaning and disinfection of the	ribed in Part I: 			
Notes This c (zebra	Anima I, the u II.2.1 II.2.2 II.2.3	Regulation (EC) No 85	3/2004. Inarian, hereby certify, that the fresh meat desc m wild animals that were killed between (dd/mm/yyyy) (²) inside the territory/ies with co- n wild animals which after killing were transpor y afterwards] (¹) to an approved game-handling en no case/outbreak of African horse sickness uch diseases, the preparation of meat for expo tt, and the total cleaning and disinfection of the prepared without contact with other meats not o	ribed in Part I: 			

	Health information	II.a. Certificate reference numb	per	II.b.
art I:				
	x reference I 8. Provide the code	e of territory as appearing in Part 1 of A	Annex II to Rec	ulation (EU) No 206/2010
		name and address of the dispatch est	-	
				ht number (aircraft) or name (ship) is to be
-		I reloading, the consignor must inform	the BIP of entr	ry into the Union.
		priate HS code: 02.08.90 or 05.04.		
	•	ross weight and total net weight.		
		or boxes, the container number and th nodity: Indicate 'carcass-whole', 'carca		· · · · ·
				ozen, indicate the date of freezing (mm/yy)
	the cuts/pieces.	an appropriate, indicate matured of t		ozen, indicate the date of neezing (him/yy)
– Bo	x reference I.28: <i>Abattoir</i> : any at	battoir or game handling establishmen	ıt.	
Part II:	:			
	ep as appropriate.			
				unted either prior to the date of authorisation boxes I.7 and I.8, or during a period where
				this third country, territory or part thereof.
³) Co	de of the territory as it appears i	in Part 1 of Annex II to Regulation (EU)) No 206/2010	
Official	lveterinarian			
	Name (in capital letters):		Qualification	and title:
	Date:		Signature:	
	Stamp:			

ANNEX III

Model	TRANS	SIT/ST	ORAGE

	COL	JNTRY	Veterinary certificate to EU				
	l.1.	Consignor	I.2. Certificate reference number I.2.a.				
		Name	I.3. Central Competent Authority				
		Address					
eut		Tel. No	I.4. Local Competent Authority				
ŭ	I.5.	Consignee	I.6. Person responsible for the consignment in EU				
nsiç		Name	Name				
o p		Address	Address				
tche		Postal code	Postal code				
ispa		Tel. No	Tel. No				
Part I: Details of dispatched consignment	1.7.	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	l.11.	Place of origin	I.12. Place of destination				
Ë		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				
ŀ	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				
-	1.23	Identification of container/seal number	I.24. Type of packaging				
	I.25	Commodities certified for:					
-	1.26	For transit through EU to 3 rd Country ISO code	1.27.				
-	1.00						
		Identification of the commodities Species Nature of Treatment Approval nu Scientific name) commodity type	mber establishments Number Net of packages weight				
		Abattoir	Cutting manufacturing plant/ plant				

