This document is meant purely as a documentation tool and the institutions do not assume any liability for its contents

►<u>B</u>

▶<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)

Amended by:

Official Journal

		No	page	date
► <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

Corrected by:

- ▶<u>C1</u> Corrigendum, OJ L 146, 11.6.2010, p. 1 (206/2010)
- ► <u>C2</u> Corrigendum, OJ L 49, 24.2.2011, p. 53 (144/2011)
- ► <u>C3</u> Corrigendum, OJ L 63, 10.3.2011, p. 28 (144/2011)

COMMISSION REGULATION (EU) No 206/2010

of 12 March 2010

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

(Text with EEA relevance)

THE EUROPEAN COMMISSION,

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

Having regard to Council Directive 92/65/EEC of 13 July 1992 laying down animal health requirements governing trade in and imports into the Community of animals, semen, ova and embryos not subject to animal health requirements laid down in specific Community rules referred to in Annex A (I) to Directive 90/425/EEC (1), and in particular Articles 17(2)(b) and 17(3)(a), the first subparagraph of Article 17(3)(c), the fourth indent of Article 18(1) and Article 19 thereof,

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

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

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

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

▼B **V**C1

⁽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.

▼<u>M8</u>

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.

▼<u>C1</u>

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 $\left(*\right)$

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 North-easterly to a point 119° longitude, 50°45' latitude Southerly to a point on the Canada/United States border 118°15' 	BOV-X, OVI-X, OVI-Y RUM (**)	А	IVb IX V
CH – Switzerland	CH-0	longitude, 49° latitude Whole country	(***)		
			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	B	
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

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	
	RU-0	Whole country				
RU – Russia	RU-1	Whole country except the region of Kaliningrad				
	RU-2	Region of Kaliningrad	BOV-X-TRANSIT-RU		X	

(*) 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.

- **'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.
- **'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.

^(*) Delete country as applicable.

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

⁽¹⁾ OJ 121, 29.7.1964, p. 1977/64.

⁽²⁾ OJ L 46, 19.2.1991, p. 19.

- **'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.
'POR-X':	Model of veterinary certificate for domestic porcine animals (Sus scrofa) intended for breeding and/or production after importation.
'POR-Y':	Model of veterinary certificate for domestic porcine animals (Sus scrofa) intended for immediate slaughter after importation.
'RUM':	Model of veterinary certificate for animals of the order Artiodactyla (excluding bovine animals (including Bubalus and Bison species and their cross-breeds), Ovis aries, Capra hircus, Suidae and Tayassuidae), and of the families Rhinocerotidae and Elephantidae.
'SUI':	Model of veterinary certificate for non-domestic Suidae, 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.

'A':	guarantees regarding Bluetongue and Epizootic-haem orrhagic-disease tests on animals certified according to the model of veterinary certificates BOV-X (poin II.2.8 B), OVI-X (point II.2.6 D) and RUM (poin
	II.2.6).
'B':	guarantees regarding Swine-vesicular-disease and Classical-swine-fever tests on animals certificat according to the model of veterinary certificate POR-X (point II.2.4 B) and SUI (point II.2.4 B).
'C':	guarantees regarding Brucellosis test on animal certified according to the model of veterinary certificates POR-X (point II.2.4 C) and SUI (point

Model BOV-X

col	INTR	1				Veterinary ce	ertificate to EU	
	1.1.	Consignor		I.2. Certificat	e reference No	l.2.a.		
		Name Address	I.3. Central competent authority					
		Tel.				-		
Ħ				I.4. Local col	mpetent authority			
Part I: Details of dispatched consignment	1.5.	Consignee		I.6.				
Isign		Name Address						
cor								
chec		Postal code Tel.						
spat	1.7.		Code	I.9. Country	of ISO code	e I.10. Region of	Code	
of di	1.7.		Joue	destinatio		destination	Oode	
ils c								
Deta	1.11.	Place of origin		l.12.				
÷		Name Approval number						
Ра		Address						
	113	Place of loading		I.14. Date of departure				
					opurturo			
		Address Approval number						
	l.15.	Means of transport		I.16. Entry BIF	P in EU			
		Aeroplane 🗌 Ship 🗌 Railway wagon 🗌						
		Road vehicle Other I	F	l.17.				
		Documentary references						
	I.18.	Description of commodity	L		I.19. Commodity	code (HS code)		
				l	01.02			
						I.20. Quantity		
	1.21.					I.22. Number of packag	jes	
	1.23.	Seal/Container No				1.24.		
	1.05							
	1.25.	Commodities certified for:	_					
		Breeding	F	attening 🔲				
	1.26.			I.27. For impo	rt or admission inf	to EU		
	1.28.	Identification of the commodities	I					
		Species Breed Ide	entificatior	n k	dentification	Age	Sex	
			system		number	-		

• 1010

col	JNTRY						Model BOV-X		
	11.	Health	n information			II.a. Certificate reference number	II.b.		
	11.1.	Public	Health Attesta	tion	·				
		I, the	undersigned offic	cial v	eterinarian, hereby certify, that th	ne animals described in this certificate	и:		
Part II: Certification		II.1.1. come from holdings which have been free from any official prohibition on health grounds, for the past 42 days in the c brucellosis, for the past 30 days in the case of anthrax and for the past six months in the case of rabies, and, have not b contact with animals from holdings which did not satisfy these conditions;							
t II: Ce		II.1.2.	have not receiv	ed:					
Par			— any stilbene	or t	hyrostatic substances,				
					ogenic, gestagenic or β- agonist rective 96/22/EC);	substances for purposes other than t	herapeutic or zootechnic treatment		
	-	II.1.3.	with regard to b	ovin	e spongiform encephalopathy (B	SE):			
			(¹) (²) either	[(a)		a permanent identification system ena 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 iding 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 ena Id are not exposed bovine animals a Regulation (EC) No 999/2001;			
				(b)	meal and greaves derived from	date from which the ban on the feedin ruminants had been effectively enforc n after the date of the feed ban.]			
			(¹) (⁴) or	[(a)		a permanent identification system ena 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 the reaves derived from ruminants had be digenous case if born after the date of	en effectively enforced or after the		
	II.2.	Anima	al Health attesta	ation	:				
		I, the	undersigned offic	ial v	eterinarian, hereby certify, that th	ne animals described above meet the	following requirements:		
		II.2.1.	they come from	the	territory with code:	(⁵) which, at the date o	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 ter that date, and authorised to exp No/, of	ort these animals by Commission		
				(b)		m rinderpest, Rift valley fever, contagio norrhagic disease, and for six months			
				(c)		s, no vaccination against the diseases i f domestic cloven-hoofed animals vac			
			(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	orn bluetongue, and the animals have y for bluetongue and epizootic haemoi taken at the beginning of the isolation (dd/mm/yyyy) and on tithin 10 days before export;]	rrhagic disease, carried out on two n/quarantine period and at least 28		

COUNTRY					Model BOV-X			
П.	Health	information		II.a. Certificate reference number	II.b.			
		(¹) or	inactivated vaccine, at least 60 serotype/s (inse demonstrated through a surve holding(s) of origin described u	hths from bluetongue, and the anima o days before the date of dispatch to <i>int 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.		ained in the territory described under p without contact with imported cloven-	point 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 ren reference I.11.		lays before dispatch in the holding(s) of origin described under box			
		(a) in and around which, in an area with a 150 km radius, there has been no case/outbreak of epizootic haemorrhag during the previous 60 days,						
		rinderpest,		n radius, there has been no case/ou bus bovine pleuropneumonia, lumpy sk				
	II.2.4.		nimals to be killed under a national pr eases referred to under point II.2.1,(a	ogramme for the eradication of diseas)) and (b);	es, nor have they been vaccinated			
	II.2.5.		m herds that are not restricted und I enzootic bovine leukosis;	er the national legislation pertaining	to the eradication of tuberculosis,			
	II.2.6.	they come fror	n 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) or	[have been subjected to an intrade 30 days before dispatch to the Unic	ermal 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	been 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	[II.2.8.			or the control of enzootic bovine leukos / test of this disease during the past t				
(¹) or	[11.2.8.	they come fror	n herds recognised as officially enzoo	otic-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 individu samples taken within the past 30 da	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 assembl described under point II.2.1,]	y centre described under box referen	ce I.13 situated within the territory			

			Il o Cortificato referer e much	Model BOV-				
П.	Health	information	II.a. Certificate reference number	II.b.				
		and, until dispatched to the Union:						
	 (a) they did not come in contact with other cloven-hoofed animals not complying with the health requirements as described in 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 been a case/outbreak of any of the diseases referred to in point II.2.1.;							
	II.2.10. any transport vehicles or containers in which they were loaded were cleaned and disinfected before loading with an official authorised disinfectant;							
	II.2.11.	they were examined by an official veterinarian withi	n 24 hours of loading and showed no	o clinical sign of disease;				
	II.2.12.	they have been loaded for dispatch to the Union or under box reference I.15 above that were cleaned a so constructed that faeces, urine, litter or fodder or	ind disinfected before loading with an	officially authorised disinfectant an				
II.3.	Anima	l transport attestation						
	loading	indersigned official veterinarian, hereby certify, that to g in accordance with the relevant provisions of Regul re fit for the intended transport.						
(¹) (¹¹) [II.4.	Specif	ic requirements						
	II.4.1.	According to official information, no clinical or pa recorded in the holding(s) of origin referred to in bo						
	II.4.2.	the animals referred to in box reference I.28.:						
		 (a) have been isolated in accommodation approve dispatch for export, 	d by the competent authority for the	e last 30 days immediately prior to				
		(b) have been subjected to a serological test for IB results, and all animals in isolation have also give		er entry into isolation, with negative				
		(c) have not been vaccinated against IBR.]						
Notes								
This certifica production.	ate is m	eant for domestic bovine animals (including <i>Bubalus</i>	and Bison species and their cross-b	reeds) intended for breeding and/o				
		animals must be conveyed without delay to the holdi ment outside the holding, except in the case of a di		ain for a minimum period of 30 day				
Part I:								
— Box refe	rence I.	8.: Provide the code of territory as appearing in Part	1 of Annex I to Regulation (EU) No	206/2010.				
 Box reference I.13.: The assembly 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. 								
— 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 refe	rence I.	23.: For containers or boxes, the container number a	and the seal number (if applicable) sh	ould be included.				
— Box refe	rence I.	28.: Identification system: The animals must bear:						
	dividual conder)	number which permits tracing of their premises of c	rigin. Specify the identification system	n (such as tag, tattoos, brand, chip				
	or toa t	hat includes the ISO code of the exporting country	The individual number must permit	traging of their promises of origin				

COUNTRY Model BC									
11.	Health information	II.a. Certificate reference number	II.b.						
	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, crossbreed.								
Pa	Part II:								
(1)	Keep as appropriate.								
(2)	Only if the animals were born and continuously reared in a country No 999/2001 as a country or region posing a negligible BSE risk ar								
(3)	Only if the country or region of origin is categorised in accordance posing a controlled BSE risk and is listed as such in Decision 2007		lo 999/2001 as a country or region						
(4)	Only if the country or region of origin has not been categorised in ac categorised as a country or region with undetermined BSE risk and								
(5)	Code of the territory as it appears in Part 1 of Annex I to Regulatio	n (EU) No 206/2010.							
(6)	Officially tuberculosis/brucellosis-free regions and herds as laid dow regions and herds as laid down in Chapter I of Annex D to Directive		; and enzootic-bovine-leukosis-free						
(^{6a})	Only for officially enzootic-bovine-leukosis-free herds recognised as Directive 64/432/EEC for the purpose of exports to the EU of live ar column 6 of Part 1 of Annex I to Regulation (EU) No 206/2010, app	nimals according to the model certifica	ate BOV-X from the territory that, in						
(7)	Only for a territory that, in column 6 of Part 1 of Annex I to Regulatic "III", as regards brucellosis, and/or "IVa" as regards enzootic bovine		e entry "II", as regards tuberculosis,						
(8)	Tests carried out in accordance with the protocols that, for the dise No 206/2010.	ease concerned, are described in Pa	rt 6 of Annex I to Regulation (EU)						
(9)	Supplementary guarantees to be provided when required in column entry "A".	n 5 "SG" of Part 1 of Annex I to Regi	ulation (EU) No 206/2010, with the						
	Tests for bluetongue and for epizootic haemorrhagic disease in acc	cordance with Part 6 of Annex I to Re	gulation (EU) No 206/2010.						
(10)	Date of loading. Imports of these animals shall not be allowed whi 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, o	r during a period where restrictive						
(11)	When required by the EU Member State of destination or Switzerlan Agreement between the Community and the Swiss Confederation or								
(12)	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.).						
Off	Official veterinarian								
	Name (in capital letters):	Qualification and title:							
	Date:	Signature:							
	Stamp:								
1									

Model BOV-Y

col	NTR	<i>i</i>	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				
dispatched consignment	1.5.	Consignee	1.6.				
signı		Name Address					
con		Postal code					
ched		Tel.					
spate	1.7.	Country of origin ISO code I.8. Region of origin Code	I.9. Country of ISO code I.10. Region of Code				
of di			destination destination				
ails	1 1 1	Place of origin	1.12.				
Part I: Details	1.11.		1.12.				
art I		Name Approval number Address					
4							
	1.13.	Place of loading	I.14. Date of departure				
		Address Approval number					
	l.15.	Means of transport	I.16. Entry BIP in EU				
		Aeroplane Ship Railway wagon					
		Road vehicle Other					
		Identification	I.17. I.19. Commodity code (HS code) 01.02				
		Documentary references					
	l.18.	Description of commodity					
			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				
	1.20.						
	1.28.	Identification of the commodities					
		Species Breed Identification system (scientific name)	Identification number Age Sex				

cou	NTRY						Model BOV-Y
	II.	Health	information			II.a. Certificate reference number	ll.b.
	II.1.	Public	Health Attestation	ı			
		l, the u	ndersigned official	veter	narian, hereby certify, that the ar	nimals described in this certificate:	
Part II: Certification		II.1.1.	brucellosis, for the	last		ficial prohibition on health grounds, f r the last six months in the case of ra conditions;	
II: Cei		II.1.2.	have not received	:			
Part			— any stilbene o	r thyr	ostatic substances,		
			 oestrogenic, and defined in Direct 			bstances for purposes other than ther	rapeutic or zootechnic treatment (as
		II.1.3.	with regard to boy	/ine s	pongiform encephalopathy (BSE)		
			(¹) (²) <i>either</i>	[(a)		permanent identification system enab not exposed bovine animals as descrit C) No 999/2001;	
				(b)	from which the ban on the fee	us cases in the country concerned, the ding of ruminants with meat-and-bor enforced or after the date of birth on	ne meal and greaves derived from
			(¹) (³) or	[(a)		permanent identification system enab not exposed bovine animals as desc (EC) No 999/2001;	
				(b)	and-bone meal and greaves de	ne date from which the ban on the rived from ruminants had been effect case if born after the date of the fee	tively enforced or after the date of
			(¹) (⁴) or	[(a)		permanent identification system enab not exposed bovine animals as desc i (EC) No 999/2001;	
				(b)	with meat-and-bone meal and gr	wo years after the date from which ti reaves derived from ruminants had be igenous case if born after the date o	een effectively enforced or after the
	II.2.	Animal	Health Attestatio	n			
		I, the u	ndersigned official	veter	narian, hereby certify, that the ar	nimals described above meet the follo	owing requirements:
		II.2.1.	they come from the	ne tei	ritory with code:	(⁵) which, a	t the date of issuing this certificate:
			(¹) either	[(a)	has been free for 24 months fro	m foot-and-mouth disease]	
			(¹) or	[(a)	had cases/outbreaks after that	foot-and-mouth disease since at date, and authorised to export o/, of	t these animals by Commission
				(b)		n rinderpest, Rift valley fever, contagio norrhagic disease, and for six months	
				(c)		no vaccination against the diseases domestic cloven-hoofed animals vac	
			(¹) either	[(d)	has been free for 24 months from	m bluetongue;]	

Health	information			II.a. Certificate	reference number	II.b.	
	(¹) or	,	inactivated vaccine, at I serotype/s demonstrated through a	24 months from bluetor east 60 days before the (<i>insert serotype</i> , surveillance programme (r box reference I.11, and ications of the vaccine;]	e date of dispatch to /s) which are those (⁹) in an area with a	o the Union, present in the 150 km radius	against all bluetor source population around the holdin
II.2.2.				der point II.2.1 since birth ven-hoofed animals for t		ast three mor	nths before dispate
II.2.3.	they have ren	nained sin	ce birth or at least 40 da	ays before dispatch in th	e holding(s) describ	ed under box	reference I.11:
			h, in an area with a 150 60 days, and	km radius, there has be	en no case/outbreał	of epizootic	haemorrhagic dise
		y fever, bl		n radius, there has been ovine pleuropneumonia, l			
II.2.4.			be killed under a natio ferred to in point II.2.1(a	nal programme for the e) and (b);	eradication of diseas	es, nor have	they been vaccina
II.2.5.	they come fro	om herds:					
	(a) included in an official system for the control of enzootic bovine leukosis, and						
	(b) that are not restricted under the national legislation regarding eradication of tuberculosis and brucellosis, and						
	(c) recognise	d as offici	ally tuberculosis free; (⁶)				
II.2.6.	they have not	t been vac	cinated against brucellos	sis and they:			
	(¹) either	[come fr	om herds which are reco	ognised as officially bruch	ellosis free;](⁶)		
	(¹) or	[are cas	trated males of any age;]			
II.2.7.	they are individually marked on at least two places on their hindquarters as to show that they are exclusively inter immediate slaughter; (⁷)						clusively intended
II.2.8.	8. they are/were (¹) dispatched from their holding(s) of origin, without				ng through any mark	ket:	
	(¹) either	[directly	to the Union,]				
	(¹) or		officially authorised asse d under point II.2.1]	embly centre described	under box referenc	e I.13 situat	ed within the terri
	and, until dispatched to the Union:						
	(a) they did n certificate		n contact with other clove	n-hoofed animals not cor	mplying with the heal	th requiremer	nts as described in
	(b) they were not at any place where, or around which within a 10 km radius, during the previous 30 days there has case/outbreak of any of the diseases referred to in point II.2.1;						
II.2.9.	any transport authorised dis		or containers in which th	ney were loaded were c	leaned and disinfect	ed before lo	ading with an offic
II.2.10.	they were exa	amined by	an official veterinarian v	vithin 24 hours of loading	g and showed no cli	nical sign of	disease;
II.2.11.	under box ref	erence I.1	for dispatch to the Unio 5 above that were clean s, urine, litter or fodder	n on ed and disinfected before	e loading with an offi	cially authoris	ed disinfectant and

II.	Health information	II.a. Certificate reference numbe	r II.b.			
11.3.	Animal transport attestation					
	I, the undersigned official veterinarian, hereby certify, th in accordance with the relevant provisions of Regulatio the intended transport.					
Note	S					
This	certificate is meant for live bovine animals (including Bub	palus and Bison species and their cross-bre	eds) intended for immediate slaughter			
After	importation the animals must be conveyed without delay	/ to the slaughterhouse of destination to be	e slaughtered within five working days			
Part	l:					
— В	ox reference I.8: Provide the code of territory as appearin	ng in Part 1 of Annex I to Regulation (EU) N	No 206/2010.			
	ox reference I.13: The assembly centre, if any, must fulfil t o 206/2010.	the conditions for its approval, as laid down	in Part 5 of Annex I to Regulation (EU			
	ox reference I.15: Registration number (railway wagons or ase of unloading and reloading, the consignor must inform		ft) or name (ship) is to be provided. I			
— В	ox reference I.23: For containers or boxes, the container	number and the seal number (if applicable)	should be included.			
— В	ox reference I.28: Identification system: the animals must	bear:				
_	 An individual number which permits tracing of their prer transponder). 	nises of origin. Specify the identification sys	stem (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 premis						
S	pecies: Select amongst "Bos", "Bison" and "Bubalus" as a	appropriate.				
Ą	ge: Date of birth (dd/mm/yy).					
S	ex (M = male, F = female, C = castrated).					
Part	И:					
(¹) K	eep as appropriate.					
	only if the animals were born and continuously reared in a to 999/2001 as a country or region posing a negligible BS					
	only if the country or region of origin is categorised in acc osing a controlled BSE risk and is listed as such in Decis		C) No 999/2001 as a country or regio			
	only if the country or region of origin has not been catego ategorised as a country or region with undetermined BSE					
(⁵) C	ode of the territory as it appears in Part 1 of Annex I to I	Regulation (EU) No 206/2010.				
(⁶) O	fficially tuberculosis/brucellosis free regions and herds as	laid down in Annex A to Directive 64/432/	EEC.			
(⁷) T	his mark shall take the form of "L" having 13 cm in the le pplied using the technique known as "freeze-branding".	ft side and 7 cm in the bottom side with 1	cm of strength in both lines. It shall b			

COUNTRY Model E							
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 1.7 and 1.8, or during a period where restrictive measures have been adopted by the Union against imports of these animals from this third country, territory or part thereof.							
(⁹) Surveillance programme as laid down in Annex I to Commission reg	ulation (EC) No 1266/2007 (OJ L 283	, 27.10.2007, p. 37.).					
Official veterinarian							
Name (in capital letters):	Name (in capital letters): Qualification and title:						
Date:	Signature:						
Stamp:							

Model BOV-X-TRANSIT-RU

cοι	INTR	(Veterinary certificate to E				
	I.1.	Consignor Name	I.2. Certificate reference No I.2.a.				
		Address	I.3. Central competent authority				
		Tel.	I.4. Local competent authority				
Part I: Details of dispatched consignment	1.5.	Consignee Name	I.6. Person responsible for the load in EU Name				
		Address	Address Postal code Tel.				
ched c		Postal code Tel.					
lispato	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				
° of		Russia Kaliningrad	Russia				
etails	1.11.	Place of origin	1.12.				
art I: D		Name Address					
ã		Postal code					
	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 Other	1.17.				
		Identification					
		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:					
		Breeding	Fattening				
	1.26.	For transit through EU to third country	1.27.				
		Third country Russian Federation ISO code RU					
	1.28.	Identification of the commodities					
		Species Breed Identification (scientific name)	n system Identification number Age Sex				

