

Part I : Details of consignment	I.1. Consignor			I.2. IMSOC Reference		
	Name			I.2.a. Local Reference		
	Address					
	Country			ISO Code		
	I.5. Consignee			I.3. Central competent authority		
	Name			I.4. Local competent authority		
	Address					
	Country			ISO Code		
	I.7. Country of origin			I.9. Country of destination		
	ISO Code			ISO Code		
	I.8. Region of origin			I.10. Region of destination		
	Code					
	I.11. Place of Dispatch			I.12. Place of destination		
	Name			Name		
	Address			Address		
Approval Number			Approval Number			
Country			Country			
ISO Code			ISO Code			
I.13. Place of Loading			I.14. Date and time of departure			
Name						
Address						
Approval Number						
Country						
ISO Code						
I.15. Means of Transport			I.16 Entry Point			
Mode	International transport document	Identification				
I.18. Transport conditions			I.17. Accompanying documents			
Ambient <input type="checkbox"/>			Commercial document reference			
			Date of issue			
			Country			
			Place of issue			
I.19. Container No / Seal No						
I.20. Certified as						
Animal Feedingstuff <input type="checkbox"/>		Pharmaceutical use <input type="checkbox"/>		Relaying <input type="checkbox"/>		
Slaughter <input type="checkbox"/>		Technical use <input type="checkbox"/>		Production <input type="checkbox"/>		
Breeding and production <input type="checkbox"/>		Artificial reproduction <input type="checkbox"/>		Breeding <input type="checkbox"/>		
Production of petfood <input type="checkbox"/>				Human consumption <input type="checkbox"/>		
				Fattening <input type="checkbox"/>		
				Other <input type="checkbox"/>		
I.21. For transit through a third country <input type="checkbox"/>			I.22. For transit through Member State(s) <input type="checkbox"/>			
Country			Country			
ISO Code			ISO Code			
EU Exit Authority			BCP code			
EU Entry Authority			BCP code			
I.23. Total number of packages		I.24. Total quantity		I.25. Total net weight		
				I.25. Total gross weight		
I.28. Description of consignment						
1. 01 LIVE ANIMALS						
0102 Live bovine animals						
Commodity	Species	Quantity	Net weight	Package count		
Identification number			Identification system			

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,		
		- estrogenic, androgenic, gestagenic or β - agonist substances for purposes other than therapeutic or zootechnical treatment (as defined in Directive 96/22/EC);		
	II.2.	Animal health attestation		
		1, 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, for 12 months from rinderpest, Rift valley fever, contagious bovine pleuropneumonia, lumpy skin disease, peste des petits ruminants, sheep pox and goat pox and contagious caprine pleuropneumonia, and for 6 months from vesicular stomatitis,			
	(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,			
(2)	○ either [(c) has been free for 24 months from bluetongue and 12 months for epizootic haemorrhagic disease;]			
(2)(5)	○ or [(c) has been free for 24 months from bluetongue, and the animals have reacted negatively to a serological test for the detection of antibodies 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)(8)	○ or [(c) is seasonally free of bluetongue and epizootic haemorrhagic disease and the animals have been kept during the seasonally free period in the seasonally free territory since birth or for at least 60 days prior to shipment;]			
(2)(8)	○ or [(c) is seasonally free of bluetongue and epizootic haemorrhagic disease and the animals have been kept during the seasonally free period in the seasonally free territory for at least 28 days prior to shipment, and have reacted negatively to a serological test according to the WOA (formerly OIE) Manual for detection of antibodies for bluetongue, carried out at least 28 days after the start of the residence period;]			
(2)(8)	○ or [(c) is seasonally free of bluetongue and epizootic haemorrhagic disease and the animals have been kept during the seasonally free period in the seasonally free territory for at least 14 days prior to shipment, and have reacted negatively to a PCR test for bluetongue virus according to the WOA (formerly OIE) Manual, carried out at least 14 days after the start of the residence period;]			
(2) ○ either	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 Great Britain and without contact with cloven-hoofed animals imported into this territory less than six months ago;]			

