

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						
Registered equidae <input type="checkbox"/> Unregistered equidae <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.24. Total quantity			I.25. Total gross weight			
I.28. Description of consignment						
Commodity	Species	Identification system	Identification number	Age		
Gender			Quantity			

Part II: Certification	II. Health information		
	I, the undersigned official veterinarian, hereby certify, that the animal described in Box I.28.:		
	○ either(1)	[—	is a registered equine as defined in Article 2(c) of Commission Implementing Regulation (EU) 2018/659;]
	○ or(1)	[-	an unregistered equine;]
		—	was examined today and found free of clinical signs of disease and of obvious signs of ectoparasite infestation;
		—	is not intended for slaughter under a national programme of infectious or contagious disease eradication;
	○ either(1)	[—	does not come from the territory or part of the territory of a Member State or Norway which is the subject of restrictions for reasons of African horse sickness;]
	○ or(1)	[—	it comes from the territory or part of the territory of a Member State or Norway, which is subject to restrictions for reasons of African horse sickness, has remained for at least 40 days prior to dispatch in the vector proved quarantine station of (insert name of quarantine station) and has undergone a test for the detection of antibodies to the African horse sickness virus as described in Annex IV to Directive 2009/156/EC carried out simultaneously on blood samples taken on two occasions with an interval of between 21 and 30 days on (insert date) and during the 10 days prior to dispatch on (insert date)
		○ either(1)	[with negative result in each case if it was not vaccinated against African horse sickness;]
		○ or(1)	[without an increase in antibody count if it was vaccinated against African horse sickness;]
○ either(1)	[—	was not vaccinated against African horse sickness;]	
○ or(1)	[—	was vaccinated against African horse sickness on (insert date);	
	○ either(1)	[at least two months prior to certification]	
	○ or(1)	[at least two months prior to entry into the quarantine station;]	
	—	has not been obtained from a holding which was subject to prohibition for animal health reasons, which laid down at least one of the following conditions:	
	○ either(1)	[not all animals on the holding of species susceptible to the diseases mentioned hereafter were slaughtered and the prohibition lasted for at least:	
	(a)	in the case of equidae suspected of having contracted dourine	
	○ either(1)	[six months beginning on the date of the last actual or possible contact with a sick or infected with Trypanosoma equiperdum animal;]	
	○ or(1)	[in the case of a stallion until the animal is castrated;]	
	(b)	in the case of glanders, six months beginning on the day on which the equidae suffering from the disease or subjected with positive result to a test for the detection of the causative pathogen Burkholderia mallei or antibodies to that pathogen, were killed and destroyed;	
	(c)	in the case of equine encephalomyelitis of any type, six months beginning on the day on which the equidae suffering from the disease have been slaughtered, except in case of West Nile virus infection, where the period of six months begins on the day the equidae died, have been removed from the holding or fully recovered;	
	(d)	in the case of equine infectious anaemia, until the date on which, the infected animals having been slaughtered, the remaining animals have shown a negative reaction to two Coggins tests carried out three months apart;	
	(e)	in the case of vesicular stomatitis, six months from the last case;	

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	<p>(f) in the case of rabies, one month from the last case;</p> <p>(g) in the case of anthrax, 15 days from the last case.]</p> <p>○ or(1) [following cases of dourine, glanders, equine encephalomyelitis of all types, equine infectious anaemia, vesicular stomatitis, rabies or anthrax, all animals on the holding of species susceptible to the disease in question were slaughtered or killed and the prohibition lasted for 30 days or 15 days in the case of anthrax, beginning on the day on which, following the destruction of the animals, the disinfection of the premises, was satisfactorily completed;]</p> <p>— to the best of my knowledge, it has not been in contact with equidae suffering from an infectious or contagious disease in the 15 days prior to this declaration;</p> <p>— at the time of the inspection, it was fit to be transported on the intended journey in accordance with the provisions of Regulation (EC) No 1/2005.</p> <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>Part I:</p> <p>Box I.15.: Registration number (railway wagons or container and lorries), flight number (aircraft) or name (ship) and other relevant information is to be provided. In case of unloading and reloading, the consignor must inform the competent authority of GB.</p> <p>Box I.16.: Do not use this box until the end of the transitional staging period.</p> <p>Box I.23.: The container number and the seal number (if applicable) should be included.</p> <p>Box I.28.: Identification system: The animal must bear an individual identifier which permits linking the animal to the identification document as defined in Article 2(b) of Commission Implementing Regulation (EU) 2018/659. Specify the identification system (such as ear tag, tattoo, brand, transponder) and the anatomic place used on the animal. The number of the accompanying passport must be stated and the name of the competent authority which validated it.</p> <p style="padding-left: 40px;">Age: Date of birth (dd/mm/yyyy).</p> <p style="padding-left: 40px;">Sex (M = male, F = female, C = castrated).</p> <p>Part II:</p> <p>(1) Delete as appropriate.</p> <p>This health certificate shall:</p> <p>(a) be issued on the day of loading [for any equine] or on the last working day before loading [for registered equines only] of the animal for dispatch to GB;</p> <p>(b) be drawn up in at least a language understood by the certifying officer and in English;</p> <p>(c) be made out to a single consignee;</p> <p>(d) be signed and stamped in a colour different to the colour of the printing;</p> <p>(e) consist of a single sheet of paper or all sheets of paper required are part of an integrated whole and indivisible by inserting page numbers and the total number of pages, and each page shall bear the certificate reference number at the top of the page and those pages are stapled and stamped.</p>		
Certifying Officer			
Name (in capital letters)		Qualification and title	
Date of signature		Signature	
Stamp			