

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						
Production <input type="checkbox"/>		Pharmaceutical use <input type="checkbox"/>		Fattening <input type="checkbox"/>		
Other <input type="checkbox"/>		Production of petfood <input type="checkbox"/>		Human consumption <input type="checkbox"/>		
Breeding <input type="checkbox"/>		Relaying <input type="checkbox"/>		Animal Feedingstuff <input type="checkbox"/>		
Technical use <input type="checkbox"/>				Breeding and production <input type="checkbox"/>		
				Slaughter <input type="checkbox"/>		
				Artificial reproduction <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 _____ ISO Code _____			Country _____ 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						
0106 Other live animals						
Mammals:						
010619 Other						
01061900 Other						
Commodity	Species	Quantity	Net weight	Package count		
Identification number			Identification system			

Part II: Certification	II. Health information			
	II.1.	Animal health attestation		
		I, the undersigned official veterinarian responsible for the approved body, institute or centre/holding (1) of origin certify that the animals described in Part I meet the following requirements:		
	II.1.1.	They come from the country, territory or part thereof described in Box I.7.		
		(a)	where the diseases referred to in this certificate are notifiable,	
		(b)	which at the date of issuing this certificate has been free for the past 12 months from rinderpest.	
	II.1.2.	They come from the body, institute or centre/holding (1) described in Box I.11.		
		(a)	which is approved according to the requirements and conditions set out in Part 3 and 4 of Annex VI to Regulation (EU) No 206/2010;	
		(b)	which is not subjected to any restrictions relating to a national programme for the control of infectious diseases to which the animals referred to in Box I.25. are susceptible;	
		(c)	where there have been no clinical cases of the following diseases to which the animals referred to in Box I.28. are susceptible:	
		-	anthrax for the last 30 days;	
		-	foot-and-mouth disease, vesicular stomatitis, rabies, African swine fever, classical swine fever and swine vesicular disease for the past 6 months;	
		(d)	where there have been no clinical or non-clinical cases of tuberculosis and brucellosis for the past 6 months;	
		(e)	around which in an area of radius of 10 km for the last 12 months, there has been no case/outbreak of African swine fever, classical swine fever and swine vesicular disease;	
		(f)	around which in an area of 10 km radius for the past 30 days, there has been no case/outbreak of foot-and-mouth disease or vesicular stomatitis,	
	(1) ○ either	[(g)	in which they have remained since birth or for the past 6 months before dispatch to Great Britain.]	
	○ Or (1)	[(h)	have been introduced on (dd/mm/yyyy) into the territory described under point II.2.1, from the EU territory with code (5) that at that date was authorised to import the animals into Great Britain and the animals has not been in contact with imported cloven-hoofed animals from countries not subject to transitional import arrangements for the last 30 days.]	
	II.1.3.	They:		
		(a)	have not come into contact with other animals not complying with at least the same health requirements as described in this certificate since birth or for the last 30 days and during their transportation from the approved body, institute or centre/holding (1) to the place of shipment;	
		(b)	were examined by an official veterinarian within 24 hours of loading and showed no clinical sign of disease and are fit for the intended transport;	
		(c)	are not animals to be killed under a national programme for the eradication of diseases.	
	II.1.4.	Foot-and-Mouth Disease		
	either(1) ○ [(a)	They come from the country, territory or part thereof described in Box I.7. which at the date of issuing this certificate has been free for the past 12 months from foot-and-mouth disease and;]		
	or(1) ○ [(a)	They have been subjected to a virological and serological test for evidence of foot-and-mouth disease virus infection carried out in accordance with one of the prescribed tests for international trade laid down in the OIE Manual of Diagnostic Tests and Vaccines for Terrestrial Animals (OIE Terrestrial Manual), with negative results, taken in the 10 days prior to dispatch to Great Britain; and]		
		(b)	they have not been vaccinated against foot-and-mouth disease.	