COUNT	RY			Model BOV-X-TRANSIT-RU					
II.	Health info	rmation	II.a. Certificate reference number	II.b.					
II.1.	Animal He	ealth attestation							
	I, the unde	ersigned official veterinarian, hereby certify, that the a	nimals described in Part I meet the fo	llowing requirements:					
	II.1.1.	they come from the territory with code: RU-2 $(^{2})$ whi	ch, at the date of issuing this certifica	te:					
	(¹) either	[(a) has been free for 24 months from foot-and-mou	ith disease]						
	(¹) or	[(a) has been considered free from foot-and-mouth cases/outbreaks after that date, and authorised No	to export these animals by Commis-						
		(b) has been free for 12 months from rinderpest, F and epizootic haemorrhagic disease, and for 6		uropneumonia, lumpy skin disease					
		(c) where during the last 12 months, no vaccination and imports of domestic cloven-hoofed animals							
	(¹) either	[(d) has been free for 24 months from bluetongue;]							
	(¹) or	[(d) has not been free for 24 months from bluetong at least 60 days before the date of the movem are those present in the source population as de radius around the holding(s) of origin described period of time guaranteed in the specifications	ent, against all bluetongue serotype/s monstrated through a surveillance prog d under box reference I.11, and the a	(insert serotype/s) which pramme (⁴) in an area with a 150 km					
	II.1.2. they have remained in the territory described under point II.1.1. since birth, or for at least the last six months befor dispatch via the European Union and without contact with imported cloven-hoofed animals for the last 30 days;								
	II.1.3.	they have remained since birth or at least 40 days b reference $\ensuremath{I.11.}$	efore the date of dispatch in the holdi	ng(s) of origin described under box					
		 (a) in and around which, in an area with a 150 km ra during the previous 60 days, 	dius, there has been no case/outbreal	of epizootic haemorrhagic disease					
		(b) in and around which, in an area with a 10 kn rinderpest, Rift valley fever, bluetongue, contagic during the previous 40 days;							
	II.1.4.	they are not animals to be killed under a national pragainst the diseases referred to under point II.1.1.,		es, nor have they been vaccinated					
		 (a) they did not come in contact with other cloven-he this certificate, 	pofed animals not complying with the h	nealth requirements as described in					
		(b) they were not at any place where, or around wh case/outbreak of any of the diseases referred to		previous 30 days there has been a					
	II.1.5.	any transport vehicles or containers in which they w authorised disinfectant;	ere loaded were cleaned and disinfec	ted before loading with an officially					
	II.1.6 <i>.</i>	they were examined by an official veterinarian within	ו 24 hours of loading and showed no	clinical sign of disease;					
	II.1.7 <i>.</i>	they have been loaded for dispatch to the Russian the means of transport described under box referen officially authorised disinfectant and so constructed to container during transportation.	nce I.15. above that were cleaned and	disinfected before loading with an					
	II.1.8 <i>.</i>	The consignment is intended to leave the European	n Union at the designated Border Ins	pection Post Medininkai, Lithuania.					

COUNTRY Model BC										
II.		Health information	II.a. Certificate reference number	II.b.						
11.3	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.									
N	Notes									
	This certificate is meant for transit through the European Union of domestic bovine animals (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 the Russia.									
Pa	art I:									
-	- Box	reference I.8.: Provide the code of territory as appearing in Part	1 of Annex I to Commission Regulat	ion (EU) No 206/2010.						
-		reference I.13.: The assembly centre, if any, must fulfil the cond ulation (EU) No 206/2010.	litions for its approval, as laid down i	n Part 5 of Annex I to Commission						
-		reference I.15.: Registration number of road vehicle, is to be pro- ler Inspection Post of entry into the Union.	vided. In case an emergency, the cor	nsignor must immediately inform the						
-	- Вох	reference I.23.: For containers or boxes, the container number a	and the seal number (if applicable) mi	ust be included.						
-	- Box	reference I.28.: Identification system: The animals must bear:								
		An individual number which permits tracing of their premises of o ransponder).	rigin. Specify the identification system	n (such as tag, tattoos, brand, chip,						
	— A	An ear tag that includes the ISO code of the exporting country.	The individual number must permit	tracing of their premises of origin.						
-	- Box	reference I.28.: Species: Select amongst "Bos", "Bison" and "Bul	balus" as appropriate.							
-	- Box	reference I.28.: Age: Date of birth (dd/mm/yy).								
-	- Box	reference I.28.: Sex (M = male, F = female, C = castrated).								
-	- Box	reference I.28.: Breed: select purebred, crossbreed.								
Pa	art II:									
0) Kee	p as appropriate.								
(2)) Coc	le of the territory as it appears in Part 1 of Annex I to Commission	on Regulation (EU) No 206/2010.							
(3)	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)) Sur	veillance programme as laid down in Annex I to Commission Reg	gulation (EC) No 1266/2007.							
0	fficial	veterinarian								
	Nar	ne (in capital letters):	Qualifica	tion and title:						
	Dat	e:	Signatur	e:						
	Sta	np:								

Model OVI-X

cou	INTR	1		Veterinary certificate to EU			
	l.1.	Consignor Name	I.2. Certificate reference No	l.2.a.			
		Address	I.3. Central competent authority				
-		Tel.	I.4. Local competent authority				
I: Details of dispatched consignment	1.5.	Consignee Name Address	1.6.				
atched co		Postal code Tel.					
s of disp	1.7.	Country of origin ISO code I.8. Region of origin Code	I.9. Country of ISO code destination	e I.10. Region of Code destination			
Detail	l.11.	Place of origin	l.12.				
Part I: I		Name Approval number Address					
	l.13.	Place of loading	I.14. Date of departure				
		Address Approval number					
	l.15.	Means of transport	I.16. Entry BIP in EU				
		Aeroplane Ship Railway wagon Road vehicle Other					
		Identification Documentary references	l.17.				
	l.18.	Description of commodity	I.19. Commodity code (HS code)				
				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.		I.27. For import or admission into EU				
	1.28.	Identification of the commodities					
		Species Breed Identificati (scientific name) system	on Identification number	Age Sex			

οι	JNTRY						Model OVI				
	11.	Health i	nformation	I		II.a. Certificate reference number	II.b.				
	II.1.	Public	Health At	alth Attestation							
		I, the ur	ndersignec	l offici	al veterinarian, hereby certify, that th	e animals described in this certificate:					
Fart II: Ceruildauon		II.1.1. come from holdings which have been free from any official prohibition on health grounds, for the la brucellosis, for the last 30 days in the case of anthrax, for the last six months in the case of rabie contact with animals from holdings which did not satisfy these conditions;									
دe ۱		II.1.2. have not received:									
			— any	stilbe	ne or thyrostatic substances,						
					ic, androgenic, gestagenic or β- agon d in Directive 96/22/EC).	ist substances for purposes other thar	therapeutic or zootechnic treatment				
	11.2.	Animal	Health at	testa	ion						
		I, the ur	, the undersigned official veterinarian, hereby certify, that the animals described above meet the following requirements:								
		II.2.1.	they cor	me fro	m the territory with code:	(1) which, at the date of issu	ing this certificate:				
			(²) eithei	r [(a)	has been free for 24 months from f	oot-and-mouth disease]					
			(²) or	[(a)	having had cases/outbreaks after the	-and-mouth disease since hat date, and authorised to export th of (dd/mm/yyyy),]	ese animals by Commission Imple				
				(b)		nderpest, Rift valley fever, peste des p sumonia, and epizootic haemorrhagic					
				(c)		vaccination against the diseases ment stic cloven-hoofed animals vaccinate					
			(²) eithei	r [(d)	has been free for 24 months from b	bluetongue;]					
			(²) (⁹) or	[(d)	the detection of antibody for blueton samples of blood taken at the be	uetongue, and the animals have reacte gue and epizootic haemorrhagic disea eginning of the isolation/quarantine p y) 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 press programme (1^{1}) in an area with a	om bluetongue, and the animals have date of dispatch to the Union, againsi 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 n-hoofed animals for the last 30 days;					
		II.2.3.	they hav	ve rer	nained since birth or at least 40 da	ys in the holding(s) described under	box reference I.11 before dispatcl				
					round which, in an area with a 150 during the previous 60 days, and	km radius, there has been no case/	outbreak of epizootic haemorrhag				
			rind	lerpes		km radius, there has been no case/c ste des petits ruminants, sheep pox ng the provious 40 days;					

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

COUNTRY	Y			Model OVI-X				
11.	Health in	formation	II.a. Certificate reference number	II.b.				
(2	²) [II.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	ne 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 with	h the guarantees provided for in the				
(¹) eithe	ər [II.2.8.2.	are animals intended for production born in and con diagnosed;]	ntinuously reared on holdings in which	a case of scrapie has never been				
(²) (⁸) o	r [II.2.8.2.	they shall have been kept continuously since birth or following requirements for at least three years:	$^{\prime}$ for the last three years on a holding c	or holdings which have satisfied the				
		- they are subject to regular official veterinary che	cks,					
		- the animals are identified in conformity with Unic	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	[II.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.	they are/were (1) dispatched from their holding(s) of origin, without passing through any market,						
		(²) <i>either</i> [directly to the Union,]	²) <i>either</i> [directly to the Union,]					
		(²) or [to the officially authorised assembly ca described under point II.2.1.]	entre described under box reference	ə 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 h	nealth requirements as described in				
		(b) they were not at any place where, or around wh case/outbreak of any of the diseases referred to		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 disinfec	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 on described under box reference 1.15 above that we disinfectant and so constructed that faeces, urine, during transportation.	ere cleaned and disinfected before lo	bading with an officially authorised				

11.	Health information	II.a. Certificate reference number	II.b.								
II.3.	Animal transport attestation										
	I, the undersigned official veterinarian, hereby certify, that the animals described above have been treated before and at the time of loading in accordance with the relevant provisions of Regulation (EC) No 1/2005, in particular as regards watering and feeding, and they are fit for the intended transport.										
Notes											
This cei producti	rtificate is meant for live domestic ovine animals (ion.	(Ovis aries) and domestic caprine animals (Cap	ora hircus) intended for breeding or								
	portation the animals must be conveyed without delay urther movement outside the holding, except in the		nain for a minimum period of 30 days								
Part I:											
— Box	reference I.8: Provide the code of territory as appea	aring in Part 1 of Annex I to Regulation (EU) No	206/2010.								
	reference I.13: The assembly centre, if any, must ful 206/2010.	fil the conditions for its approval, as laid down in	Part 5 of Annex I to Regulation (EU)								
	reference I.15: Registration number (railway wagons of unloading and reloading, the consignor must info		or name (ship) is to be provided. In								
— Box	reference I.19: Use the appropriate HS code: 01.04	.10 or 01.04.20.									
— Box	reference I.23: For containers or boxes, the container	er number and the seal number (if applicable) sl	hould be included.								
— Box	reference 1.28: Identification system: The animals m	ust bear:									
	An individual number which permits tracing of their pransponder) and the anatomic place used in the anim		m (such as tag, tattoos, brand, chip								
— A	An ear tag that includes the ISO code of the expor	rting country. The individual number must permi	it tracing of their premises of origin								
Spec	cies: Select amongst "Ovis aries" and "Capra hircus"	" as appropriate.									
Age:	: (months).										
Sex	(M = male, F = female, C = castrated).										
Part II:											
(¹) Coo	de of the territory as it appears in Part 1 of Annex I	to Regulation (EU) No 206/2010.									
(²) Kee	ep as appropriate.										
(³) Only	y for a territory appearing with the entry "V" in colun	nn 6 of Part 1 of Annex I to Regulation (EU) No	206/2010.								
(⁴) The	e representative number of animals to be tested for t	brucellosis must, for each holding, consist of:									
—	all non-castrated male animals, which have not beer	n vaccinated against brucellosis, over six monthe	s old,								
—	all non-castrated male animals, which have been va	accinated against brucellosis, over 18 months old	I,								
—	all animals brought onto the holding since the previo	ous tests, and									
— :	25% of females which are sexually mature, within a	minimum of 50 females.									
	s must be completed when the destination is a Mer 52/EEC.	mber State or part of a Member State laid down	n in one of the Annexes of Decision								

cou	NTRY		Model OVI-X						
11.	Health information	II.a. Certificate reference number	II.b.						
(6)	In accordance with Part 6 of Annex I to Regulation (EU) No 206/2								
	Where more than one holding of origin is involved the date of the most recent test on each holding must be clearly indicated.								
(7)	Guarantees in relation to a programme of control of scrapie, as requested by the EU Member State of destination, in application of Article 15 and Chapter E of Annex IX to Regulation (EC) No 999/2001.								
(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 will exportation to the Union of the third country, territory or part them measures have been adopted by the Union against imports of the	eof referred to in boxes I.7 and I.8, c	r 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	83, 27.10.2007, p. 37.).						
Offic	ial veterinarian								
	Name (in capital letters):	Qualification and title:							
	Date: Signature:								
	Stamp:								

Model OVI-Y

col	OUNTRY								Veterinary ce	ertificate to EU		
	l.1.	Consignor Name				I.2. Certificate reference No I.2.a.						
		Address Tel.					Central competence	tent authority				
ent							Local compete	nt authority				
gnme	1.5.	Consignee				1.6.						
consi		Name Address										
hed		Postal code										
patc		Tel.				-						
Part I: Details of dispatched consignment	1.7.	Country of origin	ISO code	I.8. Region of origin	Code	1.9.	Country of destination	ISO code	I.10. Region of destination	Code		
Deta	1.11.	Place of origin				I.12.						
Part I:		Name Approval number Address										
	I.13.	B. Place of loading					Date of depart	ure				
		Address Approval number										
	l.15.	5. Means of transport				I.16.	Entry BIP in El	U				
	Aeroplane 🗌 Ship 🗌 Railway wagon 🗌											
		Road vehicle 🗌	Other [l.17.						
		Identification										
	118	Documentary refer				I.19. Commodity code (HS code)						
	1.10.	Description of con	moulty									
						I.20. Quantity						
	1.21.							1.2	2. Number of packag	ges		
	1.23.	Seal/Container No						1.2	4.			
	1.05	Commodition contif	ind for									
	1.20.	Commodities certif										
	1.26.					I.27. For import or admission into EU						
	1.28.	1.28. Identification of the commodities										
		Species (scientific name)	Bree	d Identificat system		lo	lentification num	nber	Age	Sex		

col	JNTRY				Model OVI-Y						
	II.	Health information	วท	II.a. Certificate reference number	II.b.						
	II.1.	Public Health A	Attestation								
		I, the undersigne	undersigned official veterinarian, hereby certify, that the animals described in this certificate:								
Part II: Certification		brucellosi	m holdings which have been free from an s, for the last 30 days in the case of anthra als from holdings which did not satisfy the	x, for the last six months in the case of r							
II: Cer		II.1.2. have not	received:								
Part		— any st	ilbene or thyrostatic substances,								
			genic, androgenic, gestagenic or β- agonis d in Directive 96/22/EC).	t substances for purposes other than the	rapeutic or zootechnic treatment (as						
	II.2.	Animal Health a	attestation								
		I, the undersigne	ed official veterinarian, hereby certify, that	the animals described above meet the	following requirements:						
		II.2.1. they come this certifi	e from the territory with code:		(¹) which, at the date of issuing						
		(²) either	[(a) has been free for 24 months from	foot-and-mouth disease]							
		(²) or		t-and-mouth disease since s after that date, and authorised to ex 	port these animals by Commission						
			 (b) has been free for 12 months from r pox, contagious caprine pleuropneu stomatitis, 	inderpest, Rift valley fever, peste des p monia, and epizootic haemorrhagic dise							
			(c) where during the last 12 months, no carried out and imports of domestic	o vaccination against the diseases ment cloven-hoofed animals vaccinated agair							
		(²) either	[(d) has been free for 24 months from	bluetongue;]							
		(²) or	(<i>insert serotype/s</i>) which are those programme (⁵) in an area with a 15	om bluetongue, and the animals have l e date of dispatch to the Union, agains present in the source population as d i0 km radius around the holding(s) of or the immunity period of time guaranteed	t all bluetongue serotype/s lemonstrated through a surveillance rigin described under box reference						
			remained in the territory described under p and without contact with imported cloven		last three months before dispatch to						
		II.2.3. they have	e remained since birth or at least 40 da	ys before dispatch in the holding(s) d	escribed under box reference I.11:						
			d around which in an area with a 150 km g the previous 60 days, and	radius there has been no case/outbreak	< of epizootic haemorrhagic disease						
		rinder	d around which, in an area with a 10 k rpest, Rift valley fever, bluetongue, peste o ionia and vesicular stomatitis during the p	des petits ruminants, sheep pox and go	utbreak of foot-and-mouth disease, at pox; contagious caprine pleurop-						
			not animals to be killed under a national p ne diseases referred to in point II.2.1(a) an		ses, nor have they been vaccinated						
		II.2.5. they are/v	were (²) dispatched from their holding(s) of	origin, without passing through any ma	arket,						
		(²) either	[directly to the Union]								

COUNTRY Model						
II.	Health ir	nformation	II.a. Certificate reference number	II.b.		
		(²) or [to the officially authorised assemi under point II.2.1,]	bly centre described under box reference I.13 s	ituated within the territory described		
		and, until dispatched to the Union:				
		 (a) they did not come in contact with other this certificate, and 	r cloven-hoofed animals not complying with the	health requirements as described in		
		(b) they were not at any place where, or a case/outbreak of any of the diseases	around which within a 10 km radius, during the referred to in point II.2.1;	previous 30 days there has been a		
	II.2.6.	in respect of scrapie:				
(2) (3)	[II.2.6.1.	or (c) of Chapter A(I) of Annex VIII to Reg	ich benefits, for all or part of its territory, from th ulation (EC) No 999/2001, the animals comply s, as laid down in Article 2 of Regulation (EC)	with the guarantees provided for in		
(²) either	[11.2.6.2.	were born in and continuously reared on I	noldings in which a case of scrapie has never	been diagnosed;]		
(²) or	[II.2.6.2.	are domestic ovine animals of the ARR/AR from a holding where no case of scrapie I	IR prion protein genotype as defined in Annex I has been reported in the last six months;]	to Decision 2002/1003/EC, coming		
	II.2.7.	any transport vehicles or containers in whi authorised disinfectant;	ch they were loaded were cleaned and disinfed	ted before loading with an officially		
	II.2.8.	they were examined by an official veterina	arian within 24 hours of loading and showed no	o clinical sign of disease;		
	II.2.9.	described under box reference I.15 abov	Union on (dd/mm/ e that were cleaned and disinfected before in es, urine, litter or fodder could not flow or f	bading with an officially authorised		
II.3.	Animal	welfare attestation				
	loading i		y, that the animals described above have bee Regulation (EC) No 1/2005, in particular as reg			
Notes						
This certific after impor		eant for live domestic ovine animals (Ovis ar	ies) and domestic caprine animals (Capra hircu	s) intended for immediate slaughter		
After impo	tation the	e animals must be conveyed without delay	to the slaughterhouse 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	g in Part 1 of Annex I to Regulation (EU) No 2	206/2010.		
— Box ref No 206		3: The assembly centre, if any, must fulfil th	he conditions for its approval, as laid down in F	Part 5 of Annex I to Regulation (EU)		
		15: Registration number (railway wagons or g and reloading, the consignor must inform	container and lorries), flight number (aircraft) of the BIP of entry into the Union.	or name (ship) is to be provided. In		
- Box ref	erence I.1	19: Use the appropriate HS code: 01.04.10	or 01.04.20.			
- Box ref	erence I.2	23: For containers or boxes, the container r	number and the seal number (if applicable) sho	ould be included.		