11.	Health informat	ion	II.a. Certificate reference number	II.b.
.1.	Animal Health	Attestation		
	I, the undersign	ed official veteri	narian, hereby certify, that the fresh meat (described in Part I:
			r region authorized for imports into the Un e time of slaughter, and	ion as laid down in Part 1 of Annex II to Regulation
	certifica			wn in the animal health attestation in the mode [EQW] (') in Part 2 of Annex II to Regulation (EU
			s which were slaughtered and processe 	ed on
-				
	certificate is meant fo		rage in accordance with Article 12(4) or Ar	ticle 13 of Directive 97/78/EC of:
This — f	certificate is meant fo fresh meat, including r	minced meat, of:		
This — f	certificate is meant fo fresh meat, including r (1) domestic bovin	ninced meat, of: e animals (incluc	ding <i>Bubalus</i> and <i>Bison</i> species and their	cross-breeds) (Model 'BOV');
This — f	certificate is meant fo fresh meat, including r (1) domestic bovin (2) domestic ovine	ninced meat, of: e animals (incluc animals (<i>Ovis a</i> l	ding <i>Bubalus</i> and <i>Bison</i> species and their ries) or domestic caprine animals (<i>Capra I</i>	cross-breeds) (Model 'BOV');
This — f (certificate is meant fo fresh meat, including r (1) domestic bovin (2) domestic ovine (3) domestic porcir	ninced meat, of: e animals (incluc animals (<i>Ovis a</i> n ne animals (<i>Sus</i>	ding <i>Bubalus</i> and <i>Bison</i> species and their ries) or domestic caprine animals (<i>Capra i</i> scrofa) (Model 'POR');	cross-breeds) (Model 'BOV');
This — f ((((certificate is meant fo fresh meat, including r (1) domestic bovin (2) domestic ovine (3) domestic porcir fresh meat, excluding	ninced meat, of: e animals (incluc animals (<i>Ovis a</i> . ne animals (<i>Sus</i> minced meat, of	ding <i>Bubalus</i> and <i>Bison</i> species and their ries) or domestic caprine animals (<i>Capra I</i> scrofa) (Model 'POR'); :	cross-breeds) (Model 'BOV'); <i>hircus</i>) (Model 'OVI');
This — f (((((certificate is meant fo fresh meat, including r (1) domestic bovin (2) domestic ovine (3) domestic porcir fresh meat, excluding (4) domestic solipe	ninced meat, of: e animals (incluo animals (<i>Ovis a</i> ne animals (<i>Sus</i> minced meat, of ds (<i>Equus caba</i>	ding <i>Bubalus</i> and <i>Bison</i> species and their ries) or domestic caprine animals (<i>Capra I</i> scrofa) (Model 'POR'); : : Ilus, Equus asinus and their cross-breeds	cross-breeds) (Model 'BOV'); <i>hircus</i>) (Model 'OVI');
This — f (((((((((((((certificate is meant fo fresh meat, including r (1) domestic bovin (2) domestic ovine (3) domestic porcir fresh meat, excluding (4) domestic solipe fresh meat, excluding (5) farmed non-dor	ninced meat, of: e animals (incluo animals (<i>Ovis a</i> ne animals (<i>Sus</i> minced meat, of ds (<i>Equus caba</i> offal and minceo nestic animals o	ding <i>Bubalus</i> and <i>Bison</i> species and their ries) or domestic caprine animals (<i>Capra I</i> scrofa) (Model 'POR'); : <i>Ilus, Equus asinus</i> and their cross-breeds d meat, of: f the order Artiodactyla (excluding bovine a	cross-breeds) (Model 'BOV'); <i>hircus</i>) (Model 'OVI');) (Model 'EQU'); animals (including <i>Bison</i> and <i>Bubalus</i> species an
This — f (((((((((((((certificate is meant for fresh meat, including r (1) domestic boving (2) domestic ovine (3) domestic porcir fresh meat, excluding (4) domestic solipe fresh meat, excluding (5) farmed non-dor their cross-bree (Model 'RUF'); (6) wild non-domestic 	ninced meat, of: e animals (incluc animals (<i>Ovis a</i> minced meat, of ds (<i>Equus caba</i> offal and minced nestic animals o ds), <i>Ovis aries</i> , (stic animals of th	ding <i>Bubalus</i> and <i>Bison</i> species and their ries) or domestic caprine animals (<i>Capra I</i> <i>scrofa</i>) (Model 'POR'); : : <i>Ilus, Equus asinus</i> and their cross-breeds d meat, of: f the order Artiodactyla (excluding bovine a <i>Capra hircus</i> , Suidae and Tayassuidae), an ne order Artiodactyla (excluding bovine a	cross-breeds) (Model 'BOV'); hircus) (Model 'OVI');) (Model 'EQU'); animals (including <i>Bison</i> and <i>Bubalus</i> species an d of the families Rhinocerotidae and Elephantidae nimals (including <i>Bison</i> and <i>Bubalus</i> species an
This — 1 (((((((((((((certificate is meant for fresh meat, including r (1) domestic boving (2) domestic ovine (3) domestic porcir fresh meat, excluding (4) domestic solipe fresh meat, excluding (5) farmed non-dor their cross-bree (Model 'RUF'); (6) wild non-domestic their cross-bree (Model 'RUW'); 	ninced meat, of: e animals (incluo animals (<i>Ovis a</i> me animals (<i>Sus</i> minced meat, of eds (<i>Equus caba</i> offal and minceo nestic animals of ds), <i>Ovis aries</i> , (stic animals of th ds), <i>Ovis aries</i> , (ding <i>Bubalus</i> and <i>Bison</i> species and their ries) or domestic caprine animals (<i>Capra I</i> <i>scrofa</i>) (Model 'POR'); : : <i>Ilus, Equus asinus</i> and their cross-breeds d meat, of: f the order Artiodactyla (excluding bovine a <i>Capra hircus</i> , Suidae and Tayassuidae), an ne order Artiodactyla (excluding bovine a	cross-breeds) (Model 'BOV'); h <i>ircus</i>) (Model 'OVI');) (Model 'EQU'); animals (including <i>Bison</i> and <i>Bubalus</i> species and d of the families Rhinocerotidae and Elephantidae nimals (including <i>Bison</i> and <i>Bubalus</i> species and d of the families Rhinocerotidae and Elephantidae
This — 1 (((((((((((((certificate is meant for fresh meat, including r (1) domestic boving (2) domestic porcing (3) domestic porcing (3) domestic porcing (4) domestic solipe fresh meat, excluding (4) domestic solipe fresh meat, excluding (5) farmed non-domestic ross-bree (Model 'RUF'); (6) wild non-domestic ross-bree (Model 'RUV'); (7) farmed non-domestic ross-bree 	ninced meat, of: e animals (includ animals (<i>Ovis a</i> , ne animals (<i>Sus</i> minced meat, of rds (<i>Equus caba</i> offal and minced offal and minced offal and minced offal and minced stic animals of th ds), <i>Ovis aries</i> , of nestic animals b	ding <i>Bubalus</i> and <i>Bison</i> species and their ries) or domestic caprine animals (<i>Capra I</i> <i>scrofa</i>) (Model 'POR'); <i>llus, Equus asinus</i> and their cross-breeds d meat, of: f the order Artiodactyla (excluding bovine a <i>Capra hircus</i> , Suidae and Tayassuidae), an the order Artiodactyla (excluding bovine and <i>Capra hircus</i> , Suidae and Tayassuidae), an	cross-breeds) (Model 'BOV'); hircus) (Model 'OVI');) (Model 'EQU'); animals (including <i>Bison</i> and <i>Bubalus</i> species and d of the families Rhinocerotidae and Elephantidae nimals (including <i>Bison</i> and <i>Bubalus</i> species and id of the families Rhinocerotidae and Elephantidae
This — 1 (((((((((((((certificate is meant for fresh meat, including r (1) domestic boving (2) domestic porcing (3) domestic porcing (3) domestic porcing (4) domestic solipe fresh meat, excluding (4) domestic solipe fresh meat, excluding (5) farmed non-domestic their cross-brees (Model 'RUF'); (6) wild non-domestic their cross-brees (Model 'RUW'); (7) farmed non-domestic their crosses 	minced meat, of: e animals (includ animals (<i>Ovis a</i> , ne animals (<i>Sus</i> minced meat, of rds (<i>Equus caba</i> offal and minced nestic animals o ds), <i>Ovis aries</i> , (stic animals of th ds), <i>Ovis aries</i> , (nestic animals belo	ding <i>Bubalus</i> and <i>Bison</i> species and their ries) or domestic caprine animals (<i>Capra I</i> <i>scrofa</i>) (Model 'POR'); : : <i>Ilus, Equus asinus</i> and their cross-breeds d meat, of: f the order Artiodactyla (excluding bovine a <i>Capra hircus</i> , Suidae and Tayassuidae), an the order Artiodactyla (excluding bovine a <i>Capra hircus</i> , Suidae and Tayassuidae), an	cross-breeds) (Model 'BOV'); hircus) (Model 'OVI');) (Model 'EQU'); animals (including <i>Bison</i> and <i>Bubalus</i> species and d of the families Rhinocerotidae and Elephantidae nimals (including <i>Bison</i> and <i>Bubalus</i> species and d of the families Rhinocerotidae and Elephantidae apiridae families (Model 'SUF'); idae families (Model 'SUW');

