

(2) The Model 'RUM' is replaced by the following:

'Model RUM**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. No(s) of CITES	
	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"/> 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) Identification system Identification number Age Sex								

COUNTRY

Model RUM

		II.a. Certificate reference number	II.b.
Part II: Certification	II.	Health information	
	II.1.	Public Health Attestation	
	I, the undersigned official veterinarian, hereby certify, that the animals described in this certificate:		
	II.1.1.	come from a holding which has been free from any official prohibition on health grounds, for the last 42 days in the case of brucellosis 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.1.2.	have not received:	
		— any stilbene or thyrostatic substances,	
		— oestrogenic, androgenic, gestagenic or β - agonist substances for purposes other than therapeutic or zootechnic treatment (as defined in Directive 96/22/EC).	
	II.2.	Animal Health Attestation	
	I, the undersigned official veterinarian, hereby certify, that the animals described above meet the following requirements:		
	II.2.1.	they come from the territory with code: (1) which, at the date 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 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;	
	II.2.2.	they have remained	
	(2) either	[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;]	
	(2) or	[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)]	
	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:	
	(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	
	(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;	
	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:	
	(2) (4) either	[come from a herd which is recognised as officially tuberculosis free, and]	
	(2) (5) or	[have been subjected to an intradermal tuberculin test within the past 30 days with negative results, and]	
	they have not been vaccinated against brucellosis and they:		
	(2) (4) either	[come from a herd which is recognised as officially brucellosis free;]	
	(2) (5) or	[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;]	
	(2) or	[are castrated males of any age;]	

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 ⁽²⁾, 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>⁽²⁾ either [within the last three years,]</p> <p>⁽²⁾ 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>⁽²⁾ ⁽⁶⁾ [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) ⁽⁷⁾ 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. Animal transport attestation</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>⁽²⁾ ⁽⁸⁾ [II.4. Specific requirements</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 ⁽²⁾ 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>(²) [II.4.3. (further requirements and/or tests)]]</p> <p>Notes</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>Part I:</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>Boselaphus</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>Naemorrhhedus</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>Part II:</p> <p>(¹) Code of the territory as it appears in Part 1 of Annex I to Regulation (EU) No 206/2010.</p> <p>(²) Keep as appropriate.</p> <p>(³) 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>(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.</p> <p>(⁵) Tests carried out in accordance with the protocols that, for the disease concerned, are described in Part 6 of Annex I to Regulation (EU) No 206/2010. However for the tuberculin test a result of an increase in skin fold thickness of 2mm or more, or clinical signs of such as oedema, exudation, necrosis, pain and/or inflammation shall be deemed to be positive.</p> <p>(⁶) 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>(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.</p> <p>(⁸) 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>		