COUNTRY		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	- 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 appropriate the second seco	Species: Select amongst "Ovis aries" and "Capra hircus" as appropriate.							
Age: months.								
Sex (M = male, F = female, C = castrated).								
Part II:								
(1) Code of the territory as it appears in Part 1 of Annex I to Regulatio	n (EU) No 206/2010.							
(²) Keep as appropriate.								
(³) Guarantees in relation to a programme of control of scrapie, as requand Chapter E of Annex IX to Regulation (EC) No 999/2001.	lested by the EU Member State of des	stination, in application of Article 15						
(4) Date of loading. Imports of these animals shall not be allowed wh exportation to the Union of the third country, territory or part there measures have been adopted by the Union against imports of these	of referred to in boxes I.7 and I.8, o	r during a period where restrictive						
(5) Surveillance programme as laid down in Annex I to Commission Re	gulation (EC) No 1266/2007 (OJ L 28	3, 27.10.2007, p. 37.).						
Official veterinarian								
Name (in capital letters):	Qualification and title:							
Date:	Date: Signature:							
Stamp:	Stamp:							

					Mode	I POR-X					
		UNTRY				1			Veterinary ce	rtificate to EU	
	l.1.	Consignor				I.2. Certific	ate referenc	e numbe	r I.2.a.		
		Name					I.3. Central Competent Authority				
		Address					-				
		Tel. No				I.4. Local C	Jompetent A	uthonty			
ţ	I.5.	Consignee				I.6.					
nme		Name Address									
nsig											
Sol		Postal code									
chec		Tel. No									
Part I: Details of dispatched consignment	1.7.	Country ISO I.8. Region Code of origin code of origin			I.9. Countr destina		ISO code	I.10. Region of destination	Code		
ils o	l.11.	Place of origin				I.12.	I				
l: Detai		Name Approval number Address Name Approval number Address									
Part											
		Name Address	Approval number								
	I.13. Place of loading Address Approval number					I.14. Date of departure time of departure					
	I.15. Means of transport Aeroplane Ship Railway wagon				I.16. Entry BIP in EU						
		Road vehicle	Oth	er 🗌		1.17.					
		Identification: Documentary ref	erences:								
	I.18	. Description of co	mmodity			I.19. Commodity code (HS code) 01.03					
								I.20.	Quantity		
	I.21							1.22.	Number of packag	es	
	1.23	. Identification of c	ontainer/s	eal number				1.24.			
	I.25	. Commodities cer Breed		Fattening							
	1.26.					I.27. For import or admission into EU					
	1.28	. Identification of t	1								
	Species Identification (Scientific name) system				Identificatio number	n	A	ge	Sex		

	COUNTRY					Model POR-X					
	Ш.	Health	information		II.a. Certificate reference number	II.b.					
	II.1.	II.1. Public Health Attestation									
		I, the undersigned official veterinarian, hereby certify, that the animals described in this certificate:									
tion		II.1.1	case of bruce	llosis, for the	h have been free from any official prohibition o e last 30 days in the case of anthrax and for the n in contact with animals from holdings which (e past six months in the case of rabies and,					
Part II: Certification		II.1.2	have not rece	ived:							
II: Ce			 any stilbe 	ne or thyros	static substances,						
Part			-	-	nic, gestagenic or β- agonist substances for pu I in Directive 96/22/EC).	rposes other than therapeutic or zootechnic					
	II.2.	Anima	I Health attest	ation							
		I, the u	ndersigned offi	cial veterina	arian, hereby certify, that the animals described	above meet the following requirements:					
		II.2.1	they come fro	m the territo	pry with code:(1) which,	, at the date of issuing this certificate:					
			(²) either	swine	been free for 24 months from foot-and-mouth dis e fever, classical swine fever, swine vesicular nths from vesicular stomatitis, and]						
			(²) or	A	has been free [for 24 months from foot-and-mout African swine fever, vesicular exanthema, [cla disease] (²), and for 6 months from vesicular sto	ssical swine fever] (2) and [swine vesicular					
				[! 	has been considered free from [foot-and-mouth swine vesicular disease] (²), since had cases/outbreaks from that date, and authori Regulation (EU) No/, of	(dd/mm/yyyy), without having ised to export these animals by Commission					
				and	e during the last 12 months, no vaccination ag imports of domestic cloven-hoofed animals v iitted;						
		II.2.2			e territory described under point II.2.1 since bin d without contact with imported cloven-hoofed						
		II.2.3	dispatch, and	, during this	e holding(s) described under box reference I.1 period, in the holding(s) and in an area with a * putbreak of the diseases referred to in point II.2	10 km radius around the holding(s) of origin,					
		II.2.4 A			e killed under a national programme for the e seases referred to in point II.2.1;	radication of diseases, nor have they been					
(2) (3) [II.2.4 B they have been subjected within the past 30 days to a test for swine vesicular disease an swine fever antibodies with negative results in both cases];					lar disease antibodies and a test for classical						
	(²) (⁴) [II.2.4 C they have been subjected within the past 30 days to a buffered Brucella antigen test for negative results];					Ila antigen test for porcine brucellosis with					
II.2.5 they come from herds which are not restricted under the national brucellosis eradication programme;					losis eradication programme;						
II.2.6 they are/were (²) dispatched from their holding(s) of origin, without passing through any market,						ing through any market,					
			(²) either	[directly t	to the Union,]						
			(²) or	-	officially authorised assembly centre described under box reference I.13 situated within the described under point II.2.1,]						

COUNTR	Y			Model POR-X							
П.	Health	information	II.a. Certificate reference number	II.b.							
			n contact with other cloven-hoofed animals no	t 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									
			ik of any of the diseases referred to in point II.2								
	II.2.7	any transport vehicles or officially authorised disin	containers in which they were loaded were cle fectant;	eaned and disinfected before loading with an							
	II.2.8	they were examined by a	an official veterinarian within 24 hours of loadir	ng and showed no clinical sign of disease;							
	II.2.9	transport described und	for dispatch to the Union on ler box reference I.15 above that were clear fectant and so constructed that faeces, urine, ng transportation.	ned and disinfected before loading with an							
II.3.	Anima	I transport attestation									
	at the t	ime of loading in accorda	inarian, hereby certify, that the animals desc ance with the relevant provisions of Regulation are fit for the intended transport.								
(²) (⁶) [II.4.	Specif	c requirements									
	[11.4.1	Aujeszky's disease is not	tifiable in the country referred to in box referen	ice I.7;							
	II.4.2		rmation, no clinical, pathological or serologic months in the holding(s) of origin referred to hin 5 km;								
	II.4.3	the animals referred to in	box reference I.28:								
			r exportation, have remained since birth in y have remained in this(ese) holdings(s) for the second s								
			in accommodation approved by the competer export, without direct or indirect contact with o								
			d to an ELISA test for the presence of gI antii vith negative results; and, all animals in isolatio								
			nated against Aujeszky's disease and have not s not been vaccinated during the previous 12 r								
(2) (8)	[11.4.4]	(further requirements and/or tests)							
Notes											
This certif	icate is	meant for live domestic po	prcine animals (Sus scrofa) intended for breed	ing or production.							
After impo	ortation	the animals must be con	veved without delay to the holding of destina	tion where they shall remain for a minimum							

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

со	DUNTRY		Model POR-X									
II.	Health information	II.a. Certificate reference number	II.b.									
Pa	Part I:											
	 Box reference I.8: Provide the code of territory as appearing in Part 1 of Annex I to Regulation (EU) No 206/2010. 											
_	 Box reference I.13: The assembly 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. 											
_		er (railway wagons or container and lorries), f ading, the consignor must inform the BIP of e	light number (aircraft) or name (ship) is to be ntry into the Union.									
—	Box reference I.23: For containers or bo	oxes, the container number and the seal num	per (if applicable) should be included.									
—	Box reference I.28: Identification system	n: the animals must bear:										
	brand, chip, transponder).		ne identification system (such as tag, tattoos,									
	origin.	de of the exporting country. The individual nu	mber must permit tracing of their premises of									
_	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 Pa	rt 1 of Annex I to Regulation (EU) No 206/20 ⁻	0.									
(²)	Keep as appropriate.											
(3)	Supplementary guarantees to be provi with the entry 'B'.	ded when required in column 5 'SG' of Part 1	of Annex I to Regulation (EU) No 206/2010,									
(4)	Supplementary guarantees to be provi with the entry 'C'.	ded when required in column 5 'SG' of Part '	of Annex I to Regulation (EU) No 206/2010,									
(5)	for exportation to the Union of the third	I country, territory or part thereof referred to	loaded either prior to the date of authorisation n boxes I.7 and I.8, or during a period where imals from this third country, territory or part									
(⁶)	between the Community and the Swiss		vith Decision 2008/185/EC and the Agreement ucts (OJ L 114, 30.4.2002, p. 132) except for lation (EU) No 206/2010.									
(7)	To be carried out according to the stand the test used shall be the whole virus E		35/EC. In the case of pigs aged over 4 months,									
(⁸)	Further requirements requested by Finl	and in respect of transmissible gastro-enterit	s.									
Off	icial veterinarian											
	Name (in capital letters):	Qualificatio	on and title:									
	Date:	Signature:										
	Stamp:											

▼<u>C1</u>

			Mode	I POR-Y					
		JNTRY					Veterinary cer	tificate to EU	
	l.1.	Consignor		I.2. Certific	ate reference	e numbe	r I.2.a.		
		Name		I.3. Central Competent Authority					
		Address		I.4. Local C		uthority			
		Tel. No		1.4. Local C	ompetent At	linomy			
ent	1.5.	Consignee		I.6.					
m		Name							
nsig		Address							
d Co		Postal code							
che		Tel. No							
Part I: Details of dispatched consignment	1.7.	Country ISO I.8. Region of origin code of origin	Code	I.9. Country destina		ISO code	I.10. Region of destination	Code	
ils o	I.11.	Place of origin	I.12.						
I: Deta		Name Approval numb Address							
Part		Name Approval numb Address							
		Name Approval numb Address	per						
	I.13	Place of loading Address Approval numb	I.14. Date of departure time of departure						
	I.15	Means of transport	I.16. Entry BIP in EU						
			ay wagon 🗌						
		Road vehicle Other		1.17.					
		Identification: Documentary references:							
	I.18	Description of commodity		I.19. Commodity code (HS code) 01.03					
						1.20.0	Quantity		
	I.21					1.22.1	Number of package	es	
	1.23	Identification of container/seal number				1.24.			
	1.25	Commodities certified for: Slaughter							
	1.26		I.27. For import or admission into EU						
	1.28	Identification of the commodities		1					
		Species Identifica (Scientific name) system	Identification number	1	A	ge	Sex		

	COUNTRY Model I				Model POR-Y			
	Ш.	Health	information		II.a. Certificate reference number	II.b.		
tion	II.1.	Public	Health Attesta	ation				
		I, the undersigned official veterinarian, hereby certify, that the animals described in this certificate:						
		II.1.1	case of brucel	losis, for th	h have been free from any official prohibition o e last 30 days in the case of anthrax and for th n in contact with animals from holdings which t	e past six months in the case of rabies and,		
Part II: Certification		II.1.2	have not recei	ved:				
ll: Ce			 any stilber 	ne 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 attest	ation				
		I, the u	ndersigned offic	cial veterina	arian, hereby certify, that the animals described	above meet the following requirements:		
		II.2.1	they come fror	m the territo	pry with code:(1) which	, at the date of issuing this certificate:		
			(²) either	swine	been free for 24 months from foot-and-mouth dis e fever, classical swine fever, swine vesicular onths from vesicular stomatitis, and]			
			(²) or	L , , , , , ,	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		
				[: C	has been considered free from [foot-and-mout 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 an imports of domestic cloven-hoofed animals v nitted.			
		II.2.2			e territory described under point II.2.1 since birt d without contact with imported cloven-hoofed			
		II.2.3	dispatch, and,	during this	e holding(s) described under box reference I.1 period, in the holding(s) and in an area with a putbreak of the diseases referred to in point II.2	10 km radius around the holding(s) of origin,		
		II.2.4	•		e killed under a national programme for the e seases referred to in point II.2.1;	radication of diseases, nor have they been		
		II.2.5	they are/were	(²) dispatch	ned from their holding(s) of origin, without pass	ing through any market,		
			(²) either	[directly t	to the Union,]			
			(²) or	-	ficially authorised assembly centre described described under point II.2.1,]	under box reference I.13 situated within the		
			and, until disp	atched to th	he Union:			
			., .		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.			

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

COU	COUNTRY Model POR-1								
II.	Health information	II.a. Certificate reference number	II.b.						
Part	Part II:								
(1) C	(1) Code of the territory as it appears in Part 1 of Annex I to Regulation (EU) No 206/2010.								
	eep as appropriate.								
fc re	or 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						
(4) V	/hen required by the EU Member State	e of destination, in accordance with Decisio	on 2008/185/EC.						
Offici	al veterinarian								
	Name (in capital letters):	Qualifica	tion and title:						
	Date:	Signature	e:						
	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			
ıt		Tel.	I.4. Local competent authority			
nmer	l.5.	Consignee	1.6.			
onsig		Name Address				
atched co		Postal code Tel.				
Part I: Details of dispatched consignment	I.7.	Country of origin ISO code I.8. Region of origin Code	I.9. Country of ISO code I.10. Region of Code destination			
etail	l.11.	Place of origin	1.12.			
Part I: C		Name Approval number Address				
	l.13.	Place of loading	I.14. Date of departure			
		Address Approval number				
	l.15.	Means of transport	I.16. Entry BIP in EU			
		Aeroplane Ship Railway wagon				
		Road vehicle Other I Identification Documentary references	I.17. No(s) of CITES			
	l.18.	Description of commodity	I.19. Commodity code (HS code)			
			I.20. Quantity			
	I.21.		I.22. Number of packages			
	1.23.	Seal/Container No	1.24.			
	I.25.	Commodities certified for:				
		Breeding E Fattening	Slaughter			
	I.26.		I.27. For import or admission into EU			
	1.28.	Identification of the commodities	<u> </u>			
		Species Identification system Identifi (scientific name)	cation number Age Sex			

col	INTRY					Model RUM				
	11.	Health	information		II.a. Certificate reference number	II.b.				
	II.1.	Public	Health Attest	tation						
		l, the ι	undersigned of	ficial veterinarian, hereby certify, that the	e animals described in this certificate:					
tion		II.1.1.	come from a holding which has been free from any official prohibition on health grounds, for the last 42 days in the case of brucellosis and tuberculosis, for the last 30 days in the case of anthrax, for the last six months in the case of rabies, and, have not been in contact with animals from holdings which did not satisfy these conditions;							
II: Certification		II.1.2.	have not rece	eived:						
≕ E			— any stilbe	ne or thyrostatic substances,						
Part				ic, androgenic, gestagenic or β- agonist d in Directive 96/22/EC).	t substances for purposes other than	therapeutic or zootechnic treatment				
	11.2.	Anima	l Health Attes	station						
		l, the ι	undersigned of	ficial veterinarian, hereby certify, that the	e animals described above meet the t	following requirements:				
		II.2.1.	they come fro	om the territory with code:	(¹) which, at the d	ate of issuing this certificate:				
		(a) has been free for 24 months from foot-and-mouth disease and bluetongue, for 12 months from rinderpest, Rift valley fever, contagious bovine pleuropneumonia, lumpy skin disease, peste des petits ruminants, sheep pox and goat pox, contagious caprine pleuropneumonia and epizootic haemorrhagic disease and for six months from vesicular stomatitis, and								
 (b) where during the last 12 months, no vaccination against foot-and-mouth disease, rinderpest, Rift valley fe bovine pleuropneumonia, lumpy skin disease, peste des petits ruminants, sheep pox and goat pox, com pleuropneumonia and epizootic haemorrhagic disease and during the last 24 months no vaccination against been carried out and imports of cloven-hoofed animals vaccinated against these diseases are not permitted. II.2.2. they have remained 					and goat pox, contagious caprine vaccination against bluetongue has					
			(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 the relevant species listed Part 7 of Annex I to Regulation (EU) No 206/2010 and they were imported directly under the conditions specific for each species in Part 7 of Annex I to Regulation (EU) No 206/2010 from a third country during a period of le than six months prior to embarkation to the Union and in any case they have been separated from other anima not of the same health status after being released in the exporting country and before exportation to the Union (³)]						
		II.2.3.	they have re reference I.1	mained since birth or at least 40 days 1 and I.13:	before dispatch in the holding/establ	lishment (²) described under boxes				
				round which in an area of radius of 1 agic disease during the previous 60 da		break of bluetongue and epizootic				
				ound which in an area of 10 km radius, t ing the previous 40 days;	here has been no case/outbreak of the	e other diseases referred to in point				
		II.2.4.		animals to be killed under a national pro of the diseases referred to in point II.2.1		es, nor have they been vaccinated				
			(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 no	t been vaccinated against brucellosis a	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 ag agglutination per ml, within the past 3		icella count of less than 30 IU of				
			(²) or	[are castrated males of any age;]						

COUN	TRY			Model RUM
п.	Health	information	II.a. Certificate reference number	II.b.
	II.2.5.	according to my knowledge and to the written declara	tion made by the owner, the animals	:
		(a) do not come from holdings/establishments $(^{2}),$ an which the following diseases have been clinically		nals of a holding/establishment, in
		 (i) contagious agalactia of sheep or goats (<i>Myco mycoides</i> '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/encephali	tis,	
		(²) <i>either</i> [within the last three years,]		
			the infected animals were slaughtered asts carried out at least six months a	
		(b) are included in an official system for notification o	f these diseases, and	
		(c) have been free from clinical or other evidence of	tuberculosis and brucellosis during th	e three years prior to export;
	(²) (⁶) [II.2.6.	the animals have reacted negatively to a serological rhagic-disease, carried out on two occasions on samp at least 28 days later on	les of blood taken at the beginning of	the isolation/quarantine period and
	II.2.7.	they are dispatched from the holding/establishment des dispatched to the Union:	cribed under boxes reference I.11 and	d I.13 directly to the Union and, until
		 (a) they did not come in contact with other cloven-hoo 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 whic case/outbreak of any of the diseases referred to i		previous 30 days there has been a
	II.2.8.	any transport vehicles or containers in which they we authorised disinfectant;	re loaded were cleaned and disinfect	ted before loading with an officially
	II.2.9.	they were examined by an official veterinarian within a	24 hours of loading and showed no c	linical sign of disease;
	II.2.10.	they have been loaded for dispatch to the Union on . under box reference I.15. above that were cleaned and constructed that faeces, urine, litter or fodder could ne	disinfected before loading with an offi	cially authorised disinfectant and so
11.3.	Anima	I transport attestation		
	loading	undersigned official veterinarian, hereby certify, that the i in accordance with the relevant provisions of Regulatio 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 I.28 .:		
		 (a) have been isolated in accommodation approved by for export, and 	the competent authority for the last 30) days immediately prior to dispatch
		(b) have been subjected to a serological test for IBR results, and all animals in isolation have also give		r entry into isolation, with negative

col	JNTRY			Model RUM				
II.	Health in	formation	II.a. Certificate reference number	II.b.				
	(c)	have not been vaccinated against IBR.;						
	(²) [II.4.3	(further requirement	s and/or tests)]]				
No	tes							
		ant for live animals of the order Artiodactyla (excludi <i>Capra hircus</i> , Suidae and Tayassuidae), and of the fa						
	After importation the animals must be conveyed without delay to the holding of destination where they shall remain for a minimum period of 30 days before further movement outside the holding, except in the case of a dispatch to a slaughterhouse.							
Pa	rt I:							
_	Box reference 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.1: No 206/2010.	3.: The assembly centre, if any, must fulfil the condit	ions for its approval, as laid down in F	art 5 of Annex I to Regulation (EU)				
		5.: Registration number (railway wagons or containe g and reloading, the consignor must inform the BIP		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	3.: For containers or boxes, the container number a	und the seal number (if applicable) she	ould be included.				
		8.: Identification system: Specify the identification systring country. The individual number must permit tra		nder). The ear tag includes the ISO				
	Age: months.							
	Sex (M = male, F	= female, C = castrated).						
	Species: Select t	he species amongst those listed for the following fa	milies:					
	Antilocapridae:	Antilocapra spp.;						
	Bovidae:	Addax spp., Aepyceros spp., Alcelaphus spp., Ami laphus spp., Budorcas spp., Capra spp. (excluding (including Beatragus), Dorcatragus spp., Gazella s Madoqua spp., Naemorhedus spp. (including Nemi spp., Oryx spp., Ourebia spp., Ovibos spp., Ovis Pseudois spp., Pseudoryx spp., Raphicerus spp., R Sylvicapra spp., Syncerus spp., Taurotragus spp.,	Capra hircus), Cephalophus spp., Co spp., Hemitragus spp., Hippotragus s orhaedus and Capricornis), Neotragus spp. (excluding Ovis aries), Pantholop ledunca spp., Rupicapra spp., Saiga s	nnochaetes spp., Damaliscus spp. pp., Kobus spp., Litocranius spp., spp., Oreamnos spp., Oreotragus s spp., Pelea spp., Procapra spp., pp., Sigmoceros-Alecelaphus spp.,				
	Camelidae:	Camelus spp., Lama spp., Vicugna spp.						
	Cervidae: Alces spp., Axis-Hyelaphus spp., Blastocerus spp., Capreolus spp., Cervus-Rucervus spp., Dama spp., Elaphurus spp., Hippocamelus spp., Hydropotes spp., Mazama spp., Megamuntiacus spp., Muntiacus spp., Odocoileus spp., Ozotoceros spp., Pudu spp., Rangifer spp.							
	Giraffidae:	Biraffidae: <i>Giraffa</i> 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.						