COUNTRY

COUNTRY Model TRANSIT/STORA								
II. Health information	II.a. Certificate reference number	II.b.						
Part I:								
 Box reference I.11: Place of origin: name Box reference I.12: Address (and approv or ship chandler shall be included. Box reference I.15: Registration number provided. In case of unloading and reloa Box reference I.19: Use the appropriate Box reference I.20: Indicate total gross w Box reference I.23: For containers or box Box reference I.28: <i>Nature of commodity</i> Box reference I.28: <i>Treatment type</i>: If fro Part II: (1) Keep as appropriate. (2) Date or dates of slaughter. Imports of thi date of authorisation for exportation to th 	(railway wagons or container and lorries), ding, the consignor must inform the BIP of a HS code: 02.01, 02.02, 02.03, 02.04, 02.05 weight and total net weight. kes, the container number and the seal num r: Indicate 'carcass-whole', 'carcass-side', 'c zen, indicate the date of freezing (mm/yy) o s meat shall not be authorised when obtain e Union of the third country, territory or part t	t. ree zone, free warehouse, customs warehouse flight number (aircraft) or name (ship) is to be entry into the Union. J. 02.06, 02.08.90, 02.09, 05.04 or 15.02. aber (if applicable) should be included. arcass-quarters', 'cuts', or 'minced meat'.						
Official veterinarian								
Name (in capital letters):	Qualificati	on and title:						
Date:	Signature	:						
Stamp:								

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)
(¹) Suspended from 5 May 2010.		

