

## II

(Non-legislative acts)

## REGULATIONS

## COMMISSION IMPLEMENTING REGULATION (EU) No 497/2012

of 7 June 2012

**amending Regulation (EU) No 206/2010 as regards the requirements for imports of animals susceptible to bluetongue**

(Text with EEA relevance)

THE EUROPEAN COMMISSION,

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

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<sup>(1)</sup>, and in particular Article 6(1), Article 7(e), and Article 13(1) thereof,

Whereas:

- (1) 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<sup>(2)</sup> lays down the list of third countries, territories or parts thereof from which live ungulate animals, including those susceptible to bluetongue, may be introduced into the Union and the veterinary certification requirements for such introduction.
- (2) In particular, with regard to animals susceptible to bluetongue, certificates BOV-X, BOV-Y, OVI-X, OVI-Y and RUM set out in Part 2 of Annex I to Regulation (EU) No 206/2010 include *inter alia* the requirement that the animals come from a territory which, at the date of issue of the certificate accompanying them had been free from bluetongue for a period of twelve months.
- (3) As a result of new technical developments, "inactivated vaccines" against bluetongue have become available

which do not pose the risk of undesired local circulation of the vaccine virus to unvaccinated cattle, sheep and goats. It is now widely accepted that vaccination with inactivated vaccines is the preferred tool for the control of bluetongue and for the prevention of clinical disease in such animals in the Union.

- (4) To ensure better control of the spread of the bluetongue virus and to reduce the burden on the agricultural sector posed by that disease, the rules on vaccination laid down in Council Directive 2000/75/EC of 20 November 2000 laying down specific provisions for the control and eradication of bluetongue<sup>(3)</sup> were recently amended by Directive 2012/5/EU of the European Parliament and of the Council<sup>(4)</sup> to take account of the recent technological developments in vaccine production.
- (5) Accordingly, Directive 2000/75/EC now provides for the use of inactivated vaccines in all parts of the EU.
- (6) As a result of the evolving epidemiological situation as regards bluetongue, and to align with the World Organisation for Animal Health (OIE) standards, Commission Regulation (EC) No 1266/2007 of 26 October 2007 on implementing rules for Council Directive 2000/75/EC as regards the control, monitoring, surveillance and restrictions on movements of certain animals of susceptible species in relation to bluetongue<sup>(5)</sup> was amended recently. The EU standards require the absence of virus circulation for a minimum period of two years in order to consider a territory free from bluetongue. The period of twelve months referred to in the relevant certificates set out in Part 2 of Annex I to Regulation (EU) No 206/2010 should therefore be amended accordingly.

<sup>(1)</sup> OJ L 139, 30.4.2004, p. 321.

<sup>(2)</sup> OJ L 73, 20.3.2010, p. 1.

<sup>(3)</sup> OJ L 327, 22.12.2000, p. 74.

<sup>(4)</sup> OJ L 81, 21.3.2012, p. 1.

<sup>(5)</sup> OJ L 283, 27.10.2007, p. 37.

- (7) Directive 2000/75/EC and Regulation (EC) No 1266/2007 apply to intra-Union movements of live ungulates of species susceptible to bluetongue. It is appropriate that the models of veterinary certificates BOV-X, BOV-Y, OVI-X, OVI-Y and RUM set out in Part 2 of Annex I to Regulation (EU) No 206/2010 be amended to align the animal health requirements for imports into the Union, as regards bluetongue, to the requirements for intra-Union movement in animals susceptible to that disease.
- (8) Regulation (EU) No 206/2010 should therefore be amended accordingly.
- (9) The measures provided for in this Regulation are in accordance with the opinion of the Standing Committee on the Food Chain and Animal Health,

HAS ADOPTED THIS REGULATION:

*Article 1*

Annex I to Regulation (EU) No 206/2010 is amended in accordance with the Annex to this Regulation.

*Article 2*

For a transitional period until 30 June 2012, consignments of live ungulates accompanied by a certificate issued before the date of entry into force of this Regulation in accordance with the models BOV-X, BOV-Y, OVI-X, OVI-Y or RUM set out in Part 2 of Annex I to Regulation (EU) No 206/2010 before the amendments introduced by this Regulation may continue to be introduced into the Union.

*Article 3*

This Regulation shall enter into force on the 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.

Done at Brussels, 7 June 2012.

*For the Commission*  
*The President*  
José Manuel BARROSO

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## ANNEX

In Annex I to Regulation (EU) No 206/2010, Part 2 is amended as follows:

(1) Models 'BOV-X', 'BOV-Y', 'OVI-X' and 'OVI-Y' are replaced by the following:

## 'Model BOV-X'

COUNTRY				Veterinary certificate to EU				
Part I: Details of dispatched consignment	I.1. Consignor Name Address Tel.			I.2. Certificate reference No		I.2.a.		
				I.3. Central competent authority				
				I.4. Local competent authority				
	I.5. Consignee Name Address Postal code Tel.			I.6.				
	I.7. Country of origin		ISO code	I.8. Region of origin		Code	I.9. Country of destination	
							I.10. Region of destination	
	I.11. Place of origin Name Address			Approval number		I.12.		
	I.13. Place of loading Address			Approval number		I.14. Date of departure		
	I.15. Means of transport Aeroplane <input type="checkbox"/> Ship <input type="checkbox"/> Railway wagon <input type="checkbox"/> Road vehicle <input type="checkbox"/> Other <input type="checkbox"/> Identification Documentary references			I.16. Entry BIP in EU		I.17.		
I.18. Description of commodity				I.19. Commodity code (HS code) <b>01.02</b>		I.20. Quantity		
I.21.						I.22. Number of packages		
I.23. Seal/Container No						I.24.		
I.25. Commodities certified for: Breeding <input type="checkbox"/> Fattening <input type="checkbox"/>								
I.26.			I.27. For import or admission into EU		<input type="checkbox"/>			
I.28. Identification of the commodities								
Species (scientific name)		Breed	Identification system	Identification number	Age	Sex		

COUNTRY

Model BOV-X

Part II: Certification	II.	Health information	II.a. Certificate reference number	II.b.
	II.1.	<p><b>Public Health Attestation</b></p> <p>I, the undersigned official veterinarian, hereby certify, that the animals described in this certificate:</p> <p>II.1.1. come from holdings which have been free from any official prohibition on health grounds, for the past 42 days in the case of 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 been in contact with animals from holdings which did not satisfy these conditions;</p> <p>II.1.2. have not received:</p> <ul style="list-style-type: none"> <li>— any stilbene or thyrostatic substances,</li> <li>— estrogenic, androgenic, gestagenic or <math>\beta</math>- agonist substances for purposes other than therapeutic or zootechnic treatment (as defined in Directive 96/22/EC);</li> </ul> <p>II.1.3. with regard to bovine spongiform encephalopathy (BSE):</p> <p>(<sup>1</sup>) (<sup>2</sup>) <i>either</i> [(a) the animals are identified by a permanent identification system enabling them to be traced back to the dam and herd of origin, and are not exposed bovine animals as described in Chapter C, part I, point (4) (b) (iv) of Annex II of Regulation (EC) No 999/2001;</p> <p>(b) if there have been BSE indigenous cases in the country concerned, the animals were born after the date from which the ban on the feeding of ruminants with meat-and-bone meal and greaves derived from ruminants had been effectively enforced or after the date of birth of the last BSE indigenous case if born after the date of the feed ban.]</p> <p>(<sup>1</sup>) (<sup>3</sup>) <i>or</i> [(a) the animals are identified by a permanent identification system enabling them to be traced back to the dam and herd of origin, and are not exposed bovine animals as described in Chapter C, Part II, point (4) (b) (iv) of Annex II of Regulation (EC) No 999/2001;</p> <p>(b) the animals were born after the date from which the ban on the feeding of ruminants with meat-and-bone meal and greaves derived from ruminants had been effectively enforced or after the date of birth of the last BSE indigenous case if born after the date of the feed ban.]</p> <p>(<sup>1</sup>) (<sup>4</sup>) <i>or</i> [(a) the animals are identified by a permanent identification system enabling them to be traced back to the dam and herd of origin, and are not exposed bovine animals as described in Chapter C, Part II, point (4) (b) (iv) of Annex II of Regulation (EC) No 999/2001;</p> <p>(b) the animals were born at least two years after the date from which the ban on the feeding of ruminants with meat-and-bone meal and greaves derived from ruminants had been effectively enforced or after the date of birth of the last BSE indigenous case if born after the date of the feed ban.]</p>		
	II.2.	<p><b>Animal Health attestation:</b></p> <p>I, the undersigned official veterinarian, hereby certify, that the animals described above meet the following requirements:</p> <p>II.2.1. they come from the territory with code: ..... (<sup>5</sup>) which, at the date of issuing this certificate:</p> <p>(<sup>1</sup>) <i>either</i> [(a) has been free for 24 months from foot-and-mouth disease]</p> <p>(<sup>1</sup>) <i>or</i> [(a) has been considered free from foot-and-mouth disease since ..... (dd/mm/yyyy), without having had cases/outbreaks after that date, and authorised to export these animals by Commission Implementing Regulation (EU) No .../..., of ..... (dd/mm/yyyy),]</p> <p>(b) has been free for 12 months from rinderpest, Rift valley fever, contagious bovine pleuropneumonia, lumpy skin disease and epizootic haemorrhagic disease, and for six months from vesicular stomatitis,</p> <p>(c) where during the last 12 months, no vaccination against the diseases mentioned in points (a) and (b) has been carried out and imports of domestic cloven-hoofed animals vaccinated against these diseases are not permitted;</p> <p>(<sup>1</sup>) <i>either</i> [(d) has been free for 24 months from bluetongue;]</p> <p>(<sup>1</sup>) (<sup>9</sup>) <i>or</i> [(d) has been free for 24 months from bluetongue, and the animals have reacted negatively to a serological test for the detection of antibody for bluetongue and epizootic haemorrhagic disease, carried out on two occasions on samples of blood taken at the beginning of the isolation/quarantine period and at least 28 days later, on ..... (dd/mm/yyyy) and on ..... (dd/mm/yyyy), the second of which must have been taken within 10 days before export;]</p>		