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

	00	UNTRY		Mod	el SUI				Votorinory	stificate to Ell
		Consignor			1.2. C	ertifica	te reference	number	~	ertificate to EU
		Name		ortinou						
		Address			I.3. C	entral (Competent A	uthority		
		Tel. No			I.4. Lo	ocal Co	ompetent Aut	hority		
ŧ	I.5.	Consignee			1.6.					
mer		Name								
sign		Address								
con		Postal code								
hed		Tel. No								
Part I: Details of dispatched consignment	1.7.	Country ISO of origin code	I.8. Region of origin	Code		ountry estinati		SO ode	I.10. Region of destination	Code
ls of	I.11.	Place of origin	I		I.12.			I		
l: Detai		Name Address	Approval number							
Part		Name Address	Approval number							
	Name Approval number Address									
	I.13	I.13. Place of loading Address Approval number				I.14. Date of departure time of departure				
	I.15. Means of transport				I.16. E	ntry BIF	P in EU			
	Aeroplane 🗌 Ship 🗌 Railway wagon 🗌									
		Road vehicle Othe	er 🗌		I.17. No(s) of CITES					
		Identification: Documentary references:			1.17.180	J(S) UI C	1125			
	I.18	. Description of commodity					I.19. Comm	odity cc	ode (HS code)	
								1.20.0	Quantity	
	I.21							1.22.1	Number of packa	ges
	I.23. Identification of container/seal number I.25. Commodities certified for:							I.24.		
	Breeding Fattening							Slau	ughter	
					I.27. F	or impo	ort or admiss	on into	EU	
	I.28. Identification of the commodities									
		Species (Scientific name)	Identification system		Identifi num			Aç	je	Sex

Model SUI

	COUNT	RY			Model SU			
	Ш.	Health	information	II.a. Certificate reference number	II.b.			
Part II: Certification	II.1. Public Health Attestation							
		I, the u	ndersigned official veterina	arian, hereby certify, that the animals describe	d in this certificate:			
		II.1.1	case of brucellosis, for th	ch has been free from any official prohibition of le last 30 days in the case of anthrax and for the en in contact with animals from holdings which	ne past six months in the case of rabies and,			
		II.1.2	have not received:					
t II: C			 any stilbene or thyros 	static substances,				
Par				enic, gestagenic or β - agonist substances for p d in Directive 96/22/EC).	urposes other than therapeutic or zootechnic			
	II.2.	Anima	I Health attestation					
		I, the u	ndersigned official veterina	arian, hereby certify, that the animals describe	d above meet the following requirements:			
		II.2.1	they come from the territo	ory with code:(1) which	n, at the date of issuing this certificate:			
				months from foot-and-mouth disease, for 12 r, swine vesicular disease and vesicular examples and vesicular example				
				t 12 months, no vaccination against these dis Is vaccinated against these diseases are not				
		II.2.2		e territory described under point II.2.1 since b I without contact with cloven-hoofed animals in				
		II.2.3	dispatch, and, during this	e holding described under boxes reference I.1 period, in the holding(s) and in an area with a putbreak of the diseases referred to in point II.	10 km radius around the holding(s) of origin,			
		II.2.4 A	vaccinated against the di	be killed under a national programme for the or seases referred to in point II.2.1 and they have test for porcine brucellosis with negative resu	e been subjected within the past 30 days to a			
	(²) (³)	[II.2.4 B		ed within the past 30 days to a test for swine bodies with negative results in both cases]	vesicular disease antibodies and a test for			
	(2) (4)	[II.2.4 C	they have been subjecte negative results]	d within the past 30 days to a buffered Bruc	ella antigen test for porcine brucellosis with			
		II.2.5	they come from holdings	which:				
				nder a national control and eradication prog eschen disease), and	gramme for brucellosis, porcine enteroviral			
			(b) are included in an off	ficial system for notification of these diseases;				
		II.2.6	they are dispatched from dispatched to the Union:	the holding described under boxes reference	I.11 and I.13 directly to the Union and, until			
			(a) they did not come in described in this cert	contact with other cloven-hoofed animals no ificate, and	t complying with the health requirements as			
				place where, or around which within a 10 km r k of any of the diseases referred to in point II.2	• • •			

COUNTRY			Model SUI				
II. Hea	Ith information	II.a. Certificate reference number	II.b.				
II.2.	II.2.7 any transport vehicles or containers in which they were loaded were cleaned and disinfected before loading with an officially authorised disinfectant;						
II.2.	3 they were examined by a	an official veterinarian within 24 hours of loading	g and showed no clinical sign of disease;				
II.2.	II.2.9 they have been loaded for dispatch to the Union on						
ll.3. Ani	nal transport attestation						
time		arian, hereby certify, that the animals described ith the relevant provisions of Regulation (EC) N the intended transport.					
(²) (⁶) [II.4. Spe	cific requirements						
II.4.	1 Aujeszky's disease is no	tifiable in the country referred to in box reference	ce I.7;				
II.4.		rmation, no clinical, pathological or serologica nonths in the holding(s) of origin referred to in b d the holding(s);					
II.4.	3 the animals referred to in	box reference I.28:					
		r exportation, have remained since birth in 13 or they have remained in this holding for th					
		in 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 isolatior					
		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)				
Notes							
		stic Suidae (<i>Babyrousa</i> spp., <i>Hylochoerus</i> spp. p. <i>,Pecari</i> spp., <i>Tayassu</i> spp.) and Tapiridae (<i>Ta</i> j					
	After importation the animals must be conveyed without delay to the holding of destination where they shall remain for a minimum period of 30 days before further movement outside the holding, except in the case of a dispatch to a slaughterhouse.						

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

	со	UNTRY	Veterinary certificate to EU					
	l.1.	Consignor	I.2. Certificate reference number I.2.a.					
		Name	I.3. Central Competent Authority					
		Address						
		Tel. No	I.4. Local Competent Authority					
nt	I.5.	Consignee	1.6.					
nme		Name						
nsig		Address						
d col		Postal code						
chea		Tel. No						
Part I: Details of dispatched consignment	I.7.	Country ISO I.8. Region Code of origin code of origin	I.9. Country of ISO I.10. Region of Code destination code destination					
ils o	I.11.	. Place of origin	1.12.					
I: 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					
	l.15	. Means of transport Aeroplane Ship Railway wagon	I.16. Entry BIP in EU					
		Road vehicle Other	I.17. No(s) of CITES					
		Identification: Documentary references:	1.17. NO(5) OF CITES					
	I.18	. Description of commodity	I.19. Commodity code (HS code) 01.06.19					
			I.20. Quantity					
	I.21		I.22. Number of packages					
	1.23	. Identification of container/seal number	1.24.					
	1.25	Commodities certified for:	2					
		Breeding Fattening	Slaughter					
	I.26		I.27. For import or admission into EU					
	1.28	. Identification of the commodities						
		Species Identification (Scientific name) system	Identification Age Sex number					

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

	COUNT	TRY					Model CAM					
	Π.	Health	information	II.a. Certificate refere	nce number	II.b.						
Part II: Certification	II.1.	Quarantine conditions attestation										
		I, the undersigned official veterinarian, hereby certify, that the animals described in the animal health certificate (1) number										
		(date (dd/mm/yyyy) of entry (²)) in the quarantine station of St. Pierre and Miquelon under the conditions provided for in Part 7 of Annex I to Regulation (EU) No 206/2010 for a period of:										
t II: Cei		II.1.1.	Brucellosis:									
Par			(a) <i>B. abortus</i> least 42 da	Serum Agglutination Test (SAT) a	nd Rose Bengal Test (F	RBT) within two days after arri	ival and after at					
			(b) <i>B. ovis</i> : Co	mplement Fixation Test (CFT) with	nin two days after arriva	al and after at least 42 days						
			(c) <i>B. meliten</i>	sis: SAT and RBT within two days	after arrival and after at	least 42 days						
		II.1.2.	Bluetongue ar	d Epizootic haemorrhagic disease	9							
			(⁵) either	[two tests using Bluetongue co 21 days]	mpetitive Elisa test wit	hin two days after arrival an	d after at least					
			(⁵) or	[they have been quarantined fo remained free of Bluetongue ve detected].		U ,						
		II.1.3.	Tuberculosis									
				al tuberculin test according to ar in two days after arrival and after		5	vian tuberculin					
		II.1.4.		h disease: ELISA test for the det d after at least 42 days	ection of antibodies an	nd a virus neutralizaton test v	within two days					
		II.1.5.	Rinderpest: co	mpetitive ELISA test within two da	lys after arrival and afte	er at least 42 days						
		II.1.6.	Vesicular storr	atitis: ELISA or virus- neutralisatic	n test within two days a	after arrival and after at least	42 days					
		II.1.7.	Rift valley feve	r: an ELISA test or a virus neutrali	sation test within two da	ays after arrival and after at le	east 42 days					
		II.1.8.	Lumpy skin di	ease: ELISA or virus neutralisatio	n test within two days a	fter arrival and after at least 4	42 days					
		II.1.9.	Crimean Cong 42 days	o haemorrhagic fever: ELISA or vi	rus neutralisation test v	vithin two days after arrival ar	nd after at least					
		II.1.10.	Surra: blood m	icroscopy within two days after an	rival and after at least 4	2 days						
		II.1.11.	Malignant cata	rrhal fever: immunofluorescence t	est within two days afte	er arrival and after at least 42	days					
	II.2.	Supple	ementary guar	antees								
		II.2.1		s: AGID test or ELISA within two d of destination) (⁵)	ays after arrival and afte	er at least 42 days (When req	uired by the EU					

II.	Health	information		II.a. Certificate reference number	II.b.							
11.3.	Treatm	Treatments										
	They have been subjected to:											
	II.3.1. an internal and external antiparasitic treatment during the quarantine period											
	II.3.2.			5								
	11.0.2.											
		(⁵) either	la treatr	nent with streptomycin 25mg/kg]								
		(⁵) or		biotic treatment effective against Leptosp	ira spp. (specify							
	(⁵) [II.3.3.		•	vies (if requested) on and with the test result	(dd/mm/yyyy) using vaccine							
Notes												
This ce	ertificate is	meant for live	animals of t	the family Camelidae.								
Part I:												
— Во	x reference	e I.8: Provide t	he code of t	erritory as appearing in Part 1 of Annex I to	o Regulation (EU) No 206/2010.							
				tre, if any, must fulfil the conditions for its	s approval, as laid down in Part 5 of Annex I to							
		U) No 206/20										
				er (railway wagons or container and lorries ading, the consignor must inform the BIP o	s), flight number (aircraft) or name (ship) is to be of entry into the Union.							
— Во	x reference	e I.23: For con	tainers or be	oxes, the container number and the seal n	umber (if applicable) should be included.							
— Во	ox reference	e I.28: Identific	ation syster	<i>n</i> : The animals must bear:								
—				nits tracing of their premises of origin. S) and the anatomic place used in the ar	Specify the identification system (such as tag nimal.							
_	An ear ta origin.	g that include	s the ISO co	ode of the exporting country. The individual	I number must permit tracing of their premises o							
— Во	x reference	e I.28: <i>Age</i> : mo	onths.									
— Во	ox reference	e I.28: <i>Sex</i> (M	= male, F =	female, $C = castrated$).								
— Во	x reference	e I.28: Species	: Select am	ongst <i>'Camelus</i> spp.', <i>'Lama</i> spp.', 'Vicugn	<i>a</i> spp.' as appropriate.							
Part II:	:											
· /		n certificate for Regulation (E			to the Union (model 'RUM') as laid down in Part 2							
²) Da	ate in which	the last anim	al in a group	o entered the quarantine facility.								
³) Te) Tests performed in accordance with the methods described in Chapter 2 of Part 7 of Annex I to Regulation (EU) No 206/201											
(⁴) Re	esults of the	e tests perform	ied must be	attached in original to this health attestation	on.							
(⁵) Ke	ep as appr	opriate.										
() 10												

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

PART 3

Addendum for transport of animals by sea

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

Declaration by the master of the ship

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 ro (<i>Ports of call en route</i>). Moreover, during the jour with other animals on board of a lower health state	nained 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)
(stamp)	(signature of master)

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

Test protocol:

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

Mab controlColumns 1 and 2, rows G and H are the monoclonal
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.
- 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 μ l to all wells of the plate.
- 7. Incubate at 37 °C for 60 minutes on an orbital shaker. Wash three times with PBS and blot dry.
- 8. Thaw the O-Phenylenediamine dihydrochloride (OPD) and immediately before use add 5 μl of 30 % hydrogen peroxide to each 10 ml of OPD. Add 50 μl to all wells of the plate. Allow colour to develop for approximately 10 minutes and stop the reaction with 1 Molar sulphuric acid (50 μl per well). Colour should develop in the Mab control wells and in those wells containing sera with no antibody to BTV.
- 9. Examine and record the plates either visually or using a spectrophotometric reader.

Analysis of results:

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

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

Percentage inhibition (PI) value = 100 – (OD 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.

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.
	vine rhinotracheitis (IBR) / infectious ustular vulvo-vaginitis (IPV)
A. The serum neut protocol:	ralisation 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 test	recognised in the framework of Decision 2004/558/EC (1).
F	oot-and-mouth disease (FMD)
A. Collecting oeso	phageal/pharyngeal samples and testing shall be carried out

- A. Collecting oesophageal/pharyngeal samples and testing shall be carried out according to the following protocol:
 - Reagents: Prior to sampling, transport medium is prepared. Two ml volumes are dispensed in as many containers as there are animals to be sampled. The containers used must withstand freezing over solid $\rm CO_2$ 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.
- 6. The plates are washed and 50 μ l 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 neutralis protocol:	sation test shall be carried out according to the following

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

Transmissible gastro-enteritis (TGE)

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

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

▼<u>C1</u>

(¹) OJ L 59, 4.3.2008, p. 19.

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

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

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

Swine vesicular disease (SVD)

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

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 Territory	iter Description of this has not a function of the set of	Veterinary ce	Veterinary certificate			
third country		Description of third country, territory or part thereof	Model(s)	SG	conditions	Closing date (²)	Opening date (³)
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,	BOV	A	1		18 March 2005
	La Rioja, Mendoza, Misiones, Part of Neuquén (excluding territory included in Part of Río Negro (excluding territory included in San Juan, San Luis, Santa Fe, Tucuman, Cordoba, La Pampa, Santiago del Estero, Chaco, Formosa, Jujuy and Salta, excluding the bu of 25 Km from the border with Bolivia and Parag extends from the Santa Catalina District in the Pro	Mendoza, Misiones, Part of Neuquén (excluding territory included in AR-4), Part of Río Negro (excluding territory included in AR-4), San Juan, San Luis,	RUF	A	1		1 December 2007
		Tucuman, Cordoba, La Pampa,	RUW	A	1		1 August 2010

VI	M2
----	----

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	СН-0	Whole country	*				
CL – Chile	CL-0	Whole country	BOV, OVI, POR, EQU, RUF, RUW, SUF				
CN – China	CN-0	Whole country	_				
CO – Colombia	СО-0	Whole country	EQU				
CR – Costa Rica	CR-0	Whole country	BOV, EQU				
CU – Cuba	CU-0	Whole country	BOV, EQU				
DZ – Algeria	DZ-0	Whole country	_				
ET – Ethiopia	ET-0	Whole country	_				
FK – Falkland Islands	FK-0	Whole country	BOV, OVI, EQU				
GL – Greenland	GL-0	Whole country	BOV, OVI, EQU, RUF, RUW				
GT – Guatemala	GT-0	Whole country	BOV, EQU				
HK – Hong Kong	НК-0	Whole country	_				
HN – Honduras	HN-0	Whole country	BOV, EQU				
HR – Croatia	HR-0	Whole country	BOV, OVI, EQU, RUF, RUW				
IL – Israel	IL-0	Whole country					

1	2	3	4	5	6	7	8
IN – India	IN-0	Whole country	_				
IS – Iceland	IS-0	Whole country	BOV, OVI, EQU, RUF, RUW				
KE – Kenya	KE-0	Whole country	_				
MA – Morocco	MA-0	Whole country	EQU				
ME – Montenegro	ME-0	Whole country	BOV, OVI, EQU				
MG – Madagascar	MG-0	Whole country	_				
MK – Former Yugoslav Republic of Macedonia (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></u>							
1	2	3	4	5	6	7	8
US – United State	s US-0	Whole country	BOV, OVI, POR, EQU,SUF, SUW, RUF, RUW	G			
UY – Uruguay	UY-0	Whole country	EQU				
			BOV,	А	1		1 November 2001
			OVI	А	1		
13							
ZA - South Africa	a 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	
12							
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).
- ^cH²: 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.
- 'J': guarantees regarding the movement of bovine, ovine and caprine animals from holdings to the slaughterhouse, which allow them to pass via an assembly centre (including markets) before being transported directly to slaughter.

Model BOV

cou	NTRY					Veterinary certificate to	o EU
	1.1.	Consignor		I.2. Certificate	reference No	l.2.a.	_
		Name		I.3. Central co	I.3. Central competent authority		
		Address				·	
ŧ		Tel.		I.4. Local com	petent authority		
nme	1.5.	Consignee		I.6.			
onsig		Name					
х х		Address			_		
atche		Postal code					
dispatched consignment		Tel.					
s of	1.7.	Country of origin ISO code	I.8. Region of origin Code	I.9. Country o destination		I.10. Region of Code destination	
Part I: Details of				destination			
÷.	1.11.	Place of origin	1	I.12.			\geq
Pai		Name					
		Address					
	1.10	Diago of looding		I.14. Date of de	parture		
	1.13.	Place of loading		1.14. Date of de	parture		
	l.15.	Means of transport		I.16. Entry BIP	in EU		
		Aeroplane 🗌 Ship 🗌	· · · —				
		Road vehicle D Other [Identification		l.17.			
		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:					
		Human consumption 🔲					
	1.26.			I.27. For import	or admission in	to EU	
	1.00	Identification of the second during					
	1.28.	Identification of the commodities					
		Species Nature (scientific name) commo	lity type	Approval number		packages weigh	
			Aba	ttoir Cutting	piant Cold	d store	

	COUNT	RY			Model BOV			
	11.	Health information		II.a. Certificate reference number	II.b.			
	11.1.	Public Health Attesta	tion					
I, the undersigned official veterinarian declare that I am aware of the relevant requirements of Regulations (EC) No 852/2004, (EC) No 853/2004, (EC) No 854/2004 and (EC) No 999/2001 and certify that the meat of dom- described in Part I was produced in accordance with those requirements, in particular that:								
Part II: Certification	II.1.1.	the [meat] [minced mea with Regulation (EC) N	at] (¹) comes from (an) establishment(s) ir o 852/2004;	nplementing a programme based on th	e HACCP principles in accordance			
ll: Ce	II.1.2.	the meat has been obt	ained in compliance with Section I of An	nex III to Regulation (EC) No 853/200	4;			
Part			neat has been produced in compliance wi arature of not more than – 18 °C;]	th Section V of Annex III to Regulation	(EC) No 853/2004 and frozen to an			
			een found fit for human consumption foll ction I and Chapters I and IX of Section					
			earcass or parts of the carcass have beer x I to Regulation (EC) No 854/2004;]	n marked with a health mark in accorda	ance with Chapter III of Section I of			
			packages of [meat] [minced meat] (¹) have x II to Regulation (EC) No 853/2004;]	e been marked with an identification m	ark in accordance with Section I of			
		II.1.6. the [meat] [minc foodstuffs;	ed meat] $(^1)$ satisfies the relevant criteria	a set out in Regulation (EC) No 2073/	2005 on microbiological criteria for			
			covering live animals and products there n particular Article 29 thereof, are fulfilled		nitted in accordance with Directive			
			ed meat] (¹) has been stored and transp Annex III to Regulation (EC) No 853/2004		requirements of Sections I and V			
		II.1.9. with regard to be	ovine spongiform encephalopathy (BSE):	:				
		(¹) <i>either</i> [II.1.§	9.1. for imports from a country or a 2007/453/EC:	region with a negligible BSE risk	and listed as such in Decision			
			 (a) the country or region is classific country or region posing a negl 	ed in accordance with Article 5(2) of F ligible BSE risk;	Regulation (EC) No 999/2001 as a			
			(b) the animals from which the bovi slaughtered in a country with a	ine meat or minced meat was derived v negligible BSE risk (¹³);	were born, continuously reared and			
			(¹) [(c) if in the country or region there	have been BSE indigenous cases:				
				n after the date from which the ban on reaves derived from ruminants had be				
				inced meat does not contain and is not V to Regulation (EC) No 999/2001, of bovine animals.]]]				
		(¹) or [II.1.9.	.2. for imports from a country or a 2007/453/EC:	region with a controlled BSE risk	and listed as such in Decision			
			 (a) the country or region is classific country or region posing a contribution 	ed in accordance with Article 5(2) of F rolled BSE risk;	Regulation (EC) No 999/2001 as a			

	ſRY	1		Model BO
II.	Health information	1	II.a. Certificate reference number	ll.b.
		stunning by means of gas inject	vine meat or minced meat was derive ted into the cranial cavity or killed by t ntral nervous tissue by means of ar ty;	the same method or slaughtered by
		(¹) either [(c) the bovine meat or minced m defined in Annex V to Regulal bones of bovine animals.]	eat does not contain and is not deriv tion (EC) No 999/2001, or mechanic	
		quarters contain no specified ganglia. The carcasses or w	s or half carcasses cut into no mor risk material other than the vertel vholesale cuts of carcasses of bo d by a blue stripe on the label	oral column, including dorsal root vine animals containing vertebral
	(¹) <i>or</i> [II.1.9.3	 for imports from a country or a region which (EC) No 999/2001 or has been categorised a Decision 2007/453/EC: 		
		 (a) the country or region has not been categorized as a country or region 		of Regulation (EC) No 999/2001 or
		(b) the animals from which the bovine meat of greaves derived from ruminants;	or minced meat was derived have no	t been fed meat-and-bone meal or
		(c) the animals from which the bovine meat o means of gas injected into the cranial c stunning of central nervous tissue by me cavity;	avity or killed by the same method	or slaughtered by laceration after
	(¹) <i>eith</i>	er [(d) the bovine meat or minced meat was no	ot derived from:	
		(i) specified risk material as defined in	Annex V to Regulation (EC) No 999,	/2001;
		(ii) nervous and lymphatic tissues expos	sed during the deboning process;	
		(iii) mechanically separated meat obtaine	ed from bones of bovine animals.]	
	(¹) or	[(d) the carcasses, half carcasses or half car no specified risk material other than the wholesale cuts of carcasses of bovine stripe on the label referred to in Regulat	ne vertebral column, including dorsa animals containing vertebral colum	al root ganglia. The carcasses o
	Pa	ulfils the requirements of Regulation (EC) No 1 rliament and of the Council as regards specia reden of certain meat and eggs;]		
II.2.	Animal Health at	ttestation		
	I, the undersigne	d official veterinarian, hereby certify, that the fre	sh meat described in Part I:	
	II.2.1. has be	een obtained in the territory/ies with code:	(²) which, at	the date of issuing this certificate:
		has been free for 12 months from rinderpest, and place, and	d during the same period no vaccina	tion against this disease has taken
		as been free for 12 months from foot-and-mouth as taken place;]	disease, and during the same period	no vaccination against this disease
		as been considered free from foot-and-mouth di fterwards, and authorised to export this meat by		

