CHAPTER 4(A)

Health certificate

For the import of blood and blood products from equidae to be used outside the feed chain, for dispatch to or for transit through $^{(2)}$ the European Union

COUNTRY: UNITED STATES

Veterinary certificate to EU

	I.1. Consignor Name Address	I.2. Certificate reference No I.2.a.		
		I.3. Central competent authority APHIS-VS		
	Tel.	I.4. Local competent authority		
	1.5. Consignee Border inspection post through which consignment is intended to leave the EU	I.6. Person responsible for the load in EU Name Address Postal code Tel.		
Part I : Details of dispatched consignment	Name Address			
	Postal code Tel.			
	I.7. Country of ISO code I.8. Region of Code origin US US-0	I.9. Country of ISO I.10. Region of eode destination code destination		
atche	I.11. Place of origin	I.12. Place of destination		
s of disp	Name Approval number Address	Custom warehouse Name Approval number Address		
Detail		Postal code		
Part I:				
	Name Approval number Address			
	Name Approval number Address			
	I.13. Place of loading	I.14. Date of departure		
	I.15. Means of transport	I.16. Entry BIP in EU		
	Aeroplane □ Ship □ Railway wagon □ Road vehicle □ Other □	1.17.		
	Identification			
	Documentation references			
	I.18. Description of commodity	I.19. Commodity code (HS code)		
		I.20. Quantity		
	I.21. Temperature of product	I.22. Number of packages		
	Ambient ☐ Chilled ☐ Frozen ☐			

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COUNTRY: UNITED STATES

Blood and blood products from equidae for purposes outside the feed chain

	I.2. Certificate reference No	I.2.a.
I.23. Seal/Container No	I.24. Type of packaging	
I.25. Commodities certified for:	_	
Technical use □		
I.26. For transit through EU to third country	I.27. For import or admission into EU	
Third country ISO code		
I.28. Identification of the commodities		
Species (Scientific name)	Approval number of establishments Manufacturing plant	

Page ____ of ____

II.	Health information	II.a. Certificate reference No	II.b.			
	I, the undersigned official veterinarian, declare that I have read and understood Regulation (EC) No 1069/2009 of the European Parliament and of the Council ^(1a) and in particular Article 8(c) and Article 8(d) and Article 10 thereof, and Commission Regulation (EU) No 142/2011 ^(1b) , and in particular Chapter IV of Annex XIII thereto, and certify that the blood or blood products of equidae described above:					
II.1. II.2.	consist of blood or blood products from equida consist exclusively of blood or blood products		·			
II.3.	have been obtained from animals that originate from the EU Member States or from a third country, territory or part thereof listed in the column 'third countries' lists' of row No 3 of Table 2 in Section 1 of Chapter II of Annex XIV to Regulation (EU) No 142/2011 where the following diseases are compulsorily notifiable: African horse sickness, dourine, glanders (<i>Burkholderia mallei</i>), equine encephalomyelitis (all types including Venezuelan equine encephalomyelitis), equine infectious anaemia, vesicular stomatitis, rabies, anthrax;					
II.4.	have been derived from blood from equidae, which was collected under the supervision of a veterinarian in slaughterhouses approved in accordance with Regulation (EC) No 853/2004 of the European Parliament and of the Council ⁽³⁾ , in slaughterhouses approved and supervised by the competent authority of the country of collection and in facilities approved and supervised by the competent authority of the country of collection for the purpose of collecting blood from equidae for the production of blood products for purposes other than feeding for farmed animals;					
II.5.	have been derived from blood which was colle	cted from equidae:				
II.5.1.	which on inspection on the date of blood collection did not show clinical signs of any of the compulsorily notifiable diseases listed in Annex I to Council Directive 2009/156/EC ⁽⁴⁾ , and of equine influenza, equine piroplasmosis, equine rhinopneumonitis and equine viral arteritis listed in point 4 of Article 1.2.3 of the Terrestrial Animal Health Code of the World Organisation for Animal Health (OIE), 2010 edition:					
II.5.2.	which have been kept for at least 30 days prior to the date of and during blood collection on holdings under veterinary supervision which were not subject to a prohibition order pursuant to Article 4(5) or restrictions for African horse sickness in accordance with Article 5 of Directive 2009/156/EC:					
II.5.3.	which had no contact with equidae from a holding which was subject to a prohibition order for animal health reasons pursuant to Article 4(5) of Directive 2009/156/EC;					
II.5.4.	for which the period for the prohibition order follows: (2) either [not all the animals of species]	referred to in points II.5.2. and II.5.3 has susceptible to the disease located on the				
		eriod of prohibition must be at least: anders (<i>Burkholderia mallei)</i> , beginning on t sease are slaughtered	the date on which t			
	- six months in the case of eq	uine encephalomyelitis of any type, including on the date on which the equidae infected				
		ous anaemia, until the date on which, the interemental remaining animals have shown a negate months apart,				
	 six months from the date of t 	the last recorded case of vesicular stomatitis	s,			
	 one month from the date of t 	the last recorded case of rabies,				
	 15 days from the date of the 	last recorded case of anthrax;]				
	slaughtered and the premises we days, beginning on the date on w	sceptible to the disease located on the re disinfected, in which case the period of particle that the sanimals were slaughtered and the ere the period of prohibition shall be 15 days	prohibition must be premises disinfecte			
II.6.	blood products come from an establishment or plant approved or registered by the competent authority of the third country meeting the specific conditions set out in Article 23 or 24 of Regulation (EC) No 1069/2009;					
II.7.	since birth if less than three mont	d which fulfils the conditions referred in II.4 s which have been kept for a period of at length of the have been kept for a period of at length of the half of collection on hole period and the period	east three months, dings under vetering			
	(a) African horse sickness for (b) Venezuelan equine encept	two years; halomyelitis for a period of at least two year	e:			
	(c) glanders		J ,			
	(2) either [for a period of the	rree years;]				