## COUNTRY

## Model BOV-X

II.	Health information	II.a. Certificate reference number	II.b.
	<p>(<sup>1</sup>) <i>or</i> [(d) has not been free for 24 months from bluetongue, and the animals have been vaccinated with an inactivated vaccine, at least 60 days before the date of dispatch to the Union, against all bluetongue serotype/s ..... (<i>insert serotype/s</i>) which are those present in the source population as demonstrated through a surveillance programme<sup>(12)</sup> in an area with a 150 km radius around the holding(s) of origin described under box reference I.11, and the animals are still within the immunity period of time guaranteed in the specifications of the vaccine;]</p> <p>II.2.2. they have remained in the territory described under point II.2.1 since birth, or for at least the last six months before dispatch to the Union and without contact with imported cloven-hoofed animals for the last 30 days;</p> <p>II.2.3. they have remained since birth or at least 40 days before dispatch in the holding(s) of origin described under box reference I.11.:</p> <p>(a) in and around which, in an area with a 150 km radius, there has been no case/outbreak of epizootic haemorrhagic disease during the previous 60 days,</p> <p>(b) in and around which, in an area with a 10 km radius, there has been no case/outbreak of foot-and-mouth disease, rinderpest, Rift valley fever, bluetongue, contagious bovine pleuropneumonia, lumpy skin disease and, vesicular stomatitis during the previous 40 days;</p> <p>II.2.4. they are not animals to be killed under a national programme for the eradication of diseases, nor have they been vaccinated against the diseases referred to under point II.2.1.(a) and (b);</p> <p>II.2.5. they come from herds that are not restricted under the national legislation pertaining to the eradication of tuberculosis, brucellosis and enzootic bovine leukosis;</p> <p>II.2.6. they come from herds recognised as officially tuberculosis-free<sup>(6)</sup>;</p> <p><i>and</i> (<sup>1</sup>) (<sup>7</sup>) <i>either</i> [come from a region which is recognised as officially tuberculosis-free<sup>(6)</sup>];</p> <p>(<sup>1</sup>) <i>or</i> [have been subjected to an intradermal tuberculin test<sup>(8)</sup> carried out with negative results within the past 30 days before dispatch to the Union;]</p> <p>(<sup>1</sup>) <i>or</i> [are less than six weeks old;]</p> <p>II.2.7. they have not been vaccinated against brucellosis and come from herds recognised as officially brucellosis-free<sup>(6)</sup>;</p> <p><i>and</i> (<sup>1</sup>) (<sup>7</sup>) <i>either</i> [come from a region which is recognised as officially brucellosis-free<sup>(6)</sup>];</p> <p>(<sup>1</sup>) <i>or</i> [have been subjected to at least one test for bovine brucellosis<sup>(8)</sup> carried out on samples taken within the past 30 days before dispatch to the Union,]</p> <p>(<sup>1</sup>) <i>or</i> [are less than 12 months old,]</p> <p>(<sup>1</sup>) <i>or</i> [are castrated males of any age,]</p> <p>(<sup>1</sup>) <i>either</i> [II.2.8. they come from herds included in an official system for the control of enzootic bovine leukosis, and in which there has been no evidence either clinical or as a result of a laboratory test of this disease during the past two years,]</p> <p>(<sup>1</sup>) <i>or</i> [II.2.8. they come from herds recognised as officially enzootic-bovine-leukosis-free<sup>(6)</sup> (<sup>6a</sup>),]</p> <p><i>and</i> (<sup>1</sup>) (<sup>7</sup>) <i>either</i> [come from a region which is recognised as officially enzootic-bovine-leukosis-free<sup>(6)</sup>];</p> <p>(<sup>1</sup>) <i>or</i> [have been subjected to an individual test for enzootic bovine leukosis<sup>(8)</sup> carried out with negative result on samples taken within the past 30 days before dispatch to the Union,]</p> <p>(<sup>1</sup>) <i>or</i> [are less than 12 months old;]</p> <p>II.2.9. they are/were (<sup>1</sup>) dispatched from their holding(s) of origin, without passing through any market:</p> <p>(<sup>1</sup>) <i>either</i> [directly to the Union,]</p> <p>(<sup>1</sup>) <i>or</i> [to the officially authorised assembly centre described under box reference I.13 situated within the territory described under point II.2.1,]</p>		

## COUNTRY

Model BOV-X

II. Health information	II.a. Certificate reference number	II.b.
<p>and, until dispatched to the Union:</p> <p>(a) they did not come in contact with other cloven-hoofed animals not complying with the health requirements as described in this certificate,</p> <p>(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.;</p> <p>II.2.10. any transport vehicles or containers in which they were loaded were cleaned and disinfected before loading with an officially authorised disinfectant;</p> <p>II.2.11. they were examined by an official veterinarian within 24 hours of loading and showed no clinical sign of disease;</p> <p>II.2.12. they have been loaded for dispatch to the Union on ..... (dd/mm/yyyy) <sup>(10)</sup> in the means of transport described under box reference I.15 above that were cleaned and disinfected before loading with an officially authorised disinfectant and so constructed that faeces, urine, litter or fodder could not flow or fall out of the vehicle or container during transportation.</p>		
<p><b>II.3. Animal transport attestation</b></p> <p>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.</p>		
<p><sup>(1)</sup> <sup>(11)</sup> <b>II.4. Specific requirements</b></p> <p>II.4.1. According to official information, no clinical or pathological evidence of infectious bovine rhinotracheitis (IBR) has been recorded in the holding(s) of origin referred to in box reference I.11, for the last 12 months;</p> <p>II.4.2. the animals referred to in box reference I.28.:</p> <p>(a) have been isolated in accommodation approved by the competent authority for the last 30 days immediately prior to dispatch for export,</p> <p>(b) have been subjected to a serological test for IBR on sera taken at least 21 days after entry into isolation, with negative results, and all animals in isolation have also given negative results to this test,</p> <p>(c) have not been vaccinated against IBR.]</p>		
<p><b>Notes</b></p>		
<p>This certificate is meant for domestic bovine animals (including <i>Bubalus</i> and <i>Bison</i> species and their cross-breeds) intended for breeding and/or production.</p>		
<p>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.</p>		
<p><b>Part I:</b></p>		
<p>— Box reference I.8.: Provide the code of territory as appearing in Part 1 of Annex I to Regulation (EU) No 206/2010.</p>		
<p>— 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.</p>		
<p>— 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.</p>		
<p>— Box reference I.23.: For containers or boxes, the container number and the seal number (if applicable) should be included.</p>		
<p>— Box reference I.28.: <i>Identification system:</i> The animals must bear:</p>		
<p>— An individual number which permits tracing of their premises of origin. Specify the identification system (such as tag, tattoos, brand, chip, transponder).</p>		
<p>— An ear tag that includes the ISO code of the exporting country. The individual number must permit tracing of their premises of origin.</p>		