COUNT	COUNTRY Model B					
۱۱.	Health info	ormati	on	II.a. Certificate reference number	II.b.	
	(¹) (⁵) or		vaccination programmes against foot-and-mouth animals;]	disease are being officially carried ou	t and controlled in domestic bovine	
	(¹) (⁶) or	herds where the efficacy of this a regular serological surveillance and mouth virus circulation;]				
	(¹)(⁶) or		has been free for 12 months from foot-and-mout has taken place and is controlled by t demonstrating the absence of foot and mouth ir	he competent veterinary authority		
	II.2.2.	has	been obtained from animals that:			
		(¹)	either [have remained in the territory described slaughter;]	d under point II.2.1 since birth, or for a	t least the last three months before	
		(¹)	or [have been introduced on			
		(1)	or [have been introduced on	(dd/mm/yyyy) into the territory descr	ibed under point II.2.1, from the EU	
	II.2.3.	has	been obtained from animals coming from holdi	ings in which:		
		(a)	None of the animals present therein have been	n vaccinated against [foot-and-mouth o	lisease or] (⁷) rinderpest, and	
	(¹) either	[(b)	in these holdings, and in the holdings situated i mouth disease or rinderpest during the previou		been no case/outbreak of foot-and-	
	(¹) (⁸) or	[(b	there is no official restriction for animal health r vicinity within 25 km, there has been no case/ days, and,			
		(c)	they have remained for at least 40 days before	e direct dispatch to the slaughterhouse	ə;]	
	(¹) (¹⁴) or	[(c)	they have remained for at least 40 days bef veterinary authority without coming into conta directly to a slaughterhouse;]			
	(¹) (⁹) or	[(b)	there is no official restriction for animal health r vicinity within 10 km, there has been no case/ months, and			
		(c)	they have remained for at least 40 days before	e direct dispatch to the slaughterhouse	ə;]	
	(1) (6)	[(d)	animals have not been introduced during the la	ast 3 months from areas not approved	I by the EU;	
		(e)	animals are identified and registered in the nation	onal System of Identification and Certif	ication of Origin for bovine animals;	
		(f)	the holdings in question are listed as approve official report, in TRACES (¹⁰) and inspections relevant requirements provided for in Regulatio	are regularly carried out by the comp		
	II.2.4. has	beer	obtained from animals which:			
	(a)		been transported from their holdings in vehicles ut contact with other animals which did not com			

II.	Health	n info	ormatio	n II.a. Certificate reference number II.b.
		(b)		slaughterhouse, have passed ante-mortem health inspection during the 24 hours before slaughter and, in particular, h
			SHOW	The evidence of the diseases referred to in point in.2.1,
		(c)		been slaughtered on (dd/mm/yyyy) or between (dd/mm/yyyy) and nm/yyyy) (¹¹);
	(1) (12) [(d) have reacted negatively to an official intra-dermal tuberculosis test carried out within 3 months before slaughter;]			
	(1) (6)	[(e)	at the the U	slaughterhouse have been kept prior to slaughter completely separate from animals the meat of which is not intended nion].
	II.2.5.	refe imp	erred to ortation	obtained in an establishment around which, within a radius of 10 km, there has been no case/outbreak of the disea o in point II.2.1 during the previous 30 days or, in the event of a case/outbreak of disease, the preparation of meat n to the Union has been authorised only after slaughter of all animals present, removal of all meat, and the total clear ection of the establishment under the control of an official veterinarian;
	II.2.6.			
		(¹)	either	[has been obtained and prepared without contact with other meats not complying with the conditions required in certificate.]
		(¹) (⁸) or	[contains [boneless meat] [and] [minced meat] $(^1)$, obtained only from de-boned meat other than offal that was obtain from carcasses in which the main accessible lymphatic glands have been removed, which have been submitted maturation at a temperature above + 2 °C for at least 24 hours before the bones were removed and in which the value of the meat was below 6.0 when tested electronically in the middle of the longissimus-dorsi muscle a maturation and before de-boning, and
				has been kept strictly separate from meat not conforming to the requirements referred to in this certificate during stages of its production, de-boning and storage until it has been packed in boxes or cartons for further storage dedicated areas.]
		(1)	(⁹) or	[contains [boneless meat] [and] [minced meat] (¹), obtained only from de-boned meat other than offal that was obtain from carcasses in which the main accessible lymphatic glands have been removed, which have been submitted maturation at a temperature above + 2 °C for at least 24 hours before the bones were removed, and
				has been kept strictly separate from meat not conforming to the requirements referred to in this certificate during stages of its production, de-boning and storage until it has been packed in boxes or cartons for further storage dedicated areas.]
1.3.	Anima	al w	elfare a	attestation
				d official veterinarian, hereby certify, that the fresh meat described in Part I derives from animals which have been trea ouse before and at the time of slaughter or killing in accordance with the relevant provisions of Union legislation.
Notes				
	ertificate preeds).		meant	for fresh meat, including minced meat, of domestic bovine animals (including Bison and Bubalus species and the
Fresh	meat m	eans	all ani	imal parts fit for human consumption whether fresh, chilled or frozen.
Part I				
— Box	< referer	nce I	.8: Pro	wide the code of territory as appearing in Part 1 of Annex II to Regulation (EU) No 206/2010.
— Box	< referer	nce I	.11: Pl	ace of origin: name and address of the dispatch establishment.
				egistration number (railway wagons or container and lorries), flight number (aircraft) or name (ship) is to be provided I reloading, the consignor must inform the BIP of entry into the Union.
_	/ referer		10.116	se the appropriate HS code: 02.01, 02.02, 02.06 or 05.04. In addition, for those territories of origin without the entry "A

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

Model OVI

cou	NTRY		Veterinary certificate to		
	l.1.	Consignor	I.2. Certificate reference No I.2.a.		
		Name	I.3. Central competent authority		
		Address			
ant		Tel.	I.4. Local competent authority		
dispatched consignment	l.5.	Consignee	1.6.		
onsi		Name			
ed c		Address			
atch		Postal code			
disp		Tel.			
Part I: Details of	1.7.	Country of origin ISO code I.8. Region of origin Code	B I.9. Country of ISO code I.10. Region of destination Code destination		
rt I: De	l.11.	Place of origin	l.12.		
Pa		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 I Identification	1.17.		
		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 packages		
		Ambient Chilled	Frozen		
	1.23.	Seal/Container No	I.24. Type of packaging		
	1.25.	Commodities certified for:	!		
		Human consumption 🗌			
	1.26.		I.27. For import or admission into EU		
	1.28.	Identification of the commodities	_1		
		Species Nature of Treatment A (scientific name) commodity type Abatto	Approval number of establishments Number of Net packages weight ttoir Cutting plant Cold store		

Part II: Certification

<u>co</u>	COUNTRY Model OVI						
١١.	Hea	alth info	rmatio	วท		II.a. Certificate reference number	ll.b.
11.1	1. Publi	c Heal	th Atl	testation			
	(EC)	No 85	52/200	4, (EC) No	853/2004, (EC) No 854/2004 a	aware of the relevant requirements or and (EC) No 999/2001 and certify that ce with those requirements, in particular	t the meat of domestic ovine and
	II.1.1				eat] (¹) comes from (an) establis ation (EC) No 852/2004;	hment(s) implementing a programme b	pased on the HACCP principles in
	(¹) II.1.2	. the r	neat ł	nas been obl	tained in compliance with the co	onditions set out in Section I of Annex I	II to Regulation (EC) No 853/2004;
	(¹) II.1.3				peen produced in compliance wit not more than – 18 °C;]	h Section V of Annex III to Regulation (EC) No 853/2004 and frozen to an
	II.1.4					ollowing ante and post-mortem inspection n IV of Annex I to Regulation (EC) No 8	
	ll.1.5	. (¹) eit			or parts of the carcass have bee legulation (EC) No 854/2004;]	en marked with a health mark in accorda	ance with Chapter III of Section I of
		(¹) or			os of [meat] [minced meat] (¹) hav Regulation (EC) No 853/2004;]	ve been marked with an identification m	ark in accordance with Section I of
	II.1.6	. the [r foods		[minced mea	at] $(^1)$ satisfies the relevant criter	ia set out in Regulation (EC) No 2073/	2005 on microbiological criteria for
	II.1.7				g live animals and products ther ular Article 29 thereof, are fulfille	reof provided by the residue plans subr d;	mitted in accordance with Directive
	ll.1.8				at] (¹) has been stored and trans I to Regulation (EC) No 853/2004	sported in accordance with the relevant 4;	t requirements of Sections I and V
	II.1.9	. with r	egard	to bovine sp	congiform encephalopathy (BSE):	:	
	(¹) either	[11.1.9.1	. for i	mports from	a country or a region with a neg	ligible BSE risk and listed as such in D	ecision 2007/453/EC:
			(8		y or region is classified in accord negligible BSE risk;	ance with Article 5(2) of Regulation (EC)	No 999/2001 as a country or region
			(t		als from which the meat or mince ith negligible BSE risk; (²)	ed meat was derived were born, continu	uously reared and slaughtered in a
			(¹) [(o) if in the co	ountry or region there have been	BSE indigenous cases:	
				(¹) either	[the animals were born after the meal and greaves derived from	e date from which the ban on the feedin ruminants had been enforced.]	ng of ruminants with meat-and-bone
				(¹) or		s not contain and is not derived from s 999/2001, or mechanically separated me	
	(1) <i>or</i>	[11.1.9	.2. fc	or imports fro	m a country or a region with a c	controlled BSE risk and listed as such ir	1 Decision 2007/453/EC:
			(8		y or region is classified in accorda controlled BSE risk;	ance with Article 5(2) of Regulation (EC)	No 999/2001 as a country or region
			(1	injected in	nto the cranial cavity or killed by	eat was derived have not been slaughte y the same method or slaughtered by rod-shaped instrument introduced into th	laceration after stunning of central

COUNT	RY				Model OVI
П.	Health	information		II.a. Certificate reference number	II.b
		(¹) either	[(c) the meat or minced meat does not conta Regulation (EC) No 999/2001, or mecha animals.]		
		(¹) or	(c) the carcasses, half carcasses or half ca no specified risk material other than the		
	(¹) or	[II.1.9.3.	for imports from a country or a region whicl (EC) No 999/2001 or has been categorised Decision 2007/453/EC:		
			 (a) the country or region has not been categorised as a country or region 		of Regulation (EC) No 999/2001 or
			(b) the animals from which the meat or minor derived from ruminants;	ced meat was derived have not been f	ed meat-and-bone meal or greaves
			(c) the animals from which the meat or mind of gas injected into the cranial cavity or central nervous tissue by means of an example.	killed by the same method or slaught	ered by laceration after stunning of
		(¹) either	[(d) the meat or minced meat was not derive	ed from:	
			(i) specified risk material as defined in	Annex V to Regulation (EC) No 999/2	2001;
			(ii) nervous and lymphatic tissues expo	sed during the deboning process;	
			(iii) mechanically separated meat obtain	ed from bones of domestic ovine or c	aprine animals.]
		(¹) or	[(d) the carcasses, half carcasses or half ca no specified risk material other than the		
II.2.	Animal	Health atte	estation		
	l, the u	ndersigned	official veterinarian, hereby certify, that the fr	resh meat described in Part I:	
	Ⅱ.2.1.	has been	obtained in the territory/ies with code:	(³) which, at the date of iss	uing this certificate:
		(a) has be and	een free for 12 months from rinderpest, and d	uring the same period no vaccination a	gainst this disease has taken place,
	(¹) eithe		een free for 12 months from foot-and-mouth aken place;]	disease, and during the same period	no vaccination against this disease
	(¹) or	break	een considered free from foot-and-mouth dis s afterwards, and authorised to export this n m/yyyy);]		
	(¹) (⁴) <i>or</i>	[(b) vaccir anima	nation programmes against foot-and-mouth c ıls;]	lisease are being officially carried out	and controlled in domestic bovine
	II.2.2.	has been	obtained from animals that:		
		(¹) either	[have remained in the territory described using territory described using territory described using the second sec	under point II.2.1 since birth, or for at	least the last three months before
		(¹) or	[have been introduced on territory with code (³) that at that date		
		(¹) or	[have been introduced on	.(dd/mm/yyyy) into the territory descrit	bed under point II.2.1, from the EU

NTRY Health	n information	II.a. Certificate reference number	Model O	
			1.0.	
11.2.3.	has been obtained from animals coming from holdings	3:		
	(a) in which none of the animals present therein have	been vaccinated against [foot-and-mouth	n disease or] (⁵) rinderpest,	
	(b) not subject to prohibition as a result of an outbrea	k of 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, during the previous 30 days;]	there has been no case/outbreak of for	ot-and-mouth disease or rinderpes	
(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 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 be	fore direct dispatch to the slaughterhouse	ə;]	
(¹) (⁸) or	[(d) where they have remained for at least 40 days veterinary authority without coming into contact wit a slaughterhouse;]			
II.2.4.	has been obtained from animals which:			
	 (a) have been transported from their holdings in vehic without contact with other animals which did not contact. 			
	(b) at the slaughterhouse, have passed ante-mortem h shown no evidence of the diseases referred to in		re slaughter and, in particular, hav	
	(c) have been slaughtered on (dd/mm/yy	yy) or between (dd/mm/yyyy) and(dd/mm/yyyy) (⁶)	
II.2.5.	has been obtained in an establishment around which, referred to in point II.2.1 during the previous 30 days importation into the Union has been authorised only after and disinfection of the establishment under the control	or, in the event of a case/outbreak of di er slaughter of all animals present, remova	isease, the preparation of meat fo	
II.2.6.				
(¹) either	[has been obtained and prepared without contact with	n other meats not complying with the cor	nditions required in this certificate	
(¹)(⁴) or	[contains [boneless meat] [and] [minced meat] $(^1)$, o carcasses in which the main accessible lymphatic g temperature above + 2 °C for at least 24 hours before 6.0 when tested electronically in the middle of the lo	lands have been removed, which have the 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 confo production, de-boning and storage until it has been p			
(¹)(⁷) or	[contains [boneless meat] [and] [minced meat] (¹), o carcasses in which the main accessible lymphatic g temperature above + 2 °C for at least 24 hours befor	lands have been removed, which have		
	has been kept strictly separate from meat not confo production, de-boning and storage until it has been p			
Animal	welfare attestation			
	dersigned official veterinarian, hereby certify, that the fre ghterhouse before and at the time of slaughter or killing			

OUNTRY Model OVI					
II. Health information	II.a. Certificate reference number	II.b.			
Notes		<u></u>			
This certificate is meant for fresh meat, including minced meat, of do	mestic ovine animals (<i>Ovis aries</i>) an	d caprine animals (Capra hircus).			
Fresh meat means all animal parts fit for human consumption whether fr	resh, chilled or frozen.				
Part I:					
- Box reference I.8: Provide the code of territory as appearing in Part	1 of Annex II to Regulation (EU) No 2	206/2010.			
- Box reference I.11: Place of origin: name and address of the dispatc	h establishment.				
 Box reference I.15: Registration number (railway wagons or container case of unloading and reloading, the consignor must inform the BIP 		r 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 ar	nd the seal number (if applicable) sho	uld be included.			
 Box reference I.28: Nature of commodity: Indicate "carcass-whole", "c meat is de-boned meat that has been minced into fragments and that adjoining fatty tissues) except heart muscle. 					
 Box reference I.28: Treatment type: If appropriate, indicate "de-bone freezing (mm/yy) of the cuts/pieces. 	əd"; 'bone in"; "matured" and/or "minc	ed". If frozen, indicate the date of			
Part 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 o	olumn 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, territory 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-b days after the date of slaughter of the animals.	neat to be provided when required in o coned meat shall not be authorised for	oolumn 5 "SG" of Part 1 of Annex II r importation into the Union until 21			
(⁸) Alternative guarantee may be provided when allowed for by the (EU) No 206/2010.	∍ entry " J " in column 5 "SG" of F	art 1 of Annex II to Regulation			
Official veterinarian					
Name (in capital letters):	Qualification and title:	:			
Date:	Signature:				
Stamp:					

▼<u>C1</u>

		lel POR			
	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			
ent	Tel. No				
gnm	I.5. Consignee	1.6.			
onsi	Name				
od co	Address				
tche	Postal code				
ispa	Tel. No				
Part I: Details of dispatched consignment	I.7. Country ISO I.8. Region Code of origin code of origin	I.9. Country of ISO I.10. Region of Code destination code destination			
Deta	I.11. Place of origin	1.12.			
irt I:	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:				
	I.18. Description of commodity	I.19. Commodity code (HS code)			
		I.20. Quantity			
	I.21. Temperature of product	I.22. Number of packages			
	Ambient Chiled	Frozen			
	I.23. Identification of container/seal number	I.24. Type of packaging			
	I.25. Commodities certified for:				
	Human consumption				
	l.26.	I.27. For import or admission into EU			
	I.28. Identification of the commodities				
	Species Nature of Treatment Ap (Scientific name) commodity type	proval number establishments Number Net of packages weight			
	(Scientine name) commounty type Abath				

Model POR

	COU	INTRY				Model POR
	П.	Health	information		II.a. Certificate reference number	II.b.
	II.1.	Public	Health Attesta	ition		
Part II: Certification		(EC) N	o 852/2004, (E0	C) No 853/2	rian, declare that I am aware of the relevant re 2004 and (EC) No 854/2004 and hereby certi ance with those requirements, in particular the	fy that the meat of domestic swine described
fication		II.1.1] (¹) comes from (an) establishment(s) implei with Regulation (EC) No 852/2004;	nenting a programme based on the HACCP
t II: Certi		II.1.2	the meat has No 853/2004;	been obtai	ned in compliance with the conditions set or	t in Section I of Annex III to Regulation (EC)
Part		II.1.3	the meat fulfils <i>Trichinella</i> in n		2 ()	ing down specific rules on official controls for
			(1) either	[has bee	n subjected to an examination by a digestion	n method with negative results]
			(1) or	[has bee No 2075	en subjected to a freezing treatment in ac /2005;]	cordance with Annex II to Regulation (EC)
			(1) <i>or</i>	holding of	ase of meat from domestic swine kept sole or category of holdings that has been official In Trichinella in accordance with Annex IV to R	ly recognized by the competent authority as
		(¹) II.1.4			en produced in accordance with Section V of h berature of not more than –18 °C;]	Annex III to Regulation (EC) No 853/2004 and
		II.1.5				and post-mortem inspections carried out in f Section IV of Annex I to Regulation (EC)
		II.1.6 (¹) either	-	cass or parts of the carcass have been mai III of Section I of Annex I to Regulation (EC) N	
			(1) <i>or</i>		kages of [meat] [minced meat] (1) have be nce with Section I of Annex II to Regulation (E	een marked with an identification mark in C) No 853/2004;]
		II.1.7	the [meat] [mir criteria for food	-	(1) satisfies the relevant criteria set out in Reg	ulation (EC) No 2073/2005 on microbiological
		II.1.8			live animals and products thereof provided b and in particular Article 29, are fulfilled.	y the residue plans submitted in accordance
		II.1.9			t] (¹) has been stored and transported in ac vely of Annex III to Regulation (EC) No 853/2	cordance with the relevant requirements of 004.
		(²) [II.1.10			of Regulation (EC) No 1688/2005 implement erning Salmonella for consignments to Finlan	
	II.2.	Anima	l Health attesta	ation		
		I, the u	ndersigned offic	cial veterina	arian, hereby certify, that the fresh meat desc	ibed in Part I :
		II.2.1	has been obta	ined in the	territory/ies with code: (³) which, at the date of issuing this certificate:
			(1) either	,	been free for 12 months from foot-and-mot sical swine fever, swine vesicular disease, and	
			(1) or		nas been free for 12 months from rinderpest, Africal swine fever] (') and [swine vesicular	

	Health	information		II.a. Certificate reference number	II.b.
				[swine vesicular disease] (1), since	outh disease] (¹), [classical swine fever] (¹) an
				orts of domestic animals vaccinated agai	nst these diseases have been carried out an nst these diseases are not permitted in thi
	II.2.2	has been ob	tained from	animals that:	
		(1) either		mained in the territory described under poin before slaughter;]	nt II.2.1 since birth, or for at least the last thre
		(1) or	point II.2		dd/mm/yyyy) into the territory described unde
		(1) <i>or</i>		een introduced on(2.1, from the EU Member State	dd/mm/yyyy) into the territory described unde
	II.2.3	has been ob	tained from	animals coming from holdings:	
		(a) in which point II.2		ne animals present therein have been va	accinated against the diseases referred to i
				, in an area of 10 km radius, there has been e previous 40 days,	no case/outbreak of the diseases referred to i
		(c) that are weeks;	not subject	to prohibition as a result of an outbreak	of porcine brucellosis during the previous si
	(1) (4)				th catering waste, are subject to official control ty for the purpose of importing pig meat into th
	II.2.4	has been ob	tained from	animals that:	
		(a) have rer	nained sepa	rate since birth from wild cloven-hoofed ani	mals,
			rhouse with		and disinfected before loading, to an approve comply with the conditions set out in points II.2.
				e, have passed ante-mortem health inspect vn no evidence of the diseases referred to ir	ion during the 24 hours before slaughter and, i n point II.2.1, and
				red on (dd/mm/yyyy) د (dd/mm/yyyy). (⁵);	or between (dd/mm/yyy)
	II.2.5	of the disea preparation	ses referred of meat for in all meat, and	I to in point II.2.1 during the previous 40 d mportation into the Union has been authori	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 presen establishment under the control of an officia
	II.2.6	has been ob certificate.	tained and p	prepared without contact with other meats no	ot complying with the conditions required in thi
.3.	Anima	I welfare atte	station		
					cribed in Part I derives from animals which hav Iling in accordance with the relevant provision

со	COUNTRY Model POF								
Ш.	Health information	II.a. Certificate reference number	II.b.						
No	Notes								
Thi	This certificate is meant for fresh meat, including minced meat, of domestic swine (Sus scrofa).								
Fre	sh meat means all animal parts fit for hur	man consumption whether fresh, chilled or fro	ozen.						
Pai	rt I:								
_	Box reference I.8: Provide the code of te	erritory as appearing in Part 1 of Annex II to R	egulation (EU) No 206/2010.						
_	Box reference I.11: Place of origin: name	e and address of the dispatch establishment.							
_	5	r (railway wagons or container and lorries), fl 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 15.0	1.						
—	Box reference I.20: Indicate total gross v	•							
—		xes, the container number and the seal numb	, , ,						
_	-	y: Indicate 'carcass-whole', 'carcass-side', 'ca							
	Minced meat is deboned meat that has muscle (including the adjoining fatty tiss	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						
Pai	rt II:								
(1)	Keep as appropriate.								
(²)	Delete if the consignment is not intende	d for import into Finland or Sweden.							
(³)	Code of the territory as it appears in Par	rt 1 of Annex II to Regulation (EU) No 206/201	0.						
(4)	Supplementary guarantees to be provid with the entry ' D '.	ded when required in column 5 'SG' of Part 1	of Annex II to Regulation (EU) No 206/2010,						
	Catering waste means: all waste from foc industrial kitchens and household kitchen	od intended for human consumption from resta ns of the farmer or persons tending pigs.	urants, catering facilities or kitchens, including						
(5)	of authorisation for importation into the L	s meat shall not be allowed when obtained fro Jnion of the third country, territory or part there been adopted by the Union against imports o	eof referred to in boxes I.7 and I.8, or during a						
Off	icial veterinarian								
	Name (in capital letters):	Qualificatio	n and title:						
	Date:	Signature:							
	Stamp:								
	ownp.								