COUNTRY: UNITED STATES

Blood and blood products from equidae for purposes outside the feed chain

In the case of blood, the approximation of the book and blood products with pathogenic agents during production, handling and packagin.	II.	Health info	ormation	II.a.	Certificate reference No	II.b.
inspection for glanders in the slaughterhouse referred to in It.4, including a careful examination of mucous membranes from the trahele, larynx, nasal cavities and sinuses and their ramifications, after splitting the head in the median plane and existing the nasal septum;] (d) in the case of blood-products other than serum—and plasma, vesicular stomatitis for six months;]] (a) In the case of blood-products other than serum—and plasma, vesicular stomatitis for six months;]] (b) In the case of blood-products of all types including Venezuelan equipe encephalomyelitis, equine infectious anaemia, vesicular stomatitis and glanders (Burkholderia malloi): (b) In the term (heat treatment at a temperature of 56°C for at least three hours;] (c) and/or—[tradiation at 25 KQy by gamma rays;] (d) and/or—[tradiation at 25 KQy by gamma rays;] (e) and/or—[theat treatment of at least 80°C throughout their substance;]] (ii) and/or—[theat treatment of at least 80°C throughout their substance;]] (iii) blood and blood products were packed in sealed impermeable containers clearly labelled 'NOT FOR HUMAN OR ANIMAL CONSUMPTION' and bearing: (b) in the case of blood, the approval number of the establishment of collection; (c) in the case of blood, the approval number of the establishment of production; (iii) in the case of blood products, the approval number of the establishment of production; (iii) in the case of blood products, the approval number of the establishment of production; (b) in the case of blood products, the approval number of the establishment of production; (c) in the case of blood products, the approval production; (d) in the case of blood products, the approval production in the surface of the approval number of the establishment of production; (d) in the case of blood products, the approval number of the establishment of production in the approval number of the approval number of the establishment of production in the continuation of the products in transit con only be stored in the continuation of th						
cavities and sinuses and their ramifications, after splitting the head in the median plane and existing the nasal septum;] (d)—in the case of blood products other than serum and plasma, vesicular stomatitis for six menethe;]] (has been subjected to at least one of the following treatments, followed by an effectiveness check, for the inactivation of possible causative pathogens for African horse sickness, equine encephalomyelitis of all types including Venezuelan equine encephalomyelitis, equine infectious anaemia, vesicular stomatitis and glanders (Burkholderia mailes): (**) (*			inspection for o	glander	s in the slaughterhouse referred to in	II.4, including a
(d) — in. the case-of-blood-products-other-than-serum and-plasma, vesicular stomatitis-for-six monthe;]] (d) — [has been subjected to at least one of the following treatments, followed by an effectiveness check, for the inactivation of possible causative pathogens for African horse sickness, equine encephalomyetitis of all types including Venezuelan equine encephalomyetitis, equine infectious anaemia, vesicular stomatitis and glanders (Burkholderia malle): (a) either [hast treatment at a temperature of 65°C for at least three hours;] (a) and/or [lenat treatment at a temperature of 65°C for at least three hours;] (a) and/or [heat treatment of at least 80°C throughout their substance;]] (b) and blood products were packed in sealed impermeable containers clearly labelled 'NOT FOR HUMAN OR ANIMAL CONSUMPTION' and bearing; (a) —in the case of blood, the approval number of the establishment of collection; (b) in the case of blood, the approval number of the establishment of production; (b) in the case of blood products, the approval number of the establishment of production; (b) in the case of blood, the approval number of the establishment of production; (b) in the case of blood, the approval number of the establishment of production; (b) in the case of blood, the approval number of the establishment of production; (c) and the products were stored in enclosed storage. (a) —in the case of blood, the experiment in the European Union: this box is to be filled in only if it is a certificate for transit commodity, it may be filled in at the certificate is for import commodity. Box reference 1.13 and 1.12. Approval number the registration number (rainer to transit commodity). It may be filled in a the certificate is for import commodity. Box reference 1.15 and 1.12. Approval number the registration number (rainer to transit commodity). It may be filled in the certificate is to fill production. Box reference 1.15 and 1.12. Approval number the registration number (rainer). The products in the consignor must		cavities and sinuses and their ramifications, after splitting the head in the median				