## COUNTRY

## Model BOV-X

II. Health information	II.a. Certificate reference number	II.b.
<p><i>Species:</i> Select amongst "Bos", "Bison" and "Bubalus" as appropriate.</p> <p><i>Age:</i> Date of birth (dd/mm/yy).</p> <p><i>Sex</i> (M = male, F = female, C = castrated).</p> <p><i>Breed:</i> select purebred, crossbreed.</p> <p><b>Part II:</b></p> <p>(<sup>1</sup>) Keep as appropriate.</p> <p>(<sup>2</sup>) Only if the animals were born and continuously reared in a country or region categorised in accordance with Article 5(2) of Regulation (EC) No 999/2001 as a country or region posing a negligible BSE risk and listed as such in Decision 2007/453/EC.</p> <p>(<sup>3</sup>) Only if the country or region of origin is categorised in accordance with Article 5(2) of Regulation (EC) No 999/2001 as a country or region posing a controlled BSE risk and is listed as such in Decision 2007/453/EC.</p> <p>(<sup>4</sup>) Only if the country or region of origin has not been categorised in accordance with Article 5 (2) of Regulation (EC) No 999/2001 or has been categorised as a country or region with undetermined BSE risk and listed as such in Decision 2007/453/EC.</p> <p>(<sup>5</sup>) Code of the territory as it appears in Part 1 of Annex I to Regulation (EU) No 206/2010.</p> <p>(<sup>6</sup>) Officially tuberculosis/brucellosis-free regions and herds as laid down in Annex A to Directive 64/432/EEC; and enzootic-bovine-leukosis-free regions and herds as laid down in Chapter I of Annex D to Directive 64/432/EEC.</p> <p>(<sup>6a</sup>) Only for officially enzootic-bovine-leukosis-free herds recognised as equivalent to the requirements as laid down in Chapter I of Annex D to Directive 64/432/EEC for the purpose of exports to the EU of live animals according to the model certificate BOV-X from the territory that, in column 6 of Part 1 of Annex I to Regulation (EU) No 206/2010, appears with the entry "IVb" as regards enzootic bovine leukosis.</p> <p>(<sup>7</sup>) Only for a territory that, in column 6 of Part 1 of Annex I to Regulation (EU) No 206/2010, appears with the entry "II", as regards tuberculosis, "III", as regards brucellosis, and/or "IVa" as regards enzootic bovine leukosis.</p> <p>(<sup>8</sup>) Tests carried out in accordance with the protocols that, for the disease concerned, are described in Part 6 of Annex I to Regulation (EU) No 206/2010.</p> <p>(<sup>9</sup>) Supplementary guarantees to be provided when required in column 5 "SG" of Part 1 of Annex I to Regulation (EU) No 206/2010, with the entry "A".</p> <p>Tests for bluetongue and for epizootic haemorrhagic disease in accordance with Part 6 of Annex I to Regulation (EU) No 206/2010.</p> <p>(<sup>10</sup>) Date of loading. Imports of these animals shall not be allowed when the animals were loaded either prior to the date of authorisation for exportation to the Union of the third country, territory or part thereof referred to in Boxes 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.</p> <p>(<sup>11</sup>) When required by the EU Member State of destination or Switzerland, in accordance with Decision 2004/558/EC and in accordance with the Agreement between the Community and the Swiss Confederation on trade in agricultural products (OJ L 114, 30.4.2002, p. 132.).</p> <p>(<sup>12</sup>) Surveillance programme as laid down in Annex I to Commission regulation (EC) No 1266/2007 (OJ L 283, 27.10.2007, p. 37.).</p>		
<p>Official veterinarian</p> <p>Name (in capital letters):</p> <p>Date:</p> <p>Stamp:</p> <p>Qualification and title:</p> <p>Signature:</p>		

## Model BOV-Y

## COUNTRY

## Veterinary certificate to EU

Part I: Details of dispatched consignment	I.1. Consignor Name Address  Tel.		I.2. Certificate reference No		I.2.a.			
			I.3. Central competent authority					
			I.4. Local competent authority					
	I.5. Consignee Name Address  Postal code Tel.		I.6.					
	I.7. Country of origin	ISO code	I.8. Region of origin	Code	I.9. Country of destination	ISO code	I.10. Region of destination	Code
	I.11. Place of origin  Name Address		Approval number		I.12.			
	I.13. Place of loading  Address		Approval number		I.14. Date of departure			
	I.15. Means of transport  Aeroplane <input type="checkbox"/> Ship <input type="checkbox"/> Railway wagon <input type="checkbox"/> Road vehicle <input type="checkbox"/> Other <input type="checkbox"/> Identification Documentary references				I.16. Entry BIP in EU		I.17.	
	I.18. Description of commodity				I.19. Commodity code (HS code) <b>01.02</b>		I.20. Quantity	
							I.22. Number of packages	
I.23. Seal/Container No				I.24.				
I.25. Commodities certified for:  Slaughter <input type="checkbox"/>								
I.26.			I.27. For import or admission into EU <input type="checkbox"/>					
I.28. Identification of the commodities								
Species (scientific name)		Breed	Identification system	Identification number	Age	Sex		

## COUNTRY

Model BOV-Y

	II. Health information	II.a. Certificate reference number	II.b.
<b>Part II: Certification</b>	<b>II.1. Public Health Attestation</b>		
	I, the undersigned official veterinarian, hereby certify, that the animals described in this certificate:		
	II.1.1.	come from holdings which have been free from any official prohibition on health grounds, for the last 42 days in the case of brucellosis, for the last 30 days in the case of anthrax, for the last six months in the case of rabies, and, have not been in contact with animals from holdings which did not satisfy these conditions;	
	II.1.2.	have not received:	
		— any stilbene or thyrostatic substances,	
		— oestrogenic, androgenic, gestagenic or $\beta$ - agonist substances for purposes other than therapeutic or zootechnic treatment (as defined in Directive 96/22/EC).	
	II.1.3.	with regard to bovine spongiform encephalopathy (BSE):	
	( <sup>1</sup> ) ( <sup>2</sup> ) either	[(a) the animals are identified by a permanent identification system enabling them to be traced back to the dam and herd of origin, and are not exposed bovine animals as described in Chapter C, part I, point (4) (b) (iv) of Annex II of Regulation (EC) No 999/2001;	
		(b) if there have been BSE indigenous cases in the country concerned, the animals were born after the date from which the ban on the feeding of ruminants with meat-and-bone meal and greaves derived from ruminants had been effectively enforced or after the date of birth of the last BSE indigenous case if born after the date of the feed ban.]	
	( <sup>1</sup> ) ( <sup>3</sup> ) or	[(a) the animals are identified by a permanent identification system enabling them to be traced back to the dam and herd of origin, and are not exposed bovine animals as described in Chapter C, Part II, point (4) (b) (iv) of Annex II of Regulation (EC) No 999/2001;	
	(b) the animals were born after the date from which the ban on the feeding of ruminants with meat-and-bone meal and greaves derived from ruminants had been effectively enforced or after the date of birth of the last BSE indigenous case if born after the date of the feed ban.]		
( <sup>1</sup> ) ( <sup>4</sup> ) or	[(a) the animals are identified by a permanent identification system enabling them to be traced back to the dam and herd of origin, and are not exposed bovine animals as described in Chapter C, Part II, point (4) (b) (iv) of Annex II of Regulation (EC) No 999/2001;		
	(b) the animals were born at least two years after the date from which the ban on the feeding of ruminants with meat-and-bone meal and greaves derived from ruminants had been effectively enforced or after the date of birth of the last BSE indigenous case if born after the date of the feed ban.]		
<b>II.2. Animal Health Attestation</b>			
I, the undersigned official veterinarian, hereby certify, that the animals described above meet the following requirements:			
II.2.1.	they come from the territory with code: ..... ( <sup>5</sup> ) which, at the date of issuing this certificate:		
( <sup>1</sup> ) either	[(a) has been free for 24 months from foot-and-mouth disease]		
( <sup>1</sup> ) or	[(a) has been considered free from foot-and-mouth disease since ..... (dd/mm/yyyy), without having had cases/outbreaks after that date, and authorised to export these animals by Commission Implementing Regulation (EU) No ...../....., of ..... (dd/mm/yyyy);]		
	(b) has been free for 12 months from rinderpest, Rift valley fever, contagious bovine pleuropneumonia, lumpy skin disease and epizootic haemorrhagic disease, and for six months from vesicular stomatitis,		
	(c) where during the last 12 months, no vaccination against the diseases mentioned in points (a) and (b) has been carried out and imports of domestic cloven-hoofed animals vaccinated against these diseases are not permitted;		
( <sup>1</sup> ) either	[(d) has been free for 24 months from bluetongue;]		