▼<u>C1</u>

PART 2

Tables of animals and the corresponding model veterinary certificates

Table 1						
'QUE':	QUE': Model of veterinary certificate for consignments of queen bees and queen bumble bees (<i>Apis mellifera and Bombus</i> spp.),					
'BEE':	Model of veterinary certif	ficate for consignments of colonie	s of bumble bees (Bombus spp.)			
Order		Family	Genera/species			
Hymeno	ptera	Apidae	Apis mellifera, Bombus spp.			

	~~		el QUE			Vatarinary contificate to E
		UNTRY	LO Cartifian		number	Veterinary certificate to El
	1.1.	Consignor	I.2. Certifica	ite reference	number	l.2.a.
		Name	I.3. Central Competent Authority			
		Address	I.4. Local Co	ompetent Au	thority	
		Tel. No				
ent	1.5.	Consignee	l.6.			
gnm		Name				
onsi		Address				
ор р		Postal code				
tche		Tel. No				
Part I: Details of dispatched consignment	1.7.	Country ISO I.8. Region Code of origin code of origin	I.9. Country destinat		SO ode	I.10. Region of Code destination
ils o	I.11.	. Place of origin	l.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 o	departure	tii	me of departure
		Address Approval number				
	I.15	. Means of transport Aeroplane Ship Railway wagon	I.16. Entry BIP in EU			
		Road vehicle		ודרכ		
		Identification: Documentary references:	1.17. No(s) of CITES			
	l.18	. Description of commodity		I.19. Comm	odity coo	de (HS code) 01.06.90
			L		1.20. Q	uantity
	I.21				1.22. N	umber of packages
	1.23	B. Identification of container/seal number			1.24.	
-	1.25	5. Commodities certified for: Breeding				
	1.26	D.	I.27. For impo	ort or admiss	ion into E	:U
ł	1.28	B. Identification of the commodities	1			
		Species Identif	ication tem			Identification number
		(Scientific name) sys	aem			number

	COUNT	RY			Model QU				
	П.	Health	information	II.a. Certificate reference number	II.b.				
	II.1.	II.1. Animal Health attestation:							
		I, 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 beet <i>turnida</i>) and the Tropilaelaps mite (<i>Tropilaelaps</i> spp.) are notifiable diseases/pests. II.1.2 they: 								
			(a) come from a breedin	g apiary, which is supervised and control	lled by the competent authority;				
			and where no such certificate. Where an kilometres have beer	occurrence has taken place within at le outbreak of American foulbrood has occ	ociated with an occurrence of American foulbrood, aast 30 days prior to the issuance of the present curred previously, all hives within a radius of three I all infected hives burned or treated and inspected ays following the last recorded case:				
			have been tested in t		f bumble bees) from which samples of the comb s laid down in the OIE Manual of Diagnostic Tests				
					t to any restrictions associated with the occurrence and where these infestations are absent;				
					bumble bees), which were inspected immediately sease including infestations affecting bees;				
			.,		ees and packaging do not contain the small hive estations, in particular <i>Tropilaelaps</i> spp., affecting				
		II.1.3		combs, and all precautions have been ta	nd food are new and have not been in contact with ken to prevent contamination with agents causing				
	Notes								
	Part I:								
		reference) attenda		es (<i>Apis mellifera and Bombus</i> spp.). Eac	ch queen bee may be accompanied by a maximum				
	Part II:								
	(') 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): Qualification and title:							
		Date:		Signat	ure:				
		Stamp:							
L									

		del BEE		
	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			
€	I.5. Consignee	1.6.		
шu	Name			
nsig	Address			
d co	Postal code			
che	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 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 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 I.17. No(s) of CITES		
	Road vehicle D Other			
	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			
	l.26.	I.27. For import or admission into EU		
	I.28. Identification of the commodities	1		
		tification Identification ystem number		

COUN	ITRY		Model BE
11.	Health information	II.a. Certificate reference number	II.b.
II.1.	Animal Health attestation:		
	I, the undersigned, hereby ce	rtify that:	
	II.1.1		
		(Bombus spp.) referred to in Part I of this certificate in a recognised establishment which is supervised	
		nt referred to in Part I of this certificate was insp d breeding stock show no clinical signs or suspicio	
	broodstock and p	mport into the Union have undergone detailed e backaging do not contain the small hive beetle (Ae rticular Tropilaelaps spp., affecting bees;	
		, containers, accompanying products and food a ood-combs, and all precautions have been taken to ons of bees.	
Notes			
Part I:			
	ox reference I.20: Number of cont mble bees.	ainers of bumble bees (<i>Bombus</i> spp.), each con	taining a colony of a maximum of 200 adult
Officia	l veterinarian /Official inspector		
	Name (in capital letters):	Qualification	n 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 (¹) 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.