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II.8. all precautions have been taken to avoid contamination of the blood and blood products with pathogenic agents during production, handling and packaging; II.9. blood and blood products were packed in sealed impermeable containers clearly labelled 'NOT FOR HUMAN OR ANIMAL CONSUMPTION' and bearing: (a) in the case of blood, the approval number of the establishment of collection; (b) in the case of blood products, the approval number of the establishment of production; the products were stored in enclosed storage. Notes Part: Box reference 1.6: Person responsible for the consignment in the European Union: this box is to be filled in only if it is a certificate for transit commodity; it may be filled in if the certificate is to import commodity. Box reference 1.1: 2-Place of destination: this box is to be filled in only if it is a certificate for transit commodity. The products were stored in enclosed storage. Box reference 1.1: 2-Place of destination: this box is to be filled in only if it is a certificate for transit commodity. The products in transit can only be stored in free zones, free warehouses and custom wenthouses. Box reference 1.1: 5: Registration number (railway wapons or container and lorries), flight number (aircraft) or name (ship) is to be provided. In case of unloading and releading, the consigion mast inform the BiP of entry into the EU. Box reference 1.2:3: For busined use: any use other than for animal consumption. Box reference 1.2:3: For busined use: any use other than for animal consumption. Box reference 1.2:3: (a) in according to whether it is a transit or an import certificate. Box reference 1.2:3: (a) in a coording to whether it is a transit or an import certificate. Box reference 1.2:3: (a) in the case of blood products, provide the approval number of the registered establishment of collection; (ii) in the case of blood products, provide the approval number of the establishment of production (iii) in the case of blood products, provide the approval number of the self-				nt at a temperature of 65°C for at least three hours;]		
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Notes Part I: Box reference I.6: Person responsible for the consignment in the European Union: this box is to be filled in only if it is a certificate for transit commodity; it may be filled in if the certificate is for import commodity. Box reference I.1 and 1/12: Approval number: the registration number of the establishment or plant, which has been issued by the competent authority. Box reference I.12: Place of destination: this box is to be filled in only if it is a certificate for transit commodity. The products in transit can only be stored in free zones, five warehouses and custom warehouses. Box reference I.15: Registration number (railway wagons or container and fornies), 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 EU. Box III by use the appropriate Harmonized System (IIIs) code under the following heading: 30.02. Box reference I.23: for bulk containers, the container number and the seal number (if applicable) must be included. Box reference I.25: exchincial use: any use other than for animal consumption. Box reference I.26 and I.27: fill in according to whether it is a transit or an import certificate. Box reference I.26: Manufacturing plant: (i) in the case of blood, provide the approval number of the registered establishment of collection.; (ii) in the case of blood products, provide the approval number of the registered establishment of production (b) Species: select amongst the following: Equus cabalus, Equus asinus, Equus cabalus*asinus. Part II: Jol L 30, 14.11.2009, p. 1. Jol L 39, 30.4.2004, p. 55. Jol L 139, 30.4.2004, p. 55. Jol L 192, 23.7.2010, p. 1. The signature and the stamp must be in a different colour to that of the printing. Note for the person responsible for the consignment in the European Union: this certificate is only for veterinary purposes and must accompany the consignment until it reaches the border inspection post. Official veterinarian/Official inspect	11.40	, ,			nber of the establishment of production;	
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