COUNTRY

Model BOV-Y

II. Health information	II.a. Certificate reference number	II.b.
<p>(<sup>1</sup>) <i>or</i>                    [(d) has not been free for 24 months from bluetongue, and the animals have been vaccinated with an inactivated vaccine, at least 60 days before the date of dispatch to the Union, against all bluetongue serotype/s ..... (<i>insert serotype/s</i>) which are those present in the source population as demonstrated through a surveillance programme (<sup>9</sup>) in an area with a 150 km radius around the holding(s) of origin described under box reference I.11, and the animals are still within the immunity period of time guaranteed in the specifications of the vaccine;]</p> <p>II.2.2. they have remained in the territory described under point II.2.1 since birth, or for at least the last three months before dispatch to the Union and without contact with imported cloven-hoofed animals for the last 30 days;</p> <p>II.2.3. they have remained since birth or at least 40 days before dispatch in the holding(s) described under box reference I.11:</p> <p>(a) in and around which, in an area with a 150 km radius, there has been no case/outbreak of epizootic haemorrhagic disease during the previous 60 days, and</p> <p>(b) in and around which, in an area with a 10 km radius, there has been no case/outbreak of foot-and-mouth disease, rinderpest, Rift valley fever, bluetongue, contagious bovine pleuropneumonia, lumpy skin disease and, vesicular stomatitis during the previous 40 days;</p> <p>II.2.4. they are not animals to be killed under a national programme for the eradication of diseases, nor have they been vaccinated against the diseases referred to in point II.2.1(a) and (b);</p> <p>II.2.5. they come from herds:</p> <p>(a) included in an official system for the control of enzootic bovine leukosis, and</p> <p>(b) that are not restricted under the national legislation regarding eradication of tuberculosis and brucellosis, and</p> <p>(c) recognised as officially tuberculosis free; (<sup>6</sup>)</p> <p>II.2.6. they have not been vaccinated against brucellosis and they:</p> <p>(<sup>1</sup>) <i>either</i>            [come from herds which are recognised as officially brucellosis free;] (<sup>6</sup>)</p> <p>(<sup>1</sup>) <i>or</i>                    [are castrated males of any age;]</p> <p>II.2.7. they are individually marked on at least two places on their hindquarters as to show that they are exclusively intended for immediate slaughter; (<sup>7</sup>)</p> <p>II.2.8. they are/were (<sup>1</sup>) dispatched from their holding(s) of origin, without passing through any market:</p> <p>(<sup>1</sup>) <i>either</i>            [directly to the Union,]</p> <p>(<sup>1</sup>) <i>or</i>                    [to the officially authorised assembly centre described under box reference I.13 situated within the territory described under point II.2.1]</p> <p>and, until dispatched to the Union:</p> <p>(a) they did not come in contact with other cloven-hoofed animals not complying with the health requirements as described in this certificate, and</p> <p>(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;</p> <p>II.2.9. any transport vehicles or containers in which they were loaded were cleaned and disinfected before loading with an officially authorised disinfectant;</p> <p>II.2.10. they were examined by an official veterinarian within 24 hours of loading and showed no clinical sign of disease;</p> <p>II.2.11. they have been loaded for dispatch to the Union on ..... (dd/mm/yyyy) (<sup>8</sup>) in the means of transport described under box reference I.15 above that were cleaned and disinfected before loading with an officially authorised disinfectant and so constructed that faeces, urine, litter or fodder could not flow or fall out of the vehicle or container during transportation.</p>		

## COUNTRY

Model BOV-Y

II. Health information	II.a. Certificate reference number	II.b.
<p data-bbox="161 322 507 344"><b>II.3. Animal transport attestation</b></p> <p data-bbox="236 387 1461 454">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.</p> <p data-bbox="161 497 220 519"><b>Notes</b></p> <p data-bbox="161 562 1461 584">This certificate is meant for live bovine animals (including <i>Bubalus</i> and <i>Bison</i> species and their cross-breeds) intended for immediate slaughter.</p> <p data-bbox="161 627 1461 649">After importation the animals must be conveyed without delay to the slaughterhouse of destination to be slaughtered within five working days.</p> <p data-bbox="161 692 225 714"><b>Part I:</b></p> <ul style="list-style-type: none"> <li data-bbox="161 757 1225 779">— Box reference I.8: Provide the code of territory as appearing in Part 1 of Annex I to Regulation (EU) No 206/2010.</li> <li data-bbox="161 822 1461 866">— 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.</li> <li data-bbox="161 909 1461 954">— Box reference I.15: Registration number (railway wagons or container and lorries), flight number (aircraft) or name (ship) is to be provided. In case of unloading and reloading, the consignor must inform the BIP of entry into the Union.</li> <li data-bbox="161 996 1294 1019">— Box reference I.23: For containers or boxes, the container number and the seal number (if applicable) should be included.</li> <li data-bbox="161 1061 1461 1238">— Box reference I.28: <i>Identification system</i>: the animals must bear: <ul style="list-style-type: none"> <li data-bbox="193 1126 1461 1171">— An individual number which permits tracing of their premises of origin. Specify the identification system (such as tag, tattoos, brand, chip, transponder).</li> <li data-bbox="193 1214 1461 1236">— An ear tag that includes the ISO code of the exporting country. The individual number must permit tracing of their premises of origin.</li> </ul> </li> </ul> <p data-bbox="193 1281 820 1303"><i>Species</i>: Select amongst “<i>Bos</i>”, “<i>Bison</i>” and “<i>Bubalus</i>” as appropriate.</p> <p data-bbox="193 1346 464 1368"><i>Age</i>: Date of birth (dd/mm/yy).</p> <p data-bbox="193 1411 587 1433"><i>Sex</i> (M = male, F = female, C = castrated).</p> <p data-bbox="161 1476 225 1498"><b>Part II:</b></p> <ul style="list-style-type: none"> <li data-bbox="161 1541 384 1563">(1) Keep as appropriate.</li> <li data-bbox="161 1606 1461 1650">(2) Only if the animals were born and continuously reared in a country or region categorised in accordance with Article 5(2) of Regulation (EC) No 999/2001 as a country or region posing a negligible BSE risk and listed as such in Decision 2007/453/EC.</li> <li data-bbox="161 1693 1461 1738">(3) Only if the country or region of origin is categorised in accordance with Article 5(2) of Regulation (EC) No 999/2001 as a country or region posing a controlled BSE risk and is listed as such in Decision 2007/453/EC.</li> <li data-bbox="161 1780 1461 1825">(4) Only if the country or region of origin has not been categorised in accordance with Article 5(2) of Regulation (EC) No 999/2001 or has been categorised as a country or region with undetermined BSE risk and is listed as such in Decision 2007/453/EC.</li> <li data-bbox="161 1868 991 1890">(5) Code of the territory as it appears in Part 1 of Annex I to Regulation (EU) No 206/2010.</li> <li data-bbox="161 1933 1150 1955">(6) Officially tuberculosis/brucellosis free regions and herds as laid down in Annex A to Directive 64/432/EEC.</li> <li data-bbox="161 1998 1461 2042">(7) This mark shall take the form of “L” having 13 cm in the left side and 7 cm in the bottom side with 1 cm of strength in both lines. It shall be applied using the technique known as “freeze-branding”.</li> </ul>		



## Model OVI-X

## COUNTRY

## Veterinary certificate to EU

Part I: Details of dispatched consignment	I.1. Consignor Name Address Tel.		I.2. Certificate reference No		I.2.a.			
			I.3. Central competent authority					
			I.4. Local competent authority					
	I.5. Consignee Name Address Postal code Tel.		I.6.					
	I.7. Country of origin	ISO code	I.8. Region of origin	Code	I.9. Country of destination	ISO code	I.10. Region of destination	Code
	I.11. Place of origin Name Address Approval number		I.12.					
	I.13. Place of loading Address Approval number		I.14. Date of departure					
	I.15. Means of transport Aeroplane <input type="checkbox"/> Ship <input type="checkbox"/> Railway wagon <input type="checkbox"/> Road vehicle <input type="checkbox"/> Other <input type="checkbox"/> Identification Documentary references		I.16. Entry BIP in EU		I.17.			
	I.18. Description of commodity				I.19. Commodity code (HS code)			
					I.20. Quantity			
I.21.				I.22. Number of packages				
I.23. Seal/Container No				I.24.				
I.25. Commodities certified for: Breeding <input type="checkbox"/> Fattening <input type="checkbox"/>								
I.26.			I.27. For import or admission into EU <input type="checkbox"/>					
I.28. Identification of the commodities								
Species (scientific name)		Breed	Identification system	Identification number	Age	Sex		

COUNTRY

Model OVI-X

Part II: Certification	II.	Health information	II.a. Certificate reference number	II.b.
	II.1.	<b>Public Health Attestation</b>		
		I, the undersigned official veterinarian, hereby certify, that the animals described in this certificate:		
	II.1.1.	come from holdings which have been free from any official prohibition on health grounds, for the last 42 days in the case of brucellosis, for the last 30 days in the case of anthrax, for the last six months in the case of rabies, and, have not been in contact with animals from holdings which did not satisfy these conditions;		
	II.1.2.	have not received:		
		— any stilbene or thyrostatic substances,		
		— oestrogenic, androgenic, gestagenic or $\beta$ - agonist substances for purposes other than therapeutic or zootechnic treatment (as defined in Directive 96/22/EC).		
	II.2.	<b>Animal Health attestation</b>		
		I, the undersigned official veterinarian, hereby certify, that the animals described above meet the following requirements:		
	II.2.1.	they come from the territory with code: ..... (1) which, at the date of issuing this certificate:		
		(2) either [(a) has been free for 24 months from foot-and-mouth disease ]		
		(2) or [(a) has been considered free from foot-and-mouth disease since ..... (dd/mm/yyyy), without having had cases/outbreaks after that date, and authorised to export these animals by Commission Implementing Regulation (EU) No .../..., of ..... (dd/mm/yyyy).]		
		(b) has been free for 12 months from rinderpest, Rift valley fever, peste des petits ruminants, sheep pox and goat pox, contagious caprine pleuropneumonia, and epizootic haemorrhagic disease, and for 6 months from vesicular stomatitis,		
		(c) where during the last 12 months, no vaccination against the diseases mentioned in points (a) and (b) has been carried out and imports of domestic cloven-hoofed animals vaccinated against these diseases are not permitted;		
		(2) either [(d) has been free for 24 months from bluetongue;]		
		(2) (9) or [(d) has been free for 24 months from bluetongue, and the animals have reacted negatively to a serological test for the detection of antibody for bluetongue and epizootic haemorrhagic disease, carried out on two occasions on samples of blood taken at the beginning of the isolation/quarantine period and at least 28 days later, on ..... (dd/mm/yyyy) and on ..... (dd/mm/yyyy), the second of which must have been taken within 10 days before export;]		
		(2) or [(d) has not been free for 24 months from bluetongue, and the animals have been vaccinated with an inactivated vaccine, at least 60 days before the date of dispatch to the Union, against all bluetongue serotype/s ... (insert serotype/s) which are those present in the source population as demonstrated through a surveillance programme (11) in an area with a 150 km radius around the holding(s) of origin described under box reference I.11, and the animals are still within the immunity period of time guaranteed in the specifications of the vaccine;]		
	II.2.2.	they have remained in the territory described under point II.2.1 since birth, or for at least the last six months before dispatch to the Union and without contact with imported cloven-hoofed animals for the last 30 days;		
	II.2.3.	they have remained since birth or at least 40 days in the holding(s) described under box reference I.11 before dispatch:		
		(a) in and around which, in an area with a 150 km radius, there has been no case/outbreak of epizootic haemorrhagic disease during the previous 60 days, and		
		(b) in and around which, in an area with a 10 km radius, there has been no case/outbreak of foot-and-mouth disease, rinderpest, Rift valley fever, bluetongue, peste des petits ruminants, sheep pox and goat pox, contagious caprine pleuropneumonia and vesicular stomatitis during the previous 40 days;		