	COUNTRY	Mode	el EQU	Veterinary certificate to EU			
	I.1. Consignor		I.2. Certificate reference				
	Name		and the second composition of a produce a story of the second				
	Address		I.3. Central Competent Authority				
Ŧ	Tel. No		I.4. Local Competent Authority				
men	I.5. Consignee		1.6.				
sign	Name						
con	Address						
hed	Postal code						
oatc	Tel. No						
Part I: Details of dispatched consignment		I.8. Region Code of origin	,	SO I.10. Region of Code ode destination			
etai	I.11. Place of origin		I.12.				
t: D	-	Approval number					
Par	Address						
·	110 Disco of loading						
	I.13. Place of loading		I.14. Date of departure				
	I.15. Means of transport Aeroplane	Railway wagon	I.16. Entry BIP in EU				
	Road vehicle Other						
	Identification: Documentary references:		1.17.				
	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				
	I.23. Identification of container/sea	l number		I.24. Type of packaging			
-	I.25. Commodities certified for: Human consumption						
	1.26.		I.27. For import or admissi	ion into EU			
	I.28. Identification of the commodit	ies	1				
		ture of Approval กเ Imodity	umber establishments	Number Net of packages weight			
		Abattoir C	Cutting plant Cold store				

COUN	IRT			Model EQ					
П.	Health	information	II.a. Certificate reference number	II.b.					
II.1.	Public Health Attestation								
	(EC) N	o 852/2004, (E	cial veterinarian, declare that I am aware of the relevant re C) No 853/2004 and (EC) No 854/2004 and hereby certif I in accordance with those requirements, in particular th	y that the meat of domestic solipeds described					
	II.1.1		nes from (an) establishment(s) implementing a prog vith Regulation (EC) No 852/2004;	ramme based on the HACCP principles in					
	II.1.2	the meat has No 853/2004;	been obtained in compliance with the conditions set o	ut in Section I of Annex III to Regulation (EC)					
	II.1.3		Is the requirements of Regulation (EC) No 2075/2005 in meat, and in particular, has been subject to an exar						
	II.1.4		been found fit for human consumption following ante with Chapter II of Section I and Chapters III and IX o						
_	II.1.5	(1) either	[the carcass or parts of the carcass have been ma Chapter III of Section I of Annex I to Regulation (EC)						
		(1) or	[the packages of meat have been marked with an ider Annex II to Regulation (EC) No 853/2004;]	ntification mark in accordance with Section I of					
	II.1.6	the meat sati foodstuffs;	isfies the relevant criteria set out in Regulation (EC)	No 2073/2005 on microbiological criteria for					
	II.1.7		es covering live animals and products thereof provided l 96/23/EC, and in particular Article 29 thereof, are fulfille						
	II.1.8		been stored and transported in accordance with the rel C) No 853/2004.	evant requirements of Section I of Annex III to					
II.2.	Anima	I Health attest	ation						
	l, the u	ndersigned offi	cial veterinarian, hereby certify, that the fresh meat desc	cribed in Part I:					
	II.2.1	has been obta	ained in the territory/ies with code:	(²);					
	II.2.2	has been obta	ained from domestic solipeds, which:						
		(1) either	[have remained in the territory described under poin months before slaughter;]	t II.2.1 since birth, or for at least the last three					
		(1) or	[have been introduced on(depoint II.2.1, from the territory with code:						
		(1) <i>or</i>	[have been introduced on(de point II.2.1, from the EU Member State						
	II.2.3	which, within previous 40 d has been aut	tained from animals which were slaughtered on (dd/mm/yyyy) and	dd/mm/yyyy) (³) in a slaughterhouse around African horse sickness or glanders during the paration of meat for importation into the Union moval of all meat, and the total cleaning and					

COUNTRY

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

				Mod	el RUF					
	со	UNTRY						Veterinary certifi	cate to EU	
	l.1.	Consignor			I.2. Certifica	ate reference	numbe	er I.2.a.		
		Name			I.3. Central Competent Authority					
		Address								
ent		Tel. No			I.4. LOCAIC	ompetent Au	thority			
un	l.5.	Consignee			1.6.					
nsiç		Name								
d co		Address								
tche		Postal code								
spat		Tel. No								
Part I: Details of dispatched consignment	1.7.	Country ISO of origin code	I.8. Region of origin	Code	I.9. Country destina		ISO code	I.10. Region of destination	Code	
Deta	l.11.	. Place of origin			I.12.		I			
Ξt		Name	Approval number							
Ра		Address								
	I.13	. Place of loading			I.14. Date of	departure				
	145	Maana of transport								
	1.15	. Means of transport Aeroplane S	hip 🗌 🛛 Railway wago	on 🗌	I.16. Entry BIP in EU					
			ner							
		Identification:			l.17.					
		Documentary references:								
	I.18	. Description of commodity			I.19. Commodity code (HS code)					
							1.00	Quantity		
							1.20.1	Quantity		
	I.21	. Temperature of product					1.22.1	Number of packages		
		Ambient	Chiled		Frozen]				
						_				
	1.23	B. Identification of container/	seal number				1.24.	Type of packaging		
	I.25	5. Commodities certified for: Human consumption								
	1.26).			I.27. For import or admission into EU					
	1.28	B. Identification of the comm	odities		1					
		Species Natu		Арр	roval number e	establishment	ts	Number	Net	
	(5	Scientific name) comm	odity type	Abotto	ir Cutting r	Nont Cold	otoro	of packages	veight	
				Abatto	ir Cutting p	olant Cold	SIDIE			

	COUNT	RY						Model RUF
	Ш.	Health	information		II.a. Certificate referenc	e number	II.b.	
	II.1.	Public	Health Attest	ation				
ation		No 178 the me and th	3/2002, (EC) Neat of farmed a eir cross-bree	lo 852/2004 Inimals of t ds), <i>Ovis a</i>	rrinarian, declare that I a 4, (EC) No 853/2004, (EC he order Artiodactyla (ex <i>ries, Capra hircus,</i> Suida I was produced in accord	C) No 854/2004 and cluding bovine anim ae and Tayassuidae	l (EC) No 999/2001 an nals (including <i>Bison a</i> e), and of the families	nd hereby certify that and <i>Bubalus</i> species Rhinocerotidae and
Part II: Certification		II.1.1			an) establishment(s) imp ion (EC) No 852/2004;	lementing a progra	amme based on the	HACCP principles in
Part II		II.1.2	the meat has No 853/2004		ned in accordance with th	e conditions set out	in Section III of Annex	III to Regulation (EC)
		II.1.3		with Chapte	d fit for human consump er II of Section I and Cha			
		II.1.4	(1) either		cass or parts of the carca III of Section I of Annex I to			k in accordance with
			(1) or		kages of meat have be I of Annex II to Regulation			in accordance with
		II.1.5	the meat sat foodstuffs;	isfies the r	elevant criteria set out in	Regulation (EC) N	o 2073/2005 on micro	obiological criteria for
		II.1.6			live animals and products and in particular Article 29			omitted in accordance
	(1) (²) [ll.1.7	with regard to	Chronic W	asting Disease (CWD):			
			animals whic other diagno	h have bee stic method	or is derived exclusively t en examined for Chronic I recognised by the com lerd where Chronic Wastin	Wasting Disease by petent authority with	/ histopathology, imm h negative results and	unohistochemistry or dis not derived from
		II.1.8	the meat has Regulation (E		d and transported in accor 2004.	rdance with the relev	vant requirements of S	ection I of Annex III to
	II.2.	Anima	I Health attes	tation				
		l, the u	ndersigned off	icial veterina	arian, hereby certify, that th	ne fresh meat descri	bed in Part I:	
		II.2.1	has been obt	ained in the	territory/ies with code:	(3)	which, at the date of i	ssuing this certificate:
				free for 12 place, and	months from rinderpest, a	and during the same	e period no vaccination	n against this disease
	(¹) either		free for 12 ise has take	months from foot-and-mo n place;]	uth disease, and du	ring the same period n	no vaccination against
	([1) or	having ha	d cases/ou	d free from foot-and-mou tbreaks afterwards, and au (dd/mm/y	uthorised to export th		
	((¹) (⁴) or		on program bovine anir	mes against foot-and-mo nals;]	outh disease are be	eing officially carried	out and controlled in

co	UNTRY			Model RUF
II.	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 th 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 i 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 break 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);]	n/yyyy) or between
	(1) or		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 de shown no evidence of the diseases referred	
		 the animals were (dd/mm/yyyy), (⁶) 	slaughtered between	(dd/mm/yyyy) and
		 the bleeding of th 	e animals was performed correctly, and	
		 the slaughtered a 	nimals were eviscerated within three hours of	f 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) (7) 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	information		II.a. Certificate reference number	II.b.
	II.2.6	of the disea preparation	ses referrec of meat for i all meat, and	d to in point II.2.1 during the previous 30 importation into the Union has been autho	dius of 10 km, there has been no case/outbrea days or, in the event of a case of disease, th prised only after slaughter of all animals preser e establishment under the control of an offici
	II.2.7				
		(1) either	-	en obtained and prepared without contact w d above.]	vith other meats not complying with the condition
		(1) (4) or	carcass submitte removed	es in which the main accessible lymphatic ed to maturation at a temperature above +	oned meat other than offal that was obtained fro c glands have been removed, which have bee 2 °C for at least 24 hours before the bones we was below 6.0 when tested electronically in the ration and before de-boning, and
			certifica		conforming to the requirements set out in th boning and storage until it has been packed reas.]
		(1) (8) or	carcass	es in which the main accessible lymphatic ed to maturation at a temperature above +	oned meat other than offal that was obtained fro c glands have been removed, which have bee 2 °C for at least 24 hours before the bones we
			certifica		conforming to the requirements set out in th boning and storage until it has been packed reas.]
This o anima	certificate is als (includin	g <i>Bison</i> and <i>B</i>	<i>ubalus</i> spec	•	apra hircus, Suidae and Tayassuidae), and of th
This c anima amilio	certificate is als (includin es Rhinocei	g <i>Bison</i> and <i>B</i> rotidae and Ele	<i>ubalus</i> spec ephantidae,	cies and their cross-breeds), Ovis aries, Ca	apra hircus, Suidae and Tayassuidae), and of th rth or for the last three months in farms.
This c anima amilie Fresh	certificate is als (includin es Rhinocei n meat mear	g <i>Bison</i> and <i>B</i> rotidae and Ele	<i>ubalus</i> spec ephantidae,	cies and their cross-breeds), Ovis aries, Ca that are domestically kept or bred since bin	apra hircus, Suidae and Tayassuidae), and of th rth or for the last three months in farms.
This c anima amilic Fresh Part I	certificate is als (includin es Rhinoce n meat mear I:	g <i>Bison</i> and <i>B</i> rotidae and Ele ns all animal pa	<i>ubalus</i> spec ephantidae, arts fit for hu	cies and their cross-breeds), Ovis aries, Ca that are domestically kept or bred since bi iman consumption whether fresh, chilled o	apra hircus, Suidae and Tayassuidae), and of th rth or for the last three months in farms. r frozen.
This c anima amilio Fresh Part I — B	certificate is als (includin es Rhinocei n meat mear n meat mear l:	g <i>Bison</i> and <i>B</i> rotidae and Ele ns all animal pa e I.8: Provide t	<i>ubalus</i> spec ephantidae, arts fit for hu he code of t	cies and their cross-breeds), Ovis aries, Ca that are domestically kept or bred since bi iman consumption whether fresh, chilled o rerritory as appearing in Part 1 of Annex II to	apra hircus, Suidae and Tayassuidae), and of th rth or for the last three months in farms. r frozen. o Regulation (EU) No 206/2010.
This c anima amilia Fresh Part I — B — B	certificate is als (includin es Rhinocei n meat mear n meat mear l: cox reference cox reference cox reference	g <i>Bison</i> and <i>B</i> rotidae and Ele ns all animal pa e I.8: Provide t e I.11: Place o e I.15: Registra	ubalus spec ephantidae, arts fit for hu he code of t f origin: nam ation numbe	cies and their cross-breeds), Ovis aries, Ca that are domestically kept or bred since bi iman consumption whether fresh, chilled o cerritory as appearing in Part 1 of Annex II to he and address of the dispatch establishme er (railway wagons or container and lorries	apra hircus, Suidae and Tayassuidae), and of th rth or for the last three months in farms. or frozen. o Regulation (EU) No 206/2010. ent. s), flight number (aircraft) or name (ship) is to b
This c anima amilia 	certificate is als (includin es Rhinocei n meat mear neat mear l: lox reference lox reference rovided. In c	g <i>Bison</i> and <i>B</i> rotidae and Ele ns all animal pa e I.8: Provide t e I.11: Place o re I.15: Registra case of unload	ubalus spec ephantidae, arts fit for hu he code of t f origin: narr ation numbe ing and relo	cies and their cross-breeds), Ovis aries, Ca that are domestically kept or bred since bi iman consumption whether fresh, chilled o rerritory as appearing in Part 1 of Annex II to he and address of the dispatch establishme er (railway wagons or container and lorries rading, the consignor must inform the BIP o	apra hircus, Suidae and Tayassuidae), and of th rth or for the last three months in farms. or frozen. o Regulation (EU) No 206/2010. ent. s), flight number (aircraft) or name (ship) is to b
This c anima amilid - Fresh - B - B p - B	certificate is als (includin es Rhinocei n meat mear l: lox referenc lox referenc lox referenc rovided. In c	g <i>Bison</i> and <i>B</i> rotidae and Ele ns all animal pa e I.8: Provide t e I.11: Place o e I.15: Registra case of unload e I.19: Use the	ubalus spec ephantidae, arts fit for hu he code of t f origin: nam ation numbe ing and relo e appropriate	cies and their cross-breeds), Ovis aries, Ca that are domestically kept or bred since bin iman consumption whether fresh, chilled o rerritory as appearing in Part 1 of Annex II to he and address of the dispatch establishme er (railway wagons or container and lorries rading, the consignor must inform the BIP o e HS code: 02.06, 02.08.90 or 05.04.	apra hircus, Suidae and Tayassuidae), and of th rth or for the last three months in farms. or frozen. o Regulation (EU) No 206/2010. ent. s), flight number (aircraft) or name (ship) is to b
This c anima amiliu Fresh - B - B - B - B - B	certificate is als (includin es Rhinocei n meat mear n meat mear l: lox referenc lox referenc rovided. In c lox referenc lox referenc	g <i>Bison</i> and <i>B</i> rotidae and Ele ns all animal pa e I.8: Provide t e I.11: Place o e I.15: Registra case of unload e I.19: Use the e I.20: Indicate	ubalus spec ephantidae, arts fit for hu he code of t f origin: nam ation numbe ing and relo e appropriate e total gross	cies and their cross-breeds), Ovis aries, Ca that are domestically kept or bred since bin iman consumption whether fresh, chilled o remitter as appearing in Part 1 of Annex II to he and address of the dispatch establishme er (railway wagons or container and lorries rading, the consignor must inform the BIP of the Scode: 02.06, 02.08.90 or 05.04. weight and total net weight.	apra hircus, Suidae and Tayassuidae), and of the rth or for the last three months in farms. or frozen. o Regulation (EU) No 206/2010. ent. s), flight number (aircraft) or name (ship) is to b of entry into the Union.
anima familia Fresh — B — B — B — B — B — B	certificate is als (includin es Rhinocei n meat mear n meat mear cox reference lox reference rovided. In co cox reference lox reference lox reference lox reference	g <i>Bison</i> and <i>B</i> rotidae and Ele ns all animal pa e I.8: Provide t e I.11: Place o e I.15: Registra case of unload e I.19: Use the e I.20: Indicate e I.23: For con	ubalus spec ephantidae, arts fit for hu he code of t f origin: nam ation numbe ing and relo e appropriate e total gross tainers or bo	cies and their cross-breeds), Ovis aries, Ca that are domestically kept or bred since bin iman consumption whether fresh, chilled o rerritory as appearing in Part 1 of Annex II to he and address of the dispatch establishme er (railway wagons or container and lorries rading, the consignor must inform the BIP o e HS code: 02.06, 02.08.90 or 05.04.	r frozen. o Regulation (EU) No 206/2010. ent. s), flight number (aircraft) or name (ship) is to b of entry into the Union. umber (if applicable) should be included.

CC	DUNTRY		Model RUF						
II.	Health information	II.a. Certificate reference number	II.b.						
Ра	rt II:								
(1) (2)	 ²) Supplementary guarantees regarding fresh meat obtained from cervids to be provided when required in column 5 'SG' of Part 1 of Annex II to Regulation (EU) No 206/2010, with the entry 'G'. 								
• •		art 1 of Annex II to Regulation (EU) No 206/201							
(')	Part 1 of Annex II to Regulation (EU)) meats from matured de-boned meat to be p No 206/2010 with the entry ' A '.	rovided when required in column 5 SG of						
(5)		rries out vaccination against foot-and-mouth Inion matured de-boned meat which fulfils the							
(6)	date of authorisation for importation in during a period where restrictive meas territory or part thereof.	his meat shall not be authorised when obtained to the Union of the third country, territory or pa ures have been adopted by the Union against	art thereof referred to in boxes I.7 and I.8, or						
(7)	Not necessary for farmed game anima								
(°)		neats from matured de-boned meat to be provi 2010, with the entry 'F'. The matured de-boned tte of slaughter of the animals.							
Of	ficial veterinarian								
	Name (in capital letters):	Qualification	and title:						
	Date:	Signature:							
	Stamp:								

					Mode	el RUW					
									Veterinary certif	icate to EU	
	1.1.	Consignor				I.2. Certific	ate reference	number	l.2.a.		
		Name				I.3. Central Competent Authority					
		Address				I.4. Local Competent Authority					
nent	1.5	Tel. No									
ignr	1.5.	Consignee				1.6.					
suos		Name									
led o		Address									
atch		Postal code									
disp	. 7	Tel. No	100		Quala			00		Quila	
Part I: Details of dispatched consignment	1.7.	Country of origin	ISO code	I.8. Region of origin	Code	I.9. Countr destina		SO ode	I.10. Region of destination	Code	
: Del	I.11.	I.11. Place of origin					1.12.				
art I		Name Address		Approval number							
٩.		, luar occ									
	I.13	. Place of loading				I.14. Date of	departure				
	l.15	. Means of transpo Aeroplane		ip 🗌 🛛 Railway wag	on 🗖	I.16. Entry B	IP in EU				
		Road vehicle		er 🗌							
			Our			1.17.					
		Identification: Documentary ref	erences:								
	I.18	. Description of co	mmodity			I.19. Commodity code (HS code)					
								l.20. Q	Quantity		
	I.21	. Temperature of p	roduct					I.22. N	lumber of packages		
		Ambient		Chiled		Frozen					
	1.23	B. Identification of c	ontainer/s	eal number				I.24. T	ype of packaging		
	I.25	i. Commodities cer Human consump									
	1.26).				I.27. For imp	oort or admissi	on into E	EU [
						roval number e	establishment	S	Number	Net	
	(3	Scientific name)	commo	odity type	Abatto	ir Cutting	plant Cold	store	of packages	weight	
					Αυαιίυ			5.018			

Model RUW

	COUNTRY			Model RUW			
	II. Healt	h information	II.a. Certificate reference number	II.b.			
ation	II.1. Publi	c Health Attestation					
	No 1 anima <i>Ovis</i>	78/2002, (EC) No 852/2004 als of the order Artiodactyla <i>aries, Capra hircus,</i> Suida	erinarian, declare that I am aware of the re I, (EC) No 853/2004 and (EC) No 854/2004 a (excluding bovine animals (including <i>Bison</i> ar e and Tayassuidae), and of the families Rhin ince with those requirements, in particular that:	nd hereby certify that the fresh meat of wild ad <i>Bubalus</i> species and their cross-breeds), occrotidae and Elephantidae described in			
Part II: Certification	II.1.1	the meat comes from accordance with Regula	(an) establishment(s) implementing a progra tion (EC) No 852/2004;	amme based on the HACCP principles in			
Part II:	II.1.2	the meat has been obta 853/2004, and in particu	ined in compliance with the conditions set out in Section IV of Annex III to Regulation ar:				
		(i) before skinning, it ha	s been stored and handled separately from ot	her food and not frozen;			
		and					
		(ii) after skinning, it has	undergone a final inspection as referred to in p	point II.1.4;			
	(¹) II.1.3		le species, the meat fulfils the requirements of controls for Trichinella in meat;]	Regulation (EC) No 2075/2005 laying down			
	II.1.4		d fit for human consumption following a post-m I and Chapters VIII and IX of Section IV of An				
	II.1.5		ase of large wild game, the carcass or parts of accordance with Chapter III of Section I of Ann				
		., .	kages of meat have been marked with an identi to Regulation (EC) No 853/2004;]	fication mark in accordance with Section I of			
	II.1.6	the meat satisfies the r foodstuffs;	elevant criteria set out in Regulation (EC) N	o 2073/2005 on microbiological criteria for			
	II.1.7	0 0	ng live animals and products thereof provided by the residue plans submitted in accordance C, and in particular Article 29 thereof, are fulfilled. Wasting Disease (CWD):				
	(1) (2) [II.1.8	with regard to Chronic W					
		have been examined fo method recognised by th	is derived exclusively from meat, excluding offal r Chronic Wasting Disease by histopathology, le competent authority with negative results an asting Disease has been confirmed in the last t	immunohistochemistry or other diagnostic dis not derived from animals coming from a			
	II.1.9	the meat has been store Regulation (EC) No 853/	d and transported in accordance with the relev 2004.	vant requirements of Section I of Annex III to			
	II.2. Anim	al Health attestation					
	I, the	undersigned official veterin	arian, hereby certify, that the fresh meat descri	bed in Part I:			
	II.2.1	has been obtained in the	e territory/ies with code:	which, at the date of issuing this certificate:			
		 (a) has been free for 12 has taken place, and 	months from rinderpest, and during the same I	e period no vaccination against this disease			
	(1) either	(b) has been free for 12 this disease has take	months from foot-and-mouth disease, and du an place;]	ring the same period no vaccination against			
l							

.

