## EUROPEAN UNION

	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						
ιt	Name				I.4. Local com	petent authority					
Jer	Address										
nn	Country		ISO Code								
Part I : Details of consignment	I.7. Country of orig	gin		ISO Code	I.9. Country of	f destination		ISO Code			
on											
ofc	I.8. Region of origi	n		Code	I.10. Region of destination						
ls (	I.11. Place of Dispa				I.12. Place of destination						
tai	Name				Name						
De	Address				Address						
	Approval Number	2			Approval Nu	mber					
t	Country		ISO Code		Country ISO Code						
Ра	I.13. Place of Load	ing			I 14 Date and	time of departure					
	Name	ing			1.14. Dute und	unite of departure					
	Address										
	Approval Number										
	Country		ISO Code								
	7.45. X ( ) ( )				MAGE - D						
	I.15. Means of Trai				I.16 Entry Poi	nt					
	Mode	International transport	Identification								
		document			-						
					-						
					-						
	I.18. Transport cor	ditions			I.17. Accompa	nying documents					
	Ambient 🗋				Commercial						
					document reference	Dat	e of issue				
					Country	Pla	ce of				
					country	issu	le				
	I.19. Container No	/ Seal No									
	I.20. Certified as										
	Registered equidae	e 🗆	Unregistered equida	e 🗆							
	I.21. For transit the	rough a third cou	ntry 🛛 ISO Code		1.22. For trans	it through Member State(s)					
	Country EU Exit										
	Authority		BCP code		Country	ISO	Code				
	EU Entry Authority		BCP code								
	I.24. Total quantity	1			I.25. Total gross weight						
	_	I.28. Description of consignment									
	1.01 LIVE ANIMA										
	0101 Live horses, asses, mules and hinnies					1					
	Commodity	Speci	es	Identification	system	Identification number	Age				
					1						
	Gender				Quantity						

## **EUROPEAN UNION**

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	II. Health information										
	I, the undersigned official veterinarian, hereby certify, that the animal described in Box I.28.:										
	o either(1)	[—	is a registered equine as defined in Article 2(c) of Commission Implementing Regulation (EU) 2018/659;]								
	• or(1)	[-	an unregis	tered equin	le;]						
tion		—	was examined today and found free of clinical signs of disease and of obvious signs of ectoparasite infestation;								
Part II: Certification		—		ot intended for slaughter under a national programme of infectious or contagious case eradication;							
II: Cej	o either(1)	[—		s not come from the territory or part of the territory of a Member State or Norway ch is the subject of restrictions for reasons of African horse sickness;]							
Part	○ or(1)	[—	subject to a days prior of quarant horse sicka simultane	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							
			o either(1)	(insert c [with neg sickness;]]	ative result in each case if it was not vaccinated against African horse						
			• or(1)		an increase in antibody count if it was vaccinated against African						
	o either(1)	[—	was not v		gainst African horse sickness;]						
	• or(1)	[—	was vacci	nated again	st African horse sickness on (insert date);						
			o either(1)	[at least ty	wo months prior to certification] ]						
			• or(1)	[at least ty	wo months prior to entry into the quarantine station;] ]						
		_			d from a holding which was subject to prohibition for animal health own at least one of the following conditions:						
			o either(1)		nimals on the holding of species susceptible to the diseases mentioned were slaughtered and the prohibition lasted for at least:						
				(a)	in the case of equidae suspected of having contracted dourine						
				o either(1)	[six months beginning on the date of the last actual or possible contact with a sick or infected with Trypanosoma equiperdum animal;]						
				o 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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Lifesth information       (f)       in the case of rabies, one month from the last case;       (g)       in the case of rabies, one month from the last case;         (g)       in the case of rabies, one month from the last case;       (g)       in the case of rabies, one month from the last case;         (g)       in the case of rabies, one month from the last case;       (g)       in the case of rabies, one month from the last case;         (g)       in the case of rabies, one month from the last case;       (g)       in the case of anthrax, all animals on the holding of species susceptible to the disease in question were saluptered 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:]         -       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;         -       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.         Notes:       References to European Union legislation which 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 gov.uk).         References to Great Britain in this certificate include Channel Islands and Isle of Man.         Part I:       Box 116:       Do not use this b	_										~,	
<ul> <li>(g) in the case of anthrax, 15 days from the last case.]</li> <li>o or(1) [following cases of dourine, glanders, equine encephalomyelitis of all types, equine infectious anamenia, 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;]</li> <li> <ul> <li>to the best of my knowledge, it has not been in contact with equidea suffering from an infectious or contagious disease in the 15 days prior to this declaration:</li></ul></li></ul>		II. Health info	rmation									
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<ul> <li>(c) be made out to a single consignee;</li> <li>(d) be signed and stamped in a colour different to the colour of the printing;</li> <li>(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.</li> <li>Certifying Officer</li> <li>Name (in capital letters)</li> <li>Qualification and title</li> <li>Date of signature</li> </ul>		(a)							before load	ling [for		
<ul> <li>(d) be signed and stamped in a colour different to the colour of the printing;</li> <li>(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.</li> <li>Certifying Officer</li> <li>Name (in capital letters)</li> <li>Date of signature</li> <li>Qualification and title Signature</li> </ul>		(b)	be drawn	up in at lea	st a language under	rstood	l by the cer	tifying officer and	in English;	,		
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		Date of signa						and title				