## COUNTRY

## Model OVI-X

II.	Health information	II.a. Certificate reference number	II.b.
II.2.4.	<p>according to my knowledge and to the written declaration made by the owner, the animals:</p> <p>(a) do not come from holdings, and have not been in contact with animals of a holding, in which the following diseases have been clinically detected:</p> <p>(i) contagious agalactia of sheep or goats (<i>Mycoplasma agalactiae</i>, <i>Mycoplasma capricolum</i>, <i>Mycoplasma mycoides</i> var. <i>mycoides</i> large colony), within the last six months,</p> <p>(ii) paratuberculosis and caseous lymphadenitis, within the last 12 months,</p> <p>(iii) pulmonary adenomatosis, within the last three years, and</p> <p>(iv) Maedi/Visna or caprine viral arthritis/encephalitis:</p> <p>(<sup>2</sup>) either [within the last three years,]</p> <p>(<sup>2</sup>) or [within the last 12 months, and all the infected animals were slaughtered and the remaining animals subsequently reacted negatively to two tests carried out at least six months apart,]</p> <p>(b) are included in an official system for notification of these diseases, and</p> <p>(c) have been free from clinical or other evidence of tuberculosis and brucellosis during the three years prior to export;</p>		
II.2.5.	they are not animals to be killed under a national programme for the eradication of diseases, nor have they been vaccinated against the diseases referred to in point II.2.1(a) and (b);		
II.2.6.	<p>they originate:</p> <p>(<sup>2</sup>) (<sup>3</sup>) either [from the territory described under box reference I.8, which has been recognised as officially brucellosis-free;]</p> <p>(<sup>2</sup>) or [from the holding(s) described under box reference I.11, where, in respect of brucellosis (<i>Brucella melitensis</i>):</p> <p>(a) all susceptible animals have been free from clinical or any signs of this disease for the last 12 months,</p> <p>(b) a representative number of the domestic ovine and caprine animals over an age of six months are submitted each year to a serological test, (<sup>4</sup>)</p> <p>(<sup>2</sup>) (<sup>5</sup>) either [(c) all domestic ovine or caprine animals have not been vaccinated against this disease, save those vaccinated with Rev. 1 vaccine more than two years ago;</p> <p>(d) the last two tests (<sup>6</sup>), separated by an interval of at least six months, carried out on ..... (dd/mm/yyyy) and on ..... (dd/mm/yyyy) on all domestic ovine and caprine animals over six months of age gave negative results, and]</p> <p>(<sup>2</sup>) or [(c) domestic ovine or caprine animals under the age of seven months are vaccinated against this disease with Rev. 1 vaccine;</p> <p>(d) the last two tests (<sup>6</sup>), separated by an interval of at least six months, carried out:</p> <p>— on ..... (dd/mm/yyyy) and on ..... (dd/mm/yyyy) on all non-vaccinated domestic ovine and caprine animals over six months of age, and</p> <p>— on ..... (dd/mm/yyyy) and on ..... (dd/mm/yyyy) on all vaccinated domestic ovine and caprine animals over 18 months of age</p> <p>gave negative results, and]</p> <p>(e) there are only domestic ovine and caprine animals that fulfil at least the above conditions and requirements;]</p>		

## COUNTRY

## Model OVI-X

II. Health information	II.a. Certificate reference number	II.b.
<p>(<sup>2</sup>) [II.2.7. the uncastrated rams have been kept continuously during the previous 60 days in a holding where no case of contagious epididymitis (<i>Brucella ovis</i>) has been diagnosed in the last 12 months and, these rams have undergone during the previous 30 days a complement fixation test to detect contagious epididymitis with a result of less than 50 IU/ml;]</p>		
<p>II.2.8. In respect of scrapie</p>		
<p>(<sup>2</sup>) (<sup>7</sup>) [II.2.8.1. if they are destined for a Member State which benefits, for all or part of its territory, from the provisions laid down in points (b) or (c) of Chapter A(I) of Annex VIII to Regulation (EC) No 999/2001, the animals comply with the guarantees provided for in the programmes referred to in those points and the animals comply with the guarantees requested by the EU Member States of destination regarding scrapie, and]</p>		
<p>(<sup>1</sup>) <i>either</i> [II.2.8.2. are animals intended for production born in and continuously reared on holdings in which a case of scrapie has never been diagnosed;]</p>		
<p>(<sup>2</sup>) (<sup>8</sup>) <i>or</i> [II.2.8.2. they shall have been kept continuously since birth or for the last three years on a holding or holdings which have satisfied the following requirements for at least three years:</p> <ul style="list-style-type: none"> <li>— they are subject to regular official veterinary checks,</li> <li>— the animals are identified in conformity with Union legislation,</li> <li>— no case of scrapie has been confirmed;</li> <li>— all animals over the age of 18 months which have died or been killed on the holdings (except the animals killed in the framework of a disease eradication campaign or slaughtered for human consumption) have been examined for scrapie in accordance with the laboratory methods laid down in point 3.2(b) of Chapter C of Annex X to Regulation (EC) No 999/2001;</li> <li>— domestic ovine and caprine animals, with the exception of domestic ovine animals of the ARR/ARR prion protein genotype have been introduced into the holding only if they come from holdings which complies with the above requirements] </li></ul>		
<p>(<sup>2</sup>) <i>or</i> [II.2.8.2. they are domestic ovine animals of the ARR/ARR prion protein genotype, as defined in Annex I to Decision 2002/1003/EC;]</p>		
<p>II.2.9. they are/were (<sup>1</sup>) dispatched from their holding(s) of origin, without passing through any market,</p>		
<p>(<sup>2</sup>) <i>either</i> [directly to the Union,]</p>		
<p>(<sup>2</sup>) <i>or</i> [to the officially authorised assembly centre described under box reference I.13 situated within the territory described under point II.2.1.]</p>		
<p>and, until dispatched to the Union:</p>		
<p>(a) they did not come in contact with other cloven-hoofed animals not complying with the health requirements as described in this certificate, and</p>		
<p>(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.;</p>		
<p>II.2.10. any transport vehicles or containers in which they were loaded were cleaned and disinfected before loading with an officially authorised disinfectant;</p>		
<p>II.2.11. they were examined by an official veterinarian within 24 hours of loading and showed no clinical sign of disease;</p>		
<p>II.2.12. they have been loaded for dispatch to the Union on ..... (dd/mm/yyyy) (<sup>10</sup>) in the means of transport described under box reference I.15 above that were cleaned and disinfected before loading with an officially authorised disinfectant and so constructed that faeces, urine, litter or fodder could not flow or fall out of the vehicle or container during transportation.</p>		