					-
I. Health	n information	II.a. Certific	cate reference number	II.b.	
(1) or	having h	had cases/outbreaks afte		ce (dd/mm/yyyy), wit export these animals by Commission Regula	
(1) (4) or		ion programmes against c bovine animals;]	t foot-and-mouth disease	are being officially carried out and controlle	ed i
II.2.2					
		nce that exceeds 20 km fr r importing this fresh mea		r or part thereof, which is not authorised during	; th
	(b) in an ar point II.2		t 60 days, there has bee	n no restrictions for the diseases referred t	to i
II.2.3	game-handlin diseases refe of meat for in	ng establishment around erred to in point II.2.1 duri aportation into the Union I	which, within a radius of ng the previous 30 days or	ed as soon as possible for chilling to an appro 10 km, there has been no case/outbreak of in the event of a case of disease, the prepara ter removal of all meat, and the total cleaning terinarian;	f th atio
II.2.4					
	(1) either	[has been obtained an required above.]	d prepared without contact	with other meats not complying with the condit	ior
	(¹) (⁴) or	carcasses in which th submitted to maturation removed and in which	e main accessible lympha on at a temperature above n the pH value of the mea	boned meat other than offal that was obtained tic glands have been removed, which have be ± 2 °C for at least 24 hours before the bones v was below 6.0 when tested electronically in uration and before de-boning, and	ve
		certificate during all s		conforming to the requirements set out in -boning and storage until it has been packe areas.]	
	(1) (6) or	carcasses in which th	e main accessible lympha	oned meat other than offal that was obtained tic glands have been removed, which have t +2 °C for at least 24 hours before the bones w	bee
		certificate during all s		conforming to the requirements set out in e-boning and storage until it has been packe areas.]	
lotes					
nimals (includir	ng <i>Bison</i> and <i>Bi</i>		cross-breeds), Ovis aries, (nimals of the order Artiodactyla (excluding bo Capra hircus, Suidae and Tayassuidae), and o	
			ption whether fresh, chilled	or frozen.	
	, unskinned ca				

со	COUNTRY Model RUV									
П.	Health information	II.a. Certificate reference number	II.b.							
	Part 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.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: Nature of commodity: Indicate 'carcass-whole', 'carcass-side', 'carcass-guarters' or 'cuts'. 									
—		opropriate, indicate 'matured' or 'unsk	kinned'. If frozen, indicate the date of freezing (mr	m/yy)						
Par	t II:									
(²) (³)	of Annex II to Regulation (EU) No 206 Code of the territory as it appears in Pa	5/2010, with the entry ' G '. rt 1 of Annex II to Regulation (EU) No								
.,	Part 1 of Annex II to Regulation (EU)	No 206/2010 with the entry 'A'.	at to be provided when required in column 5 'S							
	The matured de-boned meat shall not animals.	be authorised for importation into the	the Union until 21 days after the date of killing c	of the						
	for importation into the Union of the thi	rd country, territory or part thereof ref	Is killed or hunted either prior to the date of authoris ferred to in boxes I.7 and I.8, or during a period w s meat from this third country, territory or part ther	vhere						
(6)	Supplementary guarantees regarding m Annex II to Regulation (EU) No 206/20 the Union until 21 days after the date of	eats from matured de-boned meat to 10, with the entry ' F '. The matured de	be provided when required in column 5 'SG' of Par e-boned meat shall not be allowed for importation	rt 1 of						
Offi	cial veterinarian									
	Name (in capital letters):		ualification and title:							
	Date: Stamp:	Si	gnature:							

		.,			Mod	el SUF						
	COUNTR									Veterinary certif	ficate to EU	
	I.1. Cons	0				1.2. 0	Certifica	te referen	ce numbe	er I.2.a.		
	Nam					I.3. C	Central	Competer	t Authority	y		
	Addr	ess				I.4. Local Competent Authority						
ent	Tel. N	10				1.4.		Inpetent	Authority			
gnm	I.5. Cons	signee				I.6.						
nsi	Nam	е										
d co	Addr	Address										
tche	Postal code											
spa	Tel. N	١o										
Part I: Details of dispatched consignment	I.7. Cour of ori		ISO code	I.8. Region of origin	Code		Country lestinat		ISO code	I.10. Region of destination	Code	
Deta	I.11. Place	e of origin				I.12.						
rt I:	Nam			Approval number								
Pa	Addr	ess						_				
	I.13. Place	e of loading				I.14. [Date of o	departure				
	I.15. Mea	ns of transpo	ort			I.16. E	Entry BI	P in EU				
	Aero	plane 🗌	Shi	p 📃 🛛 Railway wage	on 🗌							
	Road	l vehicle 🗌	Othe	er 🗌								
	Ident	ification:				1.17.						
		imentary ref	erences:									
	I.18. Desc	ription of co	mmodity			I.19. Commodity code (HS code)						
							L		I.20.	Quantity		
	I.21. Tem	perature of p	roduct						1.22.	Number of packages		
		bient		Chiled 🗌		Fro	zen 🗌			g		
						1102						
	I.23. Ident	ification of c	ontainer/se	eal number					1.24.	Type of packaging		
		modities cer an consump										
	I.26. I.28. Identification of the commodities						For impo	ort or admi	ission into	EU		
		ecies ific name)	Nature commo		App Abatto		mber e utting p	stablishme lant Co	ents Id store	Number of packages	Net weight	

Martiner

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

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

COUNT	COUNTRY Model SUF									
Ш.	Health information	II.a. Certificate reference number	II.b.							
II.3.	Animal welfare attestation I, the undersigned official veterinarian, hereby certify, that the fresh meat described in Part I derives from animals which have been treated in the slaughterhouse before and at the time of slaughter or killing in accordance with the relevant provisions of Union legislation.									
Notes										
	This certificate is meant for fresh meat, excluding offal and minced meat, of wild animals belonging to the Suidae, Tayassuidae, or Tapiridae families that are domestically kept or bred since birth in farms.									
Fresh m	eat means all animal parts fit for hur	nan consumption, whether fresh, chilled or fro	izen.							
 Box prov Box prov Box Box Box Box Box Box Box Cod <li< th=""><th>reference I.11: Place of origin: name reference I.15: Registration number ided. In case of unloading and reloa reference I.19: Use the appropriate reference I.20: Indicate total gross w reference I.23: For containers or bo reference I.28: <i>Nature of commodity</i> reference I.28: <i>Treatment type</i>: If ap cuts/pieces.</th><th>erritory as appearing in Part 1 of Annex II to Re e and address of the dispatch establishment. (railway wagons or container and lorries), fli ding, the consignor must inform the BIP of en HS code: 02.03, 02.08.90 or 05.04. weight and total net weight. xes, the container number and the seal number /: Indicate 'carcass-whole', 'carcass-side', 'car opropriate indicate deboned, or bone-in. If from t 1 of Annex II to Regulation (EU) No 206/201 is meat shall not be allowed when obtained from Jion of the third country, territory or part there been adopted by the Union against imports of</th><th>ght number (aircraft) or name (ship) is to be try into the Union. er (if applicable) should be included. ccass-quarters' or 'cuts'. zen, indicate the date of freezing (mm/yy) of 0. n animals slaughtered either prior to the date of referred to in boxes 1.7 and 1.8, or during a</th></li<>	reference I.11: Place of origin: name reference I.15: Registration number ided. In case of unloading and reloa reference I.19: Use the appropriate reference I.20: Indicate total gross w reference I.23: For containers or bo reference I.28: <i>Nature of commodity</i> reference I.28: <i>Treatment type</i> : If ap cuts/pieces.	erritory as appearing in Part 1 of Annex II to Re e and address of the dispatch establishment. (railway wagons or container and lorries), fli ding, the consignor must inform the BIP of en HS code: 02.03, 02.08.90 or 05.04. weight and total net weight. xes, the container number and the seal number /: Indicate 'carcass-whole', 'carcass-side', 'car opropriate indicate deboned, or bone-in. If from t 1 of Annex II to Regulation (EU) No 206/201 is meat shall not be allowed when obtained from Jion of the third country, territory or part there been adopted by the Union against imports of	ght number (aircraft) or name (ship) is to be try into the Union. er (if applicable) should be included. ccass-quarters' or 'cuts'. zen, indicate the date of freezing (mm/yy) of 0. n animals slaughtered either prior to the date of referred to in boxes 1.7 and 1.8, or during a							
Official v	eterinarian									
	Name (in capital letters):	Qualification	and title:							
	Date:	Signature:								
	Stamp:									

		del SUW				
	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				
nent	Tel. No					
ignn	I.5. Consignee	1.6.				
ons	Name					
ed c	Address					
atch	Postal code					
lispa	Tel. No					
Part I: Details of dispatched consignment	I.7. Country ISO I.8. Region Code of origin code of origin	I.9. Country of ISO I.10. Region of Code destination code destination				
Det	I.11. Place of origin	1.12.				
art I:	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					
	Identification:	l.17.				
	Documentary references:					
	I.18. Description of commodity	I.19. Commodity code (HS code)				
		I.20. Quantity				
		1.20. Quantity				
	I.21. Temperature of product	I.22. Number of packages				
	Ambient Chiled	Frozen				
	I.23. Identification of container/seal number	I.24. Type of packaging				
	I.25. Commodities certified for:					
	Human consumption					
	l.26.	I.27. For import or admission into EU				
	I.28. Identification of the commodities					
	Species Nature of Treatment A (Scientific name) commodity type	pproval number establishments Number Net of packages weight				
	(Scientific name) commonly type					
	, 104					

Model SUW

	COUNT	RY				Model SUW				
	II.	Health	information		II.a. Certificate reference number	II.b.				
ĸ	II.1.	Public	Health Attestation	on						
		(EC) N the Su	o 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				
rtificatio		II.1.1			(an) establishment(s) implementing a progra tion (EC) No 852/2004;	mme based on the HACCP principles in				
Part II: Certification		II.1.2	the meat has be particular:	en obt	obtained in accordance with Section IV of Annex III to Regulation (EC) No 853/2004, an in					
Ъа			(i) before skinn	ing, it ha	as been stored and handled separately from oth	ner food and not frozen;				
			and							
			(ii) after skinnin	g, it has	undergone a final inspection as referred to in p	oint II.1.4;				
		II.1.3			irements of Regulation (EC) No 2075/2005 lay and in particular, has been subject to an exami					
		II.1.4			d fit for human consumption following a post-m n I and Chapters VIII and IX of Section IV of Anr					
		II.1.5			rcass or parts of the carcass have been mark r III of Section I of Annex I to Regulation (EC) No					
			(1) or [the packages of meat have been marked with an identification mark in accordance with Se Annex II to Regulation (EC) No 853/2004;]							
		II.1.6	the meat satisfic foodstuffs;	sfies the relevant criteria set out in Regulation (EC) No 2073/2005 on microbiological criteria for						
		II.1.7		he 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 stored and transported in accordance with the relevant requirements of Section I of Annex III to Regulation (EC) No 853/2004							
	II.2.	Anima	l Health attestati	on						
		l, the u	ndersigned officia	l veterir	narian, hereby certify, that the fresh meat descril	ped in Part I:				
		II.2.1	has been obtain	ed in the	e territory/ies with code:	t the date of issuing this certificate:				
			(1) either		been free for 12 months from foot-and-mout ssical swine fever, 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				
				imp	ing the last 12 months no vaccination against orts of domestic animals vaccinated against itory;					

	information		II.a. Certificate reference number	II.b.
II.2.2			wild animals that were killed between d/mm/yyyy) (3) inside the territory referred to	
			eeds 20 km from the borders of a country or nis fresh meat into the Union,	part thereof, which is not authorised during thi
	(b) in an are point II.2.		uring the last 60 days, there has been no	o restrictions for the diseases referred to i
II.2.3.A	centre, and im of 10 km, ther in the event o	nmediately a e has been f a case of	afterwards] (¹) to an approved game-handlin no case/outbreak of the diseases referred t disease, the preparation of meat for import	ed within 12 hours for chilling [to a collectio ig establishment around which, within a radiu o in point II.2.1 during the previous 40 days o ation into the Union has been authorised on e establishment under the control of an officia
(1) (4) [II.2.3.B	has been obta negative resul		arcasses on which the following test for clas	sical swine fever was carried out and provide
	(1) either	[virus iso	lation from blood (EDTA);]	
	(1) or	[virus iso	lation from samples of	
	(1) <i>or</i>	[immuno	fluorescence for viral antigen on samples of	
II.2.4	has been obta certificate.	ained and pr	repared without contact with other meats no	t complying with the conditions required in th
	s meant for fres	h meat, exc	luding offal and minced meat, of wild anim	nals belonging to the Suidae, Tayassuidae, d
nis certificate is	s meant for fres s that are killed o			nals belonging to the Suidae, Tayassuidae, o
nis certificate is apiridae families	s that are killed o	or hunted in		
nis certificate is piridae families esh meat mear	s that are killed o ns all animal par	or hunted in ts fit for hur	the wild.	rozen.
nis certificate is piridae families esh meat mear ter importation	s that are killed o ns all animal par	or hunted in ts fit for hur	the wild. nan consumption whether fresh, chilled or fi	rozen.
nis certificate is apiridae families resh meat mear iter importation art I:	s that are killed o ns all animal par , unskinned caro	or hunted in ts fit for hur casses mus	the wild. nan consumption whether fresh, chilled or fi	rozen. ng establishment of destination.
nis certificate is apiridae families resh meat mear fter importation art I: - Box referenc	s that are killed o ns all animal par , unskinned caro e I.8: Provide th	or hunted in ts fit for hur casses mus e code of te	the wild. nan consumption whether fresh, chilled or fi t be conveyed without delay to the processi	rozen. ng establishment of destination. Regulation (EU) No 206/2010.
nis certificate is apiridae families esh meat mear ter importation art I: - Box referenc - Box referenc - Box referenc	s that are killed o ns all animal par , unskinned caro e I.8: Provide th e I.11: Place of o e I.15: Registrat	or hunted in ts fit for hur casses mus e code of te origin: name tion number	the wild. nan consumption whether fresh, chilled or fr t be conveyed without delay to the processi rritory as appearing in Part 1 of Annex II to F e and address of the dispatch establishmen	rozen. ng establishment of destination. Regulation (EU) No 206/2010. t. flight number (aircraft) or name (ship) is to b
nis certificate is apiridae families resh meat mear iter importation art I: - Box referenc - Box referenc provided. In o	s that are killed on ns all animal par , unskinned card e I.8: Provide th e I.11: Place of o e I.15: Registraticase of unloadir	or hunted in ts fit for hur casses mus e code of te origin: name tion number ng and reloa	the wild. nan consumption whether fresh, chilled or fr t be conveyed without delay to the processi rritory as appearing in Part 1 of Annex II to f e and address of the dispatch establishmen • (railway wagons or container and lorries),	rozen. ng establishment of destination. Regulation (EU) No 206/2010. t. flight number (aircraft) or name (ship) is to b
his certificate is apiridae families resh meat mear fter importation art I: - Box referenc - Box referenc provided. In o - Box referenc	s that are killed on ns all animal par , unskinned card e I.8: Provide th e I.11: Place of o e I.15: Registrat case of unloadir e I.19: Use the a	or hunted in ts fit for hur casses mus e code of te origin: name tion number ng and reloa appropriate	the wild. nan consumption whether fresh, chilled or fr t be conveyed without delay to the processi rritory as appearing in Part 1 of Annex II to f e and address of the dispatch establishmen r (railway wagons or container and lorries), ding, the consignor must inform the BIP of e	rozen. ng establishment of destination. Regulation (EU) No 206/2010. t. flight number (aircraft) or name (ship) is to b
his certificate is apiridae families resh meat mear fter importation art I: - Box referenc - Box referenc provided. In o - Box referenc - Box referenc	s that are killed of ns all animal par , unskinned card e I.8: Provide th e I.11: Place of e I.15: Registrat case of unloadir e I.19: Use the a e I.20: Indicate t	or hunted in ts fit for hur casses mus e code of te origin: name tion number ng and reloa appropriate total gross v	the wild. nan consumption whether fresh, chilled or fr t be conveyed without delay to the processi rritory as appearing in Part 1 of Annex II to F e and address of the dispatch establishmen r (railway wagons or container and lorries), ding, the consignor must inform the BIP of e HS code: 02.03, 02.08.90 or 05.04.	rozen. ng establishment of destination. Regulation (EU) No 206/2010. t. flight number (aircraft) or name (ship) is to b entry into the Union.
apiridae families resh meat mear fter importation art I: - Box referenc - Box referenc provided. In o - Box referenc - Box referenc - Box referenc	s that are killed on ns all animal par , unskinned card e I.8: Provide th e I.11: Place of o e I.15: Registrat case of unloadir e I.19: Use the a e I.20: Indicate t e I.23: For conta	or hunted in ts fit for hur casses mus e code of te origin: name tion number g and reloa appropriate total gross v ainers or box	the wild. nan consumption whether fresh, chilled or fr t be conveyed without delay to the processi rritory as appearing in Part 1 of Annex II to f e and address of the dispatch establishmen • (railway wagons or container and lorries), ding, the consignor must inform the BIP of e HS code: 02.03, 02.08.90 or 05.04. weight and total net weight.	rozen. ng establishment of destination. Regulation (EU) No 206/2010. t. flight number (aircraft) or name (ship) is to b entry into the Union.
his certificate is apiridae families resh meat mear fter importation art I: - Box referenc - Box referenc - Box referenc - Box referenc - Box referenc - Box referenc - Box referenc	s that are killed of ns all animal par , unskinned card e I.8: Provide th e I.11: Place of of e I.15: Registrat case of unloadir e I.19: Use the a e I.20: Indicate t e I.23: For conta e I.28: <i>Nature o</i> e I.28: <i>Nature o</i> e I.28: <i>Treatmer</i>	or hunted in ts fit for hun casses mus e code of te origin: name tion number ag and reloa appropriate total gross v ainers or boo f commodity	the wild. nan consumption whether fresh, chilled or fi t be conveyed without delay to the processi rritory as appearing in Part 1 of Annex II to f e and address of the dispatch establishmen r (railway wagons or container and lorries), ding, the consignor must inform the BIP of e HS code: 02.03, 02.08.90 or 05.04. weight and total net weight. kes, the container number and the seal num r: Indicate 'carcass-whole', 'carcass-side', 'c	ng establishment of destination. Regulation (EU) No 206/2010. t. flight number (aircraft) or name (ship) is to b entry into the Union.