Part II: Certification	II. Health information		
	(2) ○ Or	[II.2.2.	have been introduced on (dd/mm/yyyy) into the territory described under point II.2.1, from the EU territory with code (1) that at that date was authorised to import the animals into Great Britain and the animals have not been in contact with imported cloven-hoofed animals from countries not subject to transitional import arrangements for the last 30 days.]
		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)(3)	○ either	[come from a herd which is recognised as officially tuberculosis free, and]
	(2)(4)	○ 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)(3)	○ either	[come from a herd which is recognised as officially brucellosis free;]
	(2)(4)	○ 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;]
	II.2.5.	according to my knowledge and to the written declaration made by the owner, the animals: (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: (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, (ii) paratuberculosis and caseous lymphadenitis, within the last 12 months, (iii) pulmonary adenomatosis, within the last three years, and (iv) Maedi/Visna or caprine viral arthritis/encephalitis,	
(2)	○ either	[within the last three years,]	
(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,]	
	(b)	are included in an official system for notification of these diseases, and	
	(c)	have been free from clinical or other evidence of tuberculosis and brucellosis during the three years prior to export;	
	II.2.6.	they are dispatched from the holding/establishment described under boxes reference I.11. and I.13. directly to Great Britain and, until dispatched to Great Britain:	
	(a)	they did not come in contact with other cloven-hoofed animals not complying with the health requirements as described in this certificate, and	
	(b)	they were not at any place where, or around which within a 10 km radius, during the previous 30 days there has been a case/outbreak of any of the diseases referred to in point II.2.1.;	
	II.2.7.	any transport vehicles or containers in which they were loaded were cleaned and disinfected before loading with an officially authorised disinfectant;	

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	II.2.8.	they were examined by an official veterinarian within 24 hours of loading and showed no clinical sign of disease;		
	II.2.9.	they have been loaded for dispatch to Great Britain on (dd/mm/yyyy) (6) in the means of transport described under box reference I.15. 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.		
	II.3.	Animal transport attestation		
	I, the undersigned official veterinarian, hereby certify, that the animals described above are fit for the intended transport and have been treated before and at the time of loading in accordance with the relevant provisions of Regulation (EC) No 1/2005.			
	(2)(8) <input type="checkbox"/> II.4. Specific requirements			
	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;		
	II.4.2.	the animals referred to in box reference I.28.:		
		(a) have been isolated in accommodation approved by the competent authority for the last 30 days immediately prior to dispatch for export, and		
		(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		
	(c) have not been vaccinated against IBR.			
(2)	<input type="checkbox"/> II.4.3.	(further requirements and/or tests)]]	

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	<p>Notes</p> <p>References to European Union legislation within this certificate are references to direct EU legislation which has been retained in Great Britain (retained EU law as defined in the European Union (Withdrawal) Act 2018) and can be viewed on the UK legislation website (legislation.gov.uk).</p> <p>References to Great Britain in this certificate include Channel Islands and Isle of Man.</p> <p>This certificate is meant for live animals of the order Artiodactyla (excluding bovine animals (including Bubalus and Bison species and their crossbreeds), Ovis aries, Capra hircus, 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> <ul style="list-style-type: none"> · Box reference I.8.: Provide the code of territory as it appears in a document relating to 'live ungulates' published on gov.uk, in accordance with Regulation (EU) No 206/2010.(9) · Box reference I.13.: The assembly centre, if any, must fulfil the conditions for its approval, as laid down in Part 5 of Annex 1 to Regulation (EU) No 206/2010. · Box reference I.15.: Registration number (railway wagons or container and lorries), flight number (aircraft) or name (ship) is to be provided. In case of unloading and reloading, the consignor must inform the BCP of entry into Great Britain. · Box reference I.19.: Use the appropriate HS code: 01.02, 01.04.10, 01.04.20 or 01.06.19. · Box reference I.23.: For containers or boxes, the container number and the seal number (if applicable) should be included. · Box reference I.28.: Identification system: 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. Age: months. Sex (M = male, F = female, C = castrated). Species: Select the species amongst those listed for the following families: <ul style="list-style-type: none"> Antilocapr idae: Antilocapra spp. Bovidae: Addax spp., Aepyceros spp., Alcelaphus spp., Ammodorcas spp., Ammotragus spp., Antidorcas spp., Antilope spp., Boselaphus spp., Budorcas spp., Capra spp. (excluding Capra hircus), Cephalophus spp., Connochaetes spp., Damaliscus spp. (including Beatragus), Dorcatragus spp., Gazella spp., Hemitragus spp., Hippotragus spp., Kobus spp., Litocranius spp., Madoqua spp., Naemorhedus spp. (including Nemorhaedus and Capricornis), Neotragus spp., Oreamnos spp., Oreotragus spp., Oryx spp., Ourebia spp., Ovibos spp., Ovis spp. (excluding Ovis aries), Pantholops spp., Pelea spp., Procapra spp., Pseudois spp., Pseudoryx spp., Raphicerus spp., Redunca spp., Rupicapra spp., Saiga spp., Sigmoceros-Alecelaphus spp., Sylvicapra spp., Syncerus spp., Taurotragus spp., Tetracerus spp., Tragelaphus spp. (including Boocerus). Camelidae: Camelus spp., Lama spp., Vicugna spp. 		