## COUNTRY

## Model OVI-X

II. Health information	II.a. Certificate reference number	II.b.
<p data-bbox="156 315 533 344"><b>II.3. Animal transport attestation</b></p> <p data-bbox="256 376 1461 450">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.</p> <p data-bbox="156 481 225 510"><b>Notes</b></p> <p data-bbox="156 542 1461 589">This certificate is meant for live domestic ovine animals (<i>Ovis aries</i>) and domestic caprine animals (<i>Capra hircus</i>) intended for breeding or production.</p> <p data-bbox="156 620 1461 667">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.</p> <p data-bbox="156 698 225 728"><b>Part I:</b></p> <ul style="list-style-type: none"> <li data-bbox="156 763 1225 792">— Box reference I.8: Provide the code of territory as appearing in Part 1 of Annex I to Regulation (EU) No 206/2010.</li> <li data-bbox="156 824 1461 871">— 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.</li> <li data-bbox="156 902 1461 949">— Box reference I.15: Registration number (railway wagons or container and lorries), flight number (aircraft) or name (ship) is to be provided. In case of unloading and reloading, the consignor must inform the BIP of entry into the Union.</li> <li data-bbox="156 981 842 1010">— Box reference I.19: Use the appropriate HS code: 01.04.10 or 01.04.20.</li> <li data-bbox="156 1041 1294 1070">— Box reference I.23: For containers or boxes, the container number and the seal number (if applicable) should be included.</li> <li data-bbox="156 1102 1461 1272">— Box reference I.28: <i>Identification system</i>: The animals must bear: <ul style="list-style-type: none"> <li data-bbox="188 1160 1461 1207">— 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.</li> <li data-bbox="188 1238 1461 1267">— An ear tag that includes the ISO code of the exporting country. The individual number must permit tracing of their premises of origin.</li> </ul> </li> </ul> <p data-bbox="188 1303 847 1332"><i>Species</i>: Select amongst "<i>Ovis aries</i>" and "<i>Capra hircus</i>" as appropriate.</p> <p data-bbox="188 1364 328 1393"><i>Age</i>: (months).</p> <p data-bbox="188 1424 588 1453"><i>Sex</i> (M = male, F = female, C = castrated).</p> <p data-bbox="156 1485 225 1514"><b>Part II:</b></p> <ol style="list-style-type: none"> <li data-bbox="167 1541 999 1570">(1) Code of the territory as it appears in Part 1 of Annex I to Regulation (EU) No 206/2010.</li> <li data-bbox="167 1601 392 1630">(2) Keep as appropriate.</li> <li data-bbox="167 1662 1225 1691">(3) Only for a territory appearing with the entry "V" in column 6 of Part 1 of Annex I to Regulation (EU) No 206/2010.</li> <li data-bbox="167 1722 1461 1980">(4) The representative number of animals to be tested for brucellosis must, for each holding, consist of: <ul style="list-style-type: none"> <li data-bbox="199 1778 1177 1807">— all non-castrated male animals, which have not been vaccinated against brucellosis, over six months old,</li> <li data-bbox="199 1839 1139 1868">— all non-castrated male animals, which have been vaccinated against brucellosis, over 18 months old,</li> <li data-bbox="199 1899 823 1928">— all animals brought onto the holding since the previous tests, and</li> <li data-bbox="199 1960 922 1989">— 25% of females which are sexually mature, within a minimum of 50 females.</li> </ul> </li> <li data-bbox="167 2011 1461 2058">(5) This must be completed when the destination is a Member State or part of a Member State laid down in one of the Annexes of Decision 93/52/EEC.</li> </ol>		

## COUNTRY

Model OVI-X

II. Health information	II.a. Certificate reference number	II.b.
<p>(<sup>6</sup>) In accordance with Part 6 of Annex I to Regulation (EU) No 206/2010.</p> <p>Where more than one holding of origin is involved the date of the most recent test on each holding must be clearly indicated.</p> <p>(<sup>7</sup>) 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.</p> <p>(<sup>8</sup>) In the case of animals intended, exclusively, for breeding purposes.</p> <p>(<sup>9</sup>) Supplementary guarantees to be provided when required in column 5 "SG" of Part 1 of Annex I to Regulation (EU) No 206/2010, with the entry "A". Tests for Bluetongue and for Epizootic-haemorrhagic-disease in accordance with Part 6 of Annex I to Regulation (EU) No 206/2010.</p> <p>(<sup>10</sup>) Date of loading. Imports of these animals shall not be allowed when the animals were loaded either prior to the date of authorisation for exportation to the Union of the third country, territory or part thereof referred to in boxes 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.</p> <p>(<sup>11</sup>) Surveillance programme as laid down in Annex I to Commission Regulation (EC) No 1266/2007 (OJ L 283, 27.10.2007, p. 37.).</p>		
<p>Official veterinarian</p> <p>Name (in capital letters):</p> <p>Date:</p> <p>Stamp:</p> <p>Qualification and title:</p> <p>Signature:</p>		

## Model OVI-Y

## COUNTRY

## Veterinary certificate to EU

Part I: Details of dispatched consignment	I.1. Consignor Name Address  Tel.		I.2. Certificate reference No		I.2.a.			
			I.3. Central competent authority					
			I.4. Local competent authority					
	I.5. Consignee Name Address  Postal code Tel.		I.6.					
	I.7. Country of origin	ISO code	I.8. Region of origin	Code	I.9. Country of destination	ISO code	I.10. Region of destination	Code
	I.11. Place of origin  Name Address		Approval number		I.12.			
	I.13. Place of loading  Address		Approval number		I.14. Date of departure			
	I.15. Means of transport  Aeroplane <input type="checkbox"/> Ship <input type="checkbox"/> Railway wagon <input type="checkbox"/> Road vehicle <input type="checkbox"/> Other <input type="checkbox"/> Identification Documentary references		I.16. Entry BIP in EU		I.17.			
	I.18. Description of commodity				I.19. Commodity code (HS code)		I.20. Quantity	
	I.21.				I.22. Number of packages			
I.23. Seal/Container No				I.24.				
I.25. Commodities certified for:  Slaughter <input type="checkbox"/>								
I.26.				I.27. For import or admission into EU <input type="checkbox"/>				
I.28. Identification of the commodities								
Species (scientific name)		Breed	Identification system	Identification number	Age	Sex		

COUNTRY

Model OVI-Y

Part II: Certification	II.	Health information	II.a. Certificate reference number	II.b.
	II.1.	<b>Public Health Attestation</b>		
		I, the undersigned official veterinarian, hereby certify, that the animals described in this certificate:		
	II.1.1.	come from holdings which have been free from any official prohibition on health grounds, for the last 42 days in the case of brucellosis, for the last 30 days in the case of anthrax, for the last six months in the case of rabies, and, have not been in contact with animals from holdings which did not satisfy these conditions;		
	II.1.2.	have not received:		
		— any stilbene or thyrostatic substances,		
		— oestrogenic, androgenic, gestagenic or $\beta$ - agonist substances for purposes other than therapeutic or zootechnic treatment (as defined in Directive 96/22/EC).		
	II.2.	<b>Animal Health attestation</b>		
		I, the undersigned official veterinarian, hereby certify, that the animals described above meet the following requirements:		
	II.2.1.	they come from the territory with code: ..... (1) which, at the date of issuing this certificate:		
		(2) <i>either</i> [(a) has been free for 24 months from foot-and-mouth disease ]		
		(2) <i>or</i> [(a) has been considered free from foot-and-mouth disease since ..... (dd/mm/yyyy), without having had cases/outbreaks after that date, and authorised to export these animals by Commission Implementing Regulation (EU) No ...../....., of ..... (dd/mm/yyyy).]		
		(b) has been free for 12 months from rinderpest, Rift valley fever, peste des petits ruminants, sheep pox and goat pox, contagious caprine pleuropneumonia, and epizootic haemorrhagic disease, and for 6 months from vesicular stomatitis,		
		(c) where during the last 12 months, no vaccination against the diseases mentioned in points (a) and (b) has been carried out and imports of domestic cloven-hoofed animals vaccinated against these diseases are not permitted;		
		(2) <i>either</i> [(d) has been free for 24 months from bluetongue;]		
		(2) <i>or</i> [(d) has not been free for 24 months from bluetongue, and the animals have been vaccinated with an inactivated vaccine, at least 60 days before the date of dispatch to the Union, against all bluetongue serotype/s ..... (insert serotype/s) which are those present in the source population as demonstrated through a surveillance programme (3) in an area with a 150 km radius around the holding(s) of origin described under box reference I.11., and the animals are still within the immunity period of time guaranteed in the specifications of the vaccine;]		
	II.2.2.	they have remained in the territory described under point II.2.1. since birth, or for at least the last three months before dispatch to the Union and without contact with imported cloven-hoofed animals for the last 30 days;		
	II.2.3.	they have remained since birth or at least 40 days before dispatch in the holding(s) described under box reference I.11:		
		(a) in and around which in an area with a 150 km radius there has been no case/outbreak of epizootic haemorrhagic disease during the previous 60 days, and		
		(b) in and around which, in an area with a 10 km radius, there has been no case/outbreak of foot-and-mouth disease, rinderpest, Rift valley fever, bluetongue, peste des petits ruminants, sheep pox and goat pox; contagious caprine pleuropneumonia and vesicular stomatitis during the previous 40 days;		
	II.2.4.	they are not animals to be killed under a national programme for the eradication of diseases, nor have they been vaccinated against the diseases referred to in point II.2.1(a) and (b);		
	II.2.5.	they are/were (2) dispatched from their holding(s) of origin, without passing through any market,		
		(2) <i>either</i> [directly to the Union]		