со	COUNTRY Model SUW									
II.	Health information	II.a. Certificate reference numb	er II.b.							
	t II:	1								
• • •	(1) Keep as appropriate.									
	 Code of the territory as it appears in Part 1 of Annex II to Regulation (EU) No 206/2010. 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 I.7 and I.8, or during a period where restrictive measures have been adopted by the Union against imports of this meat from this third country, territory or part thereof. 									
(4)	 (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. 									
Off	icial veterinarian									
	Name (in capital letters):		Qualification and title:							
	Date:		Signature:							
	Stamp:									

	COUNTRY	Mod	el EQW	Veterinary certificate to EU			
	I.1. Consignor		I.2. Certificate reference r				
	Name						
	Address		I.3. Central Competent Authority				
Ŧ	Tel. No		I.4. Local Competent Authority				
men	I.5. Consignee		1.6.				
sign	Name		1.0.				
con	Address						
hed	Postal code						
patc	Tel. No						
Part I: Details of dispatched consignment	I.7. Country ISO of origin code	5		SO I.10. Region of Code destination			
etai	I.11. Place of origin		I.12.				
t I: D	Name	Approval number					
Par	Address						
-	I.13. Place of loading						
	1.13. Flace 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 reference	s:	l.17.				
-	I.18. Description of commodi	ty	I.19. Commodity code (HS code)				
				I.20. Quantity			
	I.21. Temperature of product			I.22. Number of packages			
	Ambient	Chiled	Frozen				
	I.23. Identification of containe	er/seal number		I.24. Type of packaging			
-	I.25. Commodities certified for Human consumption	pr:					
	1.26.		I.27. For import or admission into EU				
ŀ	I.28. Identification of the com	modities	I				
	Species (Scientific name)	Nature of Approval no commodity	umber establishments Number Ne of packages weig				
		Abattoir C	Cutting plant Cold store				

				Model EC
Н.	Health	information	II.a. Certificate reference number	II.b.
II.1.	Public	Health Attestation		
	(EC) N	lo 852/2004, (EC) N	eterinarian, declare that I am aware of the relevant o 853/2004 and (EC) No 854/2004 and hereby o s (zebra) described in Part I was produced in ac	certify that the meat of wild solipeds belonging
	II.1.1		from (an) establishment(s) implementing a pro egulation (EC) No 852/2004;	gramme based on the HACCP principles in
	II.1.2	the meat was obta	ned in compliance with Section IV of Annex III to	Regulation (EC) No 853/2004;
	II.1.3		requirements of Regulation (EC) No 2075/2005 I in particular, has been subject to an examination	
	II.1.4		found fit for human consumption following a pos ection I and Chapters VIII and IX of Section IV of	•
	II.1.5	•	e carcass or parts of the carcass have been n hapter III of Section I of Annex I to Regulation (EC	
		•	e packages of meat have been marked with an id nex II to Regulation (EC) No 853/2004;]	entification mark in accordance with Section I d
	II.1.6	the meat satisfies foodstuffs;	the relevant criteria set out in Regulation (EC) No 2073/2005 on microbiological criteria fo
	II.1.7	0	vering live animals and products thereof provided 3/EC, and in particular Article 29 thereof, are fulfil	, ,
	II.1.8	the meat has been Regulation (EC) No	stored and transported in accordance with the roo 853/2004.	elevant requirements of Section I of Annex III t
II.2.	Anima	I Health attestation	ı	
	l, the u	ndersigned official v	eterinarian, hereby certify, that the fresh meat de	scribed in Part I:
	II.2.1		I from wild animals that were killed between (dd/mm/yyyy) (2) inside the territory/ies with c	· · · · · · · · · · · · · · · · · · ·
	II.2.2	has been obtained centre, and immed of 10 km, there has the event of a case	from wild animals which after killing were transp iately afterwards] (1) to an approved game-handling been no case/outbreak of African horse sickness of such diseases, the preparation of meat for exp meat, and the total cleaning and disinfection of the	ng establishment around which, within a radiu s or glanders during the previous 40 days or, portation to the Union has been authorised on
	II.2.2 II.2.3	has been obtained centre, and immed of 10 km, there has the event of a case after removal of all veterinarian;	iately afterwards] (1) to an approved game-handli s been no case/outbreak of African horse sicknes of such diseases, the preparation of meat for exp	ng establishment around which, within a radius or glanders during the previous 40 days or, portation to the Union has been authorised on the establishment under the control of an offici
		has been obtained centre, and immed of 10 km, there has the event of a case after removal of all veterinarian; has been obtained	iately afterwards] (¹) to an approved game-handli s been no case/outbreak of African horse sicknes of such diseases, the preparation of meat for exp meat, and the total cleaning and disinfection of the	ng establishment around which, within a radiu s or glanders during the previous 40 days or, portation to the Union has been authorised on he establishment under the control of an offici
Notes		has been obtained centre, and immed of 10 km, there has the event of a case after removal of all veterinarian; has been obtained	iately afterwards] (¹) to an approved game-handli s been no case/outbreak of African horse sicknes of such diseases, the preparation of meat for exp meat, and the total cleaning and disinfection of the	ng establishment around which, within a radiu s or glanders during the previous 40 days or, portation to the Union has been authorised on he establishment under the control of an offici
This ce	II.2.3	has been obtained centre, and immed of 10 km, there has the event of a case after removal of all veterinarian; has been obtained certificate.	iately afterwards] (¹) to an approved game-handli s been no case/outbreak of African horse sicknes of such diseases, the preparation of meat for exp meat, and the total cleaning and disinfection of the	ng establishment around which, within a radiu is or glanders during the previous 40 days or, portation to the Union has been authorised on he establishment under the control of an offici at complying with the requirements set out in th
This ce (zebra).	II.2.3	has been obtained centre, and immed of 10 km, there has the event of a case after removal of all veterinarian; has been obtained certificate.	iately afterwards] (¹) to an approved game-handli s been no case/outbreak of African horse sickness of such diseases, the preparation of meat for exp meat, and the total cleaning and disinfection of th and prepared without contact with other meats no	ng establishment around which, within a radiu is or glanders during the previous 40 days or, i cortation to the Union has been authorised online establishment under the control of an officia of complying with the requirements set out in this objects belonging to the subgenus <i>Hippotigri</i>

COUNTRY		Model EQW
II. Health information	II.a. Certificate reference number	II.b.
 provided. In case of unloading and relo Box reference I.19: Use the appropriate Box reference I.20: Indicate total gross Box reference I.23: For containers or b Box reference I.28: Nature of commod. Box reference I.28: Treatment type: If a of the cuts/pieces. Box reference I.28: Abattoir: any abatto Part II: (1) Keep as appropriate. (2) Dates. Imports of this meat shall not be a for importation into the Union of the this 	he and address of the dispatch establer (railway wagons or container and loading, the consignor must inform the e HS code: 02.08.90 or 05.04. weight and total net weight. oxes, the container number and the sity: Indicate 'carcass-whole', 'carcass- ppropriate, indicate 'matured' or 'unsite oir or game handling establishment.	ishment. orries), flight number (aircraft) or name (ship) is to be BIP of entry into the Union. eal number (if applicable) should be included. side', 'carcass-quarters' or 'cuts'. kinned'. If frozen, indicate the date of freezing (mm/yy) s killed or hunted either prior to the date of authorisation ferred to in boxes I.7 and I.8, or during a period where s meat from this third country, territory or part thereof.
Official veterinarian		
Name (in capital letters):	-	ualification and title:
Date:	Si	gnature:
Stamp:		

ANNEX III

Model TRANSIT/STORAGE

Veterinary certificate to EU

	CO	UNTRY	Veterinary certificate to EU				
	I.1.	Consignor	I.2. Certificate reference number I.2.a.				
		Name	I.3. Central Competent Authority				
		Address					
ent		Tel. No	I.4. Local Competent Authority				
muß	I.5.	Consignee	I.6. Person responsible for the consignment in EU				
onsi		Name	Name				
o p		Address	Address				
tche		Postal code	Postal code				
ispa		Tel. No	Tel. No				
Part I: Details of dispatched consignment	I.7.	Country ISO I.8. Region Code of origin code of origin	I.9. Country of ISO I.10. Region of Code destination code destination				
Deta	I.11.	Place of origin	I.12. Place of destination				
Ë		Name Approval number	Custom warehouse Ship supplier				
Ра		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				
-	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					
	I.26	. For transit through EU to 3 rd Country	1.27.				
		3rd country ISO code					
	I.28	. Identification of the commodities					
	(5	Species Nature of Treatment Approval nu Scientific name) commodity type	Imber establishments Number Net of packages weight				
			Cutting manufacturing plant/ plant				

			Model TRANSIT/STORA							
п.	Health information	II.a. Certificate reference number	II.b.							
II.1.	Animal Health Attestation									
	I, the undersigned official veterinarian, hereby certify, that the fresh meat described in Part I:									
	II.1.1 comes from a country or region authorized for imports into the Union as laid down in Part 1 of Annex II to Regulation (EU) No 206/2010 at the time of slaughter, and									
	II.1.2 complies with the relevant animal health conditions as laid down in the animal health attestation in the mode certificate [BOV] [OVI] [POR] [EQU] [RUF] [RUW] [SUF] [SUW] [EQW] (¹) in Part 2 of Annex II to Regulation (EU No 206/2010, and									
		s which were slaughtered and processed (on (dd/mm/yyyy) o (dd/mm/yyyy) (²).							
		rage in accordance with Article 12(4) or Articl	e 13 of Directive 97/78/EC of:							
This cei	rtificate is meant for transit and stor h meat, including minced meat, of:	•	e 13 of Directive 97/78/EC of:							
This cei	h meat, including minced meat, of:	•								
This cei — fres	h meat, including minced meat, of: domestic bovine animals (incluc		ss-breeds) (Model 'BOV');							
This cer — fres (1)	h meat, including minced meat, of: domestic bovine animals (incluc	ling <i>Bubalus</i> and <i>Bison</i> species and their cro ries) or domestic caprine animals (<i>Capra hirc</i>	ss-breeds) (Model 'BOV');							
This cer — fres (1) (2) (3)	h meat, including minced meat, of: domestic bovine animals (incluc domestic ovine animals (<i>Ovis an</i>	ling <i>Bubalus</i> and <i>Bison</i> species and their cro ries) or domestic caprine animals (<i>Capra hirc</i> scrofa) (Model 'POR');	ss-breeds) (Model 'BOV');							
This cer — fres (1) (2) (3)	h meat, including minced meat, of: domestic bovine animals (incluc domestic ovine animals (<i>Ovis ar</i> domestic porcine animals (<i>Sus</i> a h meat, excluding minced meat, of	ling <i>Bubalus</i> and <i>Bison</i> species and their cro ries) or domestic caprine animals (<i>Capra hirc</i> scrofa) (Model 'POR');	ss-breeds) (Model 'BOV'); : <i>us</i>) (Model 'OVI');							
This cer — fres (1) (2) (3) — fres (4)	h meat, including minced meat, of: domestic bovine animals (incluc domestic ovine animals (<i>Ovis ar</i> domestic porcine animals (<i>Sus</i> a h meat, excluding minced meat, of	ling <i>Bubalus</i> and <i>Bison</i> species and their cro ries) or domestic caprine animals (<i>Capra hirc</i> scrofa) (Model 'POR'); : ! <i>Ilus, Equus asinus</i> and their cross-breeds) (N	ss-breeds) (Model 'BOV'); : <i>us</i>) (Model 'OVI');							
This cer — fres (1) (2) (3) — fres (4)	h meat, including minced meat, of: domestic bovine animals (incluc domestic ovine animals (<i>Ovis ai</i> domestic porcine animals (<i>Sus</i> h meat, excluding minced meat, of domestic solipeds (<i>Equus cabai</i> h meat, excluding offal and minced farmed non-domestic animals of	ling <i>Bubalus</i> and <i>Bison</i> species and their cro ries) or domestic caprine animals (<i>Capra hirc</i> scrofa) (Model 'POR'); : <i>Ilus, Equus asinus</i> and their cross-breeds) (N d meat, of: f the order Artiodactyla (excluding bovine anir	ss-breeds) (Model 'BOV'); eus) (Model 'OVI'); Aodel 'EQU'); mals (including <i>Bison</i> and <i>Bubalus</i> species and							
This cer — fres (1) (2) (3) — fres (4) — fres	h meat, including minced meat, of: domestic bovine animals (includ domestic ovine animals (<i>Ovis ar</i> domestic porcine animals (<i>Sus</i> h meat, excluding minced meat, of domestic solipeds (<i>Equus cabar</i> h meat, excluding offal and minced farmed non-domestic animals of their cross-breeds), <i>Ovis aries</i> , <i>C</i> (Model 'RUF'); wild non-domestic animals of th	ling <i>Bubalus</i> and <i>Bison</i> species and their cro ries) or domestic caprine animals (<i>Capra hirc</i> <i>scrofa</i>) (Model 'POR'); : <i>Ilus, Equus asinus</i> and their cross-breeds) (N d meat, of: f the order Artiodactyla (excluding bovine anir <i>Capra hircus</i> , Suidae and Tayassuidae), and o ne order Artiodactyla (excluding bovine anim	ss-breeds) (Model 'BOV'); eus) (Model 'OVI'); Model 'EQU'); mals (including <i>Bison</i> and <i>Bubalus</i> species and f the families Rhinocerotidae and Elephantidae als (including <i>Bison</i> and <i>Bubalus</i> species and							
This cer — fres (1) (2) (3) — fres (4) — fres (5)	h meat, including minced meat, of: domestic bovine animals (incluc domestic ovine animals (<i>Ovis ar</i> domestic porcine animals (<i>Sus s</i> h meat, excluding minced meat, of domestic solipeds (<i>Equus cabar</i> h meat, excluding offal and minced farmed non-domestic animals of their cross-breeds), <i>Ovis aries</i> , <i>C</i> (Model 'RUF'); wild non-domestic animals of th their cross-breeds), <i>Ovis aries</i> , <i>C</i> (Model 'RUW');	ling <i>Bubalus</i> and <i>Bison</i> species and their cro ries) or domestic caprine animals (<i>Capra hirc</i> <i>scrofa</i>) (Model 'POR'); : <i>Ilus, Equus asinus</i> and their cross-breeds) (N d meat, of: f the order Artiodactyla (excluding bovine anir <i>Capra hircus</i> , Suidae and Tayassuidae), and o ne order Artiodactyla (excluding bovine anim	ss-breeds) (Model 'BOV'); eus) (Model 'OVI'); Model 'EQU'); mals (including <i>Bison</i> and <i>Bubalus</i> species and f the families Rhinocerotidae and Elephantidae als (including <i>Bison</i> and <i>Bubalus</i> species and f the families Rhinocerotidae and Elephantidae							
This cer — fres (1) (2) (3) — fres (4) — fres (5) (6) (7)	h meat, including minced meat, of: domestic bovine animals (incluc domestic ovine animals (<i>Ovis ar</i> domestic porcine animals (<i>Sus a</i> h meat, excluding minced meat, of domestic solipeds (<i>Equus cabar</i> h meat, excluding offal and minced farmed non-domestic animals of their cross-breeds), <i>Ovis aries</i> , <i>C</i> (Model 'RUF'); wild non-domestic animals of th their cross-breeds), <i>Ovis aries</i> , <i>C</i> (Model 'RUW'); farmed non-domestic animals b	ling <i>Bubalus</i> and <i>Bison</i> species and their cro ries) or domestic caprine animals (<i>Capra hirc</i> scrofa) (Model 'POR'); : ! ! ! ! ! ! ! ! ! ! ! ! ! ! ! ! ! !	ss-breeds) (Model 'BOV'); sus) (Model 'OVI'); Model 'EQU'); mals (including <i>Bison</i> and <i>Bubalus</i> species and f the families Rhinocerotidae and Elephantidae als (including <i>Bison</i> and <i>Bubalus</i> species and of the families Rhinocerotidae and Elephantidae							
This cer — fres (1) (2) (3) — fres (4) — fres (5) (6)	h meat, including minced meat, of: domestic bovine animals (includ domestic ovine animals (<i>Ovis ar</i> domestic porcine animals (<i>Sus</i> a h meat, excluding minced meat, of domestic solipeds (<i>Equus cabar</i> h meat, excluding offal and minced farmed non-domestic animals of their cross-breeds), <i>Ovis aries</i> , <i>C</i> (Model 'RUF'); wild non-domestic animals of th their cross-breeds), <i>Ovis aries</i> , <i>C</i> (Model 'RUW'); farmed non-domestic animals belowed animals and animals animal	ling <i>Bubalus</i> and <i>Bison</i> species and their cro ries) or domestic caprine animals (<i>Capra hirc</i> <i>scrofa</i>) (Model 'POR'); : <i>Illus, Equus asinus</i> and their cross-breeds) (N I meat, of: If the order Artiodactyla (excluding bovine anim <i>Capra hircus</i> , Suidae and Tayassuidae), and o ne order Artiodactyla (excluding bovine anim <i>Capra hircus</i> , Suidae and Tayassuidae), and o elonging to the Suidae, Tayassuidae, or Tapir	ss-breeds) (Model 'BOV'); sus) (Model 'OVI'); Model 'EQU'); mals (including <i>Bison</i> and <i>Bubalus</i> species and f the families Rhinocerotidae and Elephantidae als (including <i>Bison</i> and <i>Bubalus</i> species and of the families Rhinocerotidae and Elephantidae							

COUNTRY Model TRANSIT/STORAG				
II. Health information	II.a. Certificate reference number	II.b.		
 Part I: Box reference I.8: Provide the code of tt Box reference I.11: Place of origin: nam Box reference I.12: Address (and approor or ship chandler shall be included. Box reference I.15: Registration numbe provided. In case of unloading and reloated in the provided. In case of unloading and reloated in the provided. In case of unloading and reloated in the provided. In case of unloading and reloated in the provided. In case of unloading and reloated in the provided. In case of unloading and reloated in the provided. In case of unloading and reloated in the provided. In case of unloading and reloated in the provided. In case of unloading and reloated in the provided. In case of unloading and reloated in the provided. In case of unloading and reloated in the provided. In case of unloading and reloated in the provided. In case of unloading and reloated in the provided. In case of unloading and reloated in the provided. In case of unloading and reloated in the provided. In case of unloading and reloated in the provided. In case of unloading and reloated in the provided. In case of unloading and reloated in the provided. In case of unloading and reloated to the provided. In case of unloated to the provided. In case of unloated in the provided of authorisation for exportation to the provided. In case of unloated in the provided of the provided	erritory as appearing in Part 1 of Annex II to Re e and address of the dispatch establishment. val number if known) of the warehouse in a free r (railway wagons or container and lorries), flig ading, the consignor must inform the BIP of en HS code: 02.01, 02.02, 02.03, 02.04, 02.05, 0	gulation (EU) No 206/2010. e zone, free warehouse, customs warehouse ght number (aircraft) or name (ship) is to be try into the Union. 12.06, 02.08.90, 02.09, 05.04 or 15.02. er (if applicable) should be included. cass-quarters', 'cuts', or 'minced meat'. he cuts/pieces.		
Official veterinarian				
Name (in capital letters):	Qualification	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': Model of veterinary certificate for consignments of queen bees and queen bumble bees (Apis mellifera and Bombus spp.),						
'BEE': Model of veterinary certi	ficate for consignments of colonie	s of bumble bees (Bombus spp.)				
Order	Family	Genera/species				
Hymenoptera	Apidae	Apis mellifera, Bombus spp.				

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

	COUNT	RY				Model QUE					
	П.	Health	information	II.a. Certificate reference number	II.b.						
	II.1. Animal Health attestation:										
		I, the undersigned, hereby certify, that the animals referred to in Part I of this certificate meet the following requirem									
u		II.1.1		ory with code:(1) in which, A aps mite <i>(Tropilaelaps</i> spp.) are notifiable		mall hive beetle <i>(Aethina</i>					
licati	II.1.2 they:(a) come from a breeding apiary, which is supervised and controlled by the competent authority;										
Certif											
Part II: Certification			and where no such certificate. Where an kilometres have beer	hich is not subject to any restrictions ass occurrence has taken place within at le outbreak of American foulbrood has occ checked by the competent authority and the said competent authority within 30 d	east 30 days prior to the curred previously, all hive d all infected hives burned	issuance of the present s within a radius of three or treated and inspected					
			have been tested in t	ne from hives or colonies (in the case on he last 30 days for American foulbrood a estrial Animals with negative results;	a particular and the second particular second particular						
				at least 100 km radius which is not subjec lle <i>(Aethina tumida)</i> or <i>Tropilaelaps</i> spp,							
				ne from hives or colonies (in the case of show no clinical signs or suspicion of dia							
				ailed examinations to ensure that all be da) or their eggs and larvae, or other infe							
		II.1.3		ueen cages, accompanying products ar combs, and all precautions have been ta of bees.							
	Notes										
	Part I:										
		reference) attenda		es (Apis mellifera and Bombus spp.). Eac	ch queen bee may be acco	ompanied by a maximum					
	Part II:										
	(1) Cod	e of the t	erritory as it appears in Pa	t 1 of Annex II or Section 1 of Part 1 of A	nnex IV to Regulation (EL	J) No 206/2010.					
	Official v	reterinaria	an /Official inspector								
		Name	(in capital letters):	Qualifi	ication and title:						
		Date:		Signat	ture:						
		Stamp	:								

				Mode	el BEE				
	со	UNTRY						Veterinary ce	ertificate to EU
	1.1.	Consignor	I.2. Certific	ate reference	numbei	r I.2.a.			
		Name	I.3. Central	Competent A	Authority				
		Address							
		Tel. No			I.4. Local C	ompetent Au	thority		
t	I.5.	Consignee			I.6.				
nme		Name							
Isig		Address							
l col		Postal code							
chec		Tel. No							
Part I: Details of dispatched consignment	I.7.	Country ISO of origin code	I.8. Region of origin	Code	I.9. Country destina	/ of tion c	ISO code	I.10. Region of destination	Code
ils o	l.11.	. Place of origin			l.12.				
l: Deta		Name Address	Approval number						
Part		Name Address	Approval number						
		Name Address	Approval number						
	I.13	. Place of loading Address	Approval number		I.14. Date of	departure	t	time of departure	
	l.15	. Means of transport			I.16. Entry B	IP in EU			
		Aeroplane Shi	p 🗌 🛛 Railway wago	on 🗌					
		Road vehicle 🗌 Othe	er 🗌						
		Identification: Documentary references:			I.17. No(s) of	CITES			
	I.18	. Description of commodity				I.19. Comn	nodity co	ode (HS code)	01.06.90
							1.20.0	Quantity	
	I.21						1.22.1	Number of packag	jes
	1.23	B. Identification of container/se	eal number				1.24.		
	1.25	i. Commodities certified for: Breeding							
	1.26).			I.27. For imp	ort or admiss	ion into	EU	
	1.28	B. Identification of the commo	dities						
		Species (Scientific name)		Identifi sys				Identificatio number	on

	COUNTR	Y			Model B					
	П.	Health	information	II.a. Certificate reference number	II.b.					
II.1. Animal Health attestation:										
		I, the undersigned, hereby certify that:								
		II.1.1								
					ificate have been bred and kept under a controllec rvised and controlled by the competent authority;					
					inspected immediately prior to dispatch and al spicion of disease including infestations affecting					
			broodstock and pack		led examination to ensure that all bumble bees e (Aethina tumida) or its eggs and larvae or othe					
		II.1.2		combs, and all precautions have been ta	ood are new and have not been in contact with ken to prevent contamination with agents causing					
	Notes									
	Part I:									
		eference Ile bees.	I.20: Number of containe	ers of bumble bees (<i>Bombus</i> spp.), each	n containing a colony of a maximum of 200 adu					
	Official ve	eterinaria	n /Official inspector							
					cation and title:					
		Name (in capital letters):	Qualifi						
		Name (Date:	in capital letters):	Qualific Signat						
		Date:								
		Date:								
		Date:								

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.

▼<u>C1</u>

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