Part II: Certification	II. Health information			
	(1)	II.1.5. either	Brucellosis ○ [They come from the country, territory or part thereof described in Box I.7 which has been free for the past 12 months from brucellosis and have not been vaccinated against that disease]	
	(1)(3)	or	○ [They have been subjected, with negative results, to a buffered Brucella antigen test for porcine brucellosis taken in the 30 days prior to dispatch to Great Britain.]	
	(1)	II.1.6. either	Swine vesicular disease ○ [They come from the country, territory or part thereof described in box 1.7 which has been free for the past 12 months from swine vesicular disease.]	
	(1)	or	○ [They have been subjected, with negative results, to a virology and serology test for evidence of swine vesicular disease, as laid down and prescribed for international trade by the OIE Terrestrial Manual, taken in the 30 days prior to dispatch to Great Britain.]	
	(1)	II.1.7. either	Vesicular Stomatitis ○ [They come from the country, territory or part thereof described in Box I.7 which has been free for the last 6 months from vesicular stomatitis.]	
	(1)	or	○ [They have been subjected, with negative results, to a virology and serology test for evidence of vesicular stomatitis, as laid down and prescribed for international trade by the OIE Terrestrial Manual, taken in the 30 days prior to dispatch to Great Britain.]	
	(1)	II.1.8. either	Classical swine fever ○ [They come from the country, territory or part thereof described in Box I.7 which has been free for the past 12 months from classical swine fever.]	
	(1)	or	○ [They have been subjected to a virological and serological test for classical swine fever carried out in accordance with one of the prescribed tests for international trade laid down in the OIE Terrestrial Manual, with negative results, taken in the 30 days prior to dispatch to Great Britain.]	
	(1)	II.1.9. either	African swine fever ○ [They come from the country, territory or part thereof described in Box I.7 which has been free for the past 12 months from African swine fever.]	
	(1)	or	○ [They have been subjected, with negative results, to a virus and serology test for African swine fever, as laid down and prescribed for international trade in the OIE Terrestrial Manual, taken in the 30 days prior to dispatch to Great Britain.]	
		II.1.10.	Aujeszky's disease According to official information, no clinical, pathological or serological evidence of Aujeszky's disease has been recorded for the last 12 months in the ○ approved body, institute or centre/ ○ holding (1) and in an area with a 5 km radius around the approved body, centre or institute, and They have been subjected, with negative results, to a virology and serology test for evidence of Aujeszky's disease, as laid down and prescribed for international trade by the OIE Terrestrial Manual, taken in the 30 days prior to dispatch to Great Britain, and They have not been vaccinated against Aujeszky's disease and have not been in contact with vaccinated animals.	
(2)	II.1.11. (a)	Other vaccinations They have not been vaccinated against rinderpest, vesicular stomatitis, classical swine fever or swine vesicular disease,		
(1)	(b)	They have been vaccinated against: <input type="checkbox"/> [anthrax on the (dd/mm/yyyy) with the following vaccine(s) (name of vaccine (s) used)],		
(1)		<input type="checkbox"/> [rabies on the (dd/mm/yyyy) with the following vaccine(s) (name of vaccine (s) used)].		
	II.1.12.	Parasite treatment		

Part II: Certification	II. Health information													
	<p>II.1.13. Loading on the means of transport</p> <p>They have been treated at least twice in the 40 days prior to dispatch to Great Britain against internal and external parasites with the following product(s) . Specify the active ingredients and the doses of the products used</p> <p>They have been loaded for dispatch to Great Britain on (dd/mm/yyyy)(4) in the means of transport described in Box 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.</p>													
	Notes													
	<p>(*) Those countries subject to the transitional import arrangements include: an EU member State; Liechtenstein; Norway and Switzerland.</p> <p>References to Great Britain in this certificate include Channel Islands and Isle of Man. 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)</p> <p>This certificate is meant for animals of species listed in the note for Box I. 25. coming from an approved body, institute or centre in a third country, territory or part thereof, and destined to an approved body, institute or centre located within Great Britain.</p>													
	Part I:													
	<p>Box reference I.15.:</p> <p>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 shall inform the BCP of entry into Great Britain.</p>													
	<p>Box reference I.16.:</p> <p>Do not use this box until the end of the transitional staging period.</p>													
	<p>Box reference I.25.:</p> <p>Identification system: Specify the identification system (tag, tattoos, brand, chip, transponder). The identifier shall include the ISO code of the exporting country and permit tracing of their premises of origin.</p> <p>Age: months.</p> <p>Sex (M = male, F = female, C = castrated).</p> <p>Species Select the species amongst those listed below:</p> <table border="0"> <thead> <tr> <th>Order</th> <th>Family</th> <th>Genera/species</th> </tr> </thead> <tbody> <tr> <td>Artiodactyla</td> <td>Suidae</td> <td>Babyrousa ssp., Hylochoerus ssp., Phacochoerus ssp., Potamochoerus ssp., Sus ssp.</td> </tr> <tr> <td></td> <td>Tayassuidae</td> <td>Catagonus ssp., Pecari-Tayassu ssp.</td> </tr> <tr> <td></td> <td>Hippopotamidae</td> <td>Hexaprotodon-Choeropsis , Hippopotamus spp.</td> </tr> </tbody> </table>	Order	Family	Genera/species	Artiodactyla	Suidae	Babyrousa ssp., Hylochoerus ssp., Phacochoerus ssp., Potamochoerus ssp., Sus ssp.		Tayassuidae	Catagonus ssp., Pecari-Tayassu ssp.		Hippopotamidae	Hexaprotodon-Choeropsis , Hippopotamus spp.	
Order	Family	Genera/species												
Artiodactyla	Suidae	Babyrousa ssp., Hylochoerus ssp., Phacochoerus ssp., Potamochoerus ssp., Sus ssp.												
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	Hippopotamidae	Hexaprotodon-Choeropsis , Hippopotamus spp.												
	Part II:													
	(1) Keep as appropriate.													
	(2) Vaccination is not compulsory, but if the animals have been vaccinated, information on the vaccine(s) used and the time of vaccination must be filled in.													
	(3) 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.													
	(4) 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 country, territory or part thereof described in Boxes 1.7. and 1.8., or during a period where restrictive measures have been adopted by Great Britain against imports of these animals from that country, territory or part thereof.													
	(5) Code of the territory as it appears in Part 1 of Annex 1 to Regulation (EU) No 206/2010.													
	Certifying Officer													

Part II: Certification	II. Health information			
	Name (in capital letters)	Qualification and title		
	Date of signature	Signature		
	Stamp			