

COUNTRY:

Veterinary certificate to EU

Part I: Details of dispatched consignment	1.1. Consignor Name Address Tel.				1.2. Certificate reference No		1.2.a					
					1.3. Central competent authority							
					1.4. Local competent authority							
	1.5. Consignee Name Address Postal code Tel.				1.6. Person responsible for the load in EU Name Address Postal code Tel.							
	1.7. Country of origin		ISO code	1.8. Region of origin		Code	1.9. Country of destination		ISO code	1.10. Region of destination	Code	
	1.11. Place of origin Name Address Name Address Name Address				Approval number Approval number Approval number				1.12. Place of destination Name Address Postal code			
	1.13. Place of loading				1.14. Date of departure							
	1.15. Means of transport Aeroplane <input type="checkbox"/> Ship <input type="checkbox"/> Railway wagon <input type="checkbox"/> Road vehicle <input type="checkbox"/> Other <input type="checkbox"/> Identification Documentary references				1.16. Entry BIP in EU				1.17.			
	1.18. Description of commodity						1.19. Commodity code (HS code) 05 11 99 85					
							1.20. Quantity					
	1.21.						1.22. Number of packages					
	1.23. Seal/container No						1.24.					
	1.25. Commodities certified for: Artificial reproduction <input type="checkbox"/>											
	1.26. For transit through the EU to a third country <input type="checkbox"/> Third country ISO code					1.27. For import or admission into the EU <input type="checkbox"/>						
	1.28. Identification of the commodities											
Species (scientific name)		Category	Donor identity		Date of collection		Approval number of the team		Quantity			

COUNTRY:

Equine ova/embryos

II. Health information	II.a. Certificate reference No	II.b.
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I, the undersigned, official veterinarian, of the exporting country (2) hereby
(name of exporting country)

Part II: Certification

certify that:

- II.1. The ova (1)/embryos (1) described above:
- II.1.2. were collected (1)/produced (1) by the team (2) described in Box I.11, which has been approved and supervised in accordance with Chapter I(III) of Annex D to Directive 92/65/EEC and is subject to inspection by an official veterinarian at least once every calendar year;
- II.1.3. were collected (1)/produced (1), processed and stored in accordance with the requirements of Chapter III(II) of Annex D to Directive 92/65/EEC;
- II.1.4. were collected at a place separated from other parts of the premises or holding which is in good repair and was cleaned and disinfected prior to the collection;
- II.1.5. were examined, processed and packed in laboratory facilities which are not situated in a zone subject to prohibition or quarantine measures as set out in Box II.1.6, in a section which is separated from the section for storing equipment and materials used in contact with donor animals and from the area where the donor animals are handled;
- II.1.6. come from donor mares which:
 - II.1.6.1. were continuously resident for three months (or since entry if they were directly imported from a Member State of the European Union during the three month period) in the exporting country or, in the case of regionalisation according to Article 13 of Directive 2009/156/EC (6), in that part of the territory of the exporting country which was during that period:
 - not considered to be infected with African horse sickness in accordance with Article 5(2)(a) and (b) of Directive 2009/156/EC (6).
 - free from Venezuelan equine encephalomyelitis for at least two years,
 - free from glanders and dourine for at least six months;
 - (1) either [II.1.6.2. originated from a country of export which was on the day of collection free of vesicular stomatitis for at least six months;]
 - (1) or [II.1.6.2. were tested by a virus neutralisation test for vesicular stomatitis on a blood sample taken on (4) within 30 days prior to collection, with negative result at a serum dilution of 1 in 12;]
 - (1) either [II.1.6.3. during the past 30 days prior to collection have been located in holdings under veterinary supervision which fulfilled from the day of collection of ova (1)/embryos (1) until the date of their dispatch the conditions for a holding laid down in Article 4(5) of Directive 2009/156/EC (6), and in particular;]
 - (1) or [II.1.6.3. during the past 30 days prior to collection have been located in holdings under veterinary supervision which fulfilled from the day of collection of ova (1)/embryos (1) until, in the case of frozen ova (1)/embryos (1), the period of 30 days mandatory storage at approved premises elapsed the conditions for a holding laid down in Article 4(5) of Directive 2009/156/EC (6) and in particular;]
 - (1) either [II.1.6.3.1. not all the animals of species susceptible to the disease located on the holding were slaughtered or killed and the holding has been free:
 - from any type of equine encephalomyelitis for at least six months, beginning on the day on which the equidae suffering from the disease are slaughtered,
 - from equine infectious anaemia for at least the period required to obtain a negative result in an agar gel immunodiffusion test (Coggins test) carried out on samples taken after the infected animals were slaughtered on two occasions three months apart from each of the remaining equidae,
 - from vesicular stomatitis for at least six months from the last recorded case,
 - from rabies for at least one month from the last recorded case,
 - from anthrax for at least 15 days from the last recorded case]
 - (1) or [II.1.6.3.1. all the animals of species susceptible to the disease located in the holding have been slaughtered or killed and the premises disinfected, the holding has been free for at least 30 days from any type of equine encephalomyelitis, vesicular stomatitis and rabies 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;]

COUNTRY:

Equine ova/embryos

II. Health information	II.a. Certificate reference No	II.b.
II.1.6.4.		
II.1.6.5.		
II.1.6.6.		
II.1.6.7.		
II.1.6.8.		
II.1.6.9.		
II.1.7.		
II.1.8.		
II.2.		
II.3.		
Notes		
Part I:		
Box I.11: place of origin shall correspond to the embryo collection team or embryo production team by which the ova/embryos were collected/produced, processed, stored and approved in accordance with Article 17(3)(b) of Directive 92/65/EEC and listed on the Commission website: http://ec.europa.eu/food/animal/semen_ova/equine/index_en.htm		
Box I.22: number of packages shall correspond to the number of containers.		
Box I.23: identification of container and seal number shall be indicated.		
Box I.28. category: specify if <i>in vivo</i> derived embryos, <i>in vivo</i> derived ova, <i>in vitro</i> produced embryos or micromanipulated embryos.		
donor identity shall correspond to the official identification of the animal.		
date of collection shall be indicated in the following format: dd/mm/yyyy.		
approval number of the team: shall correspond to the embryo collection team or embryo production team by which the ova/embryos were collected/produced, processed, stored and approved in accordance with Article 17(3)(b) of Directive 92/65/EEC and listed on the Commission website: http://ec.europa.eu/food/animal/semen_ova/equine/index_en.htm		