## COUNTRY

## Model OVI-Y

II. Health information	II.a. Certificate reference number	II.b.
<p>(<sup>2</sup>) or [to the officially authorised assembly centre described under box reference I.13 situated within the territory described under point II.2.1.]</p> <p>and, until dispatched to the Union:</p> <p>(a) they did not come in contact with other cloven-hoofed animals not complying with the health requirements as described in this certificate, and</p> <p>(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;</p> <p>II.2.6. in respect of scrapie:</p> <p>(<sup>2</sup>) (<sup>3</sup>) [II.2.6.1. if they are destined for a Member State which benefits, for all or part of its territory, from the provisions laid down in points (b) or (c) of Chapter A(I) of Annex VIII to Regulation (EC) No 999/2001, the animals comply with the guarantees provided for in the programmes referred to in those points, as laid down in Article 2 of Regulation (EC) 546/2006, and]</p> <p>(<sup>2</sup>) either [II.2.6.2. were born in and continuously reared on holdings in which a case of scrapie has never been diagnosed;]</p> <p>(<sup>2</sup>) or [II.2.6.2. are domestic ovine animals of the ARR/ARR prion protein genotype as defined in Annex I to Decision 2002/1003/EC, coming from a holding where no case of scrapie has been reported in the last six months;]</p> <p>II.2.7. any transport vehicles or containers in which they were loaded were cleaned and disinfected before loading with an officially authorised disinfectant;</p> <p>II.2.8. they were examined by an official veterinarian within 24 hours of loading and showed no clinical sign of disease;</p> <p>II.2.9. they have been loaded for dispatch to the Union on ..... (dd/mm/yyyy) (<sup>4</sup>) in the means of transport described under box reference I.15 above that were cleaned and disinfected before loading with an officially authorised disinfectant and so constructed that faeces, urine, litter or fodder could not flow or fall out of the vehicle or container during transportation.</p>		
<p>II.3. <b>Animal welfare attestation</b></p> <p>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.</p>		
<p><b>Notes</b></p>		
<p>This certificate is meant for live domestic ovine animals (<i>Ovis aries</i>) and domestic caprine animals (<i>Capra hircus</i>) intended for immediate slaughter after importation.</p>		
<p>After importation the animals must be conveyed without delay to the slaughterhouse of destination to be slaughtered within five working days.</p>		
<p><b>Part I:</b></p>		
<p>— Box reference I.8: Provide the code of territory as appearing in Part 1 of Annex I to Regulation (EU) No 206/2010.</p>		
<p>— 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.</p>		
<p>— 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.</p>		
<p>— Box reference I.19: Use the appropriate HS code: 01.04.10 or 01.04.20.</p>		
<p>— Box reference I.23: For containers or boxes, the container number and the seal number (if applicable) should be included.</p>		

## COUNTRY

Model OVI-Y

II. Health information	II.a. Certificate reference number	II.b.						
<p>— Box reference I.28: <i>Identification system</i>: The animals must bear:</p> <ul style="list-style-type: none"> <li>— 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.</li> <li>— An ear tag that includes the ISO code of the exporting country. The individual number must permit tracing of their premises of origin.</li> </ul> <p><i>Species</i>: Select amongst “<i>Ovis aries</i>” and “<i>Capra hircus</i>” as appropriate.</p> <p><i>Age</i>: months.</p> <p><i>Sex</i> (M = male, F = female, C = castrated).</p> <p><b>Part II:</b></p> <p>(<sup>1</sup>) Code of the territory as it appears in Part 1 of Annex I to Regulation (EU) No 206/2010.</p> <p>(<sup>2</sup>) Keep as appropriate.</p> <p>(<sup>3</sup>) 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.</p> <p>(<sup>4</sup>) Date of loading. Imports of these animals shall not be allowed when the animals were loaded either prior to the date of authorisation for exportation to the Union of the third country, territory or part thereof referred to in boxes 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.</p> <p>(<sup>5</sup>) Surveillance programme as laid down in Annex I to Commission Regulation (EC) No 1266/2007 (OJ L 283, 27.10.2007, p. 37.).</p>								
<p>Official veterinarian</p> <table style="width: 100%; border: none;"> <tr> <td style="width: 50%; border: none;">Name (in capital letters):</td> <td style="width: 50%; border: none;">Qualification and title:</td> </tr> <tr> <td style="border: none;">Date:</td> <td style="border: none;">Signature:</td> </tr> <tr> <td style="border: none;">Stamp:</td> <td style="border: none;"></td> </tr> </table>			Name (in capital letters):	Qualification and title:	Date:	Signature:	Stamp:	
Name (in capital letters):	Qualification and title:							
Date:	Signature:							
Stamp:								



COUNTRY

Model RUM

Part II: Certification	II.	Health information	II.a. Certificate reference number	II.b.
	II.1.	<p><b>Public Health Attestation</b></p> <p>I, the undersigned official veterinarian, hereby certify, that the animals described in this certificate:</p> <p>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;</p> <p>II.1.2. have not received:</p> <ul style="list-style-type: none"> <li>— any stilbene or thyrostatic substances,</li> <li>— oestrogenic, androgenic, gestagenic or <math>\beta</math>- agonist substances for purposes other than therapeutic or zootechnic treatment (as defined in Directive 96/22/EC).</li> </ul>		
II.2.	<p><b>Animal Health Attestation</b></p> <p>I, the undersigned official veterinarian, hereby certify, that the animals described above meet the following requirements:</p> <p>II.2.1. they come from the territory with code: ..... (1) which, at the date of issuing this certificate:</p> <ul style="list-style-type: none"> <li>(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</li> <li>(b) where during the last 12 months, no vaccination against foot-and-mouth disease, 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 during the last 24 months no vaccination against bluetongue has been carried out and imports of cloven-hoofed animals vaccinated against these diseases are not permitted;</li> </ul> <p>II.2.2. they have remained</p> <ul style="list-style-type: none"> <li>(2) <i>either</i> [in the territory described under point II.2.1. since birth, or for at least the last six months before dispatch to the Union and without contact with cloven-hoofed animals imported into this territory less than six months ago;]</li> <li>(2) <i>or</i> [in the country of dispatch for at least 60 days since entry, if they are animals of the relevant species listed in Part 7 of Annex I to Regulation (EU) No 206/2010 and they were imported directly under the conditions specified for each species in Part 7 of Annex I to Regulation (EU) No 206/2010 from a third country during a period of less than six months prior to embarkation to the Union and in any case they have been separated from other animals not of the same health status after being released in the exporting country and before exportation to the Union (3)]</li> </ul> <p>II.2.3. they have remained since birth or at least 40 days before dispatch in the holding/establishment (2) described under boxes reference I.11 and I.13:</p> <ul style="list-style-type: none"> <li>(a) in and around which in an area of radius of 150 km, there has been no case/outbreak of bluetongue and epizootic haemorrhagic disease during the previous 60 days, and</li> <li>(b) in and around which in an area of 10 km radius, there has been no case/outbreak of the other diseases referred to in point II.2.1 during the previous 40 days;</li> </ul> <p>II.2.4. they are not animals to be killed under a national programme for the eradication of diseases, nor have they been vaccinated against any of the diseases referred to in point II.2.1, and they:</p> <ul style="list-style-type: none"> <li>(2) (4) <i>either</i> [come from a herd which is recognised as officially tuberculosis free, and]</li> <li>(2) (5) <i>or</i> [have been subjected to an intradermal tuberculin test within the past 30 days with negative results, and]</li> </ul> <p>they have not been vaccinated against brucellosis and they:</p> <ul style="list-style-type: none"> <li>(2) (4) <i>either</i> [come from a herd which is recognised as officially brucellosis free;]</li> <li>(2) (5) <i>or</i> [have been subjected to a serum agglutination test which showed a brucella count of less than 30 IU of agglutination per ml, within the past 30 days;]</li> <li>(2) <i>or</i> [are castrated males of any age;]</li> </ul>			