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	<p style="margin-left: 40px;">Cervidae: Alces spp., Axis-Hyelaphus spp., Blastocerus spp., Capreolus spp., Cervus-Rucervus spp., Dama spp., Elaphurus spp., Hippocamelus spp., Hydropotes spp., Mazama spp., Megamuntiacus spp., Muntiacus spp., Odocoileus spp., Ozotoceros spp., Pudu spp., Rangifer spp.</p> <p style="margin-left: 40px;">Giraffidae: Giraffa spp., Okapia spp.</p> <p style="margin-left: 40px;">Hippopotamidae: Hexaprotodon-Choeropsis spp., Hippopotamus spp.,</p> <p style="margin-left: 40px;">Moschidae Moschus spp.</p> <p style="margin-left: 40px;">Tragulidae Hyemoschus spp., Tragulus-Moschiola spp.,</p> <p style="margin-left: 40px;">Rhinocerotidae Ceratotherium spp., Dicerorhinus spp., Diceros spp., Rhinoceros spp.</p> <p style="margin-left: 40px;">Elephantidae Elephas spp., Loxodonta spp., as appropriate.</p>		
	Part II:		
	(1) Code of the territory as it appears in a document relating to 'live ungulates' published on gov.uk, in accordance with Regulation (EU) No 206/2010.(9)		
	(2) Keep as appropriate.		
	(3) 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 a document relating to 'live ungulates' published on gov.uk, in accordance with Regulation (EU) No 206/2010(9), with the entry "VII", as regards tuberculosis, "VIII", as regards brucellosis.		
	(4) Tests carried out in accordance with the protocols that, for the disease concerned, are described in Part 6 of Annex 1 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.		
	(5) Supplementary guarantees to be provided when required in column 5 "SG" of a document relating to 'live ungulates' published on gov.uk, in accordance with Regulation (EU) No 206/2010, with the entry "A".(9) Tests for Bluetongue and for Epizootic-haemorrhagic-disease in accordance with Part 6 of Annex 1 to Regulation (EU) No 206/2010.		
	(6) 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 Great Britain of the third country, territory or part thereof referred to in boxes reference I.7. and I.8., or during a period where restrictive measures have been adopted by Great Britain against imports of these animals from this third country, territory or part thereof.		
	(7) When required by Great Britain.		
	(8) Only for a territory appearing with the entry "XIII" in column 6 of a document relating to 'live ungulates' published on gov.uk, in accordance with Regulation (EU) No 206/2010(9), indicating an official bluetongue and epizootic haemorrhagic disease seasonally free status. In accordance with the WOA (formerly OIE) Terrestrial Animal Health Code, the seasonally free period is taken to conclude immediately if current climatic data or data from surveillance programme indicate an earlier resurgence of activity of adult Culicoides.		
	(9)A document relating to 'live ungulates' for EU and EFTA states published by the Secretary of State, with the consent of the Scottish and Welsh Ministers, may be found here: EU and EFTA states approved to export animals and animal products to Great Britain - data.gov.uk		
	Certifying Officer		
	Name (in capital letters)	Qualification and title	
	Date of signature	Signature	
	Stamp		