## COUNTRY

## Model RUM

II. Health information	II.a. Certificate reference number	II.b.
<p>II.2.5. according to my knowledge and to the written declaration made by the owner, the animals:</p> <p>(a) do not come from holdings/establishments <sup>(2)</sup>, and have not been in contact with animals of a holding/establishment, in which the following diseases have been clinically detected:</p> <p>(i) contagious agalactia of sheep or goats (<i>Mycoplasma agalactiae</i>, <i>Mycoplasma capricolum</i>, <i>Mycoplasma mycoides</i> var. <i>mycoides</i> 'large colony'), within the last six months,</p> <p>(ii) paratuberculosis and caseous lymphadenitis, within the last 12 months,</p> <p>(iii) pulmonary adenomatosis, within the last three years, and</p> <p>(iv) Maedi/Visna or caprine viral arthritis/encephalitis,</p> <p><sup>(2)</sup> either [within the last three years,]</p> <p><sup>(2)</sup> or [within the last 12 months, and all the infected animals were slaughtered and the remaining animals subsequently reacted negatively to two tests carried out at least six months apart.]</p> <p>(b) are included in an official system for notification of these diseases, and</p> <p>(c) have been free from clinical or other evidence of tuberculosis and brucellosis during the three years prior to export;</p> <p><sup>(2)</sup> <sup>(6)</sup> [II.2.6. the animals have reacted negatively to a serological test for the detection of antibody for bluetongue and epizootic-haemorrhagic-disease, carried out on two occasions on samples of blood taken at the beginning of the isolation/quarantine period and at least 28 days later on ..... (dd/mm/yyyy) and on ..... (dd/mm/yyyy), the second of which must have been taken within 10 days of export;]</p> <p>II.2.7. they are dispatched from the holding/establishment described under boxes reference I.11 and I.13 directly to the Union and, until dispatched to the Union:</p> <p>(a) they did not come in contact with other cloven-hoofed animals not complying with the health requirements as described in this certificate, and</p> <p>(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;</p> <p>II.2.8. any transport vehicles or containers in which they were loaded were cleaned and disinfected before loading with an officially authorised disinfectant;</p> <p>II.2.9. they were examined by an official veterinarian within 24 hours of loading and showed no clinical sign of disease;</p> <p>II.2.10. they have been loaded for dispatch to the Union on ..... (dd/mm/yyyy) <sup>(7)</sup> in the means of transport described under box reference I.15. above that were cleaned and disinfected before loading with an officially authorised disinfectant and so constructed that faeces, urine, litter or fodder could not flow or fall out of the vehicle or container during transportation.</p>		
<p>II.3. <b>Animal transport attestation</b></p> <p>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.</p>		
<p><sup>(2)</sup> <sup>(8)</sup> [II.4. <b>Specific requirements</b></p> <p>II.4.1. According to official information, no clinical or pathological evidence of infectious bovine rhinotracheitis (IBR) has been recorded in the holding/establishment <sup>(2)</sup> of origin referred to in boxes reference I.11 and I.13, for the last 12 months;</p> <p>II.4.2. the animals referred to in box reference I.28.:</p> <p>(a) have been isolated in accommodation approved by the competent authority for the last 30 days immediately prior to dispatch for export, and</p> <p>(b) have been subjected to a serological test for IBR on sera taken at least 21 days after entry into isolation, with negative results, and all animals in isolation have also given negative results to this test, and</p>		

COUNTRY		Model RUM
II.	Health information	II.a. Certificate reference number II.b.
<p>(c) have not been vaccinated against IBR.;</p> <p>(<sup>2</sup>) [II.4.3. .... (further requirements and/or tests) .....]]</p>		
<p><b>Notes</b></p> <p>This certificate is meant for live animals of the order Artiodactyla (excluding bovine animals (including <i>Bubalus</i> and <i>Bison</i> species and their cross-breeds), <i>Ovis aries</i>, <i>Capra hircus</i>, Suidae and Tayassuidae), and of the families Rhinocerotidae and Elephantidae. Use one certificate per species.</p> <p>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.</p>		
<p><b>Part I:</b></p> <p>— Box reference I.8.: Provide the code of territory as appearing in Part 1 of Annex I to Regulation (EU) No 206/2010.</p> <p>— 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.</p> <p>— 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.</p> <p>— Box reference I.19.: Use the appropriate HS code: 01.02, 01.04.10, 01.04.20 or 01.06.19.</p> <p>— Box reference I.23.: For containers or boxes, the container number and the seal number (if applicable) should be included.</p> <p>— Box reference I.28.: <i>Identification system</i>: Specify the identification system (tag, tattoos, brand, chip, transponder). The ear tag includes the ISO code of the exporting country. The individual number must permit tracing of their premises of origin.</p> <p><i>Age</i>: months.</p> <p><i>Sex</i> (M = male, F = female, C = castrated).</p> <p><i>Species</i>: Select the species amongst those listed for the following families:</p> <p>Antilocapridae: <i>Antilocapra</i> spp.;</p> <p>Bovidae: <i>Addax</i> spp., <i>Aepyceros</i> spp., <i>Alcelaphus</i> spp., <i>Ammodorcas</i> spp., <i>Ammotragus</i> spp., <i>Antidorcas</i> spp., <i>Antilope</i> spp., <i>Bose-laphus</i> spp., <i>Budorcas</i> spp., <i>Capra</i> spp. (excluding <i>Capra hircus</i>), <i>Cephalophus</i> spp., <i>Connochaetes</i> spp., <i>Damaliscus</i> spp. (including <i>Beatragus</i>), <i>Dorcatragus</i> spp., <i>Gazella</i> spp., <i>Hemitragus</i> spp., <i>Hippotragus</i> spp., <i>Kobus</i> spp., <i>Litocranius</i> spp., <i>Madoqua</i> spp., <i>Naemorhedus</i> spp. (including <i>Nemorhaedus</i> and <i>Capricornis</i>), <i>Neotragus</i> spp., <i>Oreamnos</i> spp., <i>Oreotragus</i> spp., <i>Oryx</i> spp., <i>Ourebia</i> spp., <i>Ovibos</i> spp., <i>Ovis</i> spp. (excluding <i>Ovis aries</i>), <i>Pantholops</i> spp., <i>Pelea</i> spp., <i>Procapra</i> spp., <i>Pseudois</i> spp., <i>Pseudoryx</i> spp., <i>Raphicerus</i> spp., <i>Redunca</i> spp., <i>Rupicapra</i> spp., <i>Saiga</i> spp., <i>Sigmoceros-Alecelaphus</i> spp., <i>Sylvicapra</i> spp., <i>Syncerus</i> spp., <i>Taurotragus</i> spp., <i>Tetracerus</i> spp., <i>Tragelaphus</i> spp. (including <i>Boocerus</i>).</p> <p>Camelidae: <i>Camelus</i> spp., <i>Lama</i> spp., <i>Vicugna</i> spp.</p> <p>Cervidae: <i>Alces</i> spp., <i>Axis-Hyelaphus</i> spp., <i>Blastocerus</i> spp., <i>Capreolus</i> spp., <i>Cervus-Rucervus</i> spp., <i>Dama</i> spp., <i>Elaphurus</i> spp., <i>Hippocamelus</i> spp., <i>Hydropotes</i> spp., <i>Mazama</i> spp., <i>Megamuntiacus</i> spp., <i>Muntiacus</i> spp., <i>Odocoileus</i> spp., <i>Ozotoceros</i> spp., <i>Pudu</i> spp., <i>Rangifer</i> spp.</p> <p>Giraffidae: <i>Giraffa</i> spp., <i>Okapia</i> spp.</p> <p>Hippopotamidae: <i>Hexaprotodon-Choeropsis</i> spp., <i>Hippopotamus</i> spp.,</p> <p>Moschidae: <i>Moschus</i> spp.</p> <p>Tragulidae: <i>Hyemoschus</i> spp., <i>Tragulus-Moschiola</i> spp.,</p> <p>Rhinocerotidae: <i>Ceratotherium</i> spp., <i>Dicerorhinus</i> spp., <i>Diceros</i> spp., <i>Rhinoceros</i> spp.</p> <p>Elephantidae: <i>Elephas</i> spp., <i>Loxodonta</i> spp., as appropriate.</p>		

COUNTRY		Model RUM
II.	Health information	II.a. Certificate reference number II.b.
<p><b>Part II:</b></p> <p>(<sup>1</sup>) Code of the territory as it appears in Part 1 of Annex I to Regulation (EU) No 206/2010.</p> <p>(<sup>2</sup>) Keep as appropriate.</p> <p>(<sup>3</sup>) In this case the health certificate has to be accompanied by the official document on quarantine and test conditions laid down in Part 2 of Annex I to Regulation (EU) No 206/2010 (model "CAM").</p> <p>(<sup>4</sup>) Officially tuberculosis/brucellosis free regions or herds recognised as equivalent to the requirements laid down in Annex A to Directive 64/432/EEC and which appear in column 6 of Part 1 of Annex I to Regulation (EU) No 206/2010, with the entry "VII" as regards tuberculosis, "VIII", as regards brucellosis.</p> <p>(<sup>5</sup>) Tests carried out in accordance with the protocols that, for the disease concerned, are described in Part 6 of Annex I to Regulation (EU) No 206/2010. However for the tuberculin test a result of an increase in skin fold thickness of 2mm or more, or clinical signs of such as oedema, exudation, necrosis, pain and/or inflammation shall be deemed to be positive.</p> <p>(<sup>6</sup>) Supplementary guarantees to be provided when required in column 5 "SG" of Part 1 of Annex I to Regulation (EU) No 206/2010, with the entry "A". Tests for Bluetongue and for Epizootic-haemorrhagic-disease in accordance with Part 6 of Annex I to Regulation (EU) No 206/2010.</p> <p>(<sup>7</sup>) Date of loading. Imports of these animals shall not be allowed when the animals were loaded either prior to the date of authorisation for exportation to the Union of the third country, territory or part thereof referred to in boxes 1.7. and 1.8., or during a period where restrictive measures have been adopted by the Union against imports of these animals from this third country, territory or part thereof.</p> <p>(<sup>8</sup>) When required by the EU Member State of destination.</p>		
<p>Official veterinarian</p> <p>Name (in capital letters):</p> <p>Date:</p> <p>Stamp:</p> <p>Qualification and title:</p> <p>Signature:</p>		