

Part I : Details of consignment	I.1. Consignor		I.2. IMSOC Reference		
	Name				
	Address		I.2.a. Local Reference		
	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		ISO Code	I.9. Country of destination	
				ISO Code	
I.8. Region of origin		Code	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	ISO Code	Country		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			
Chilled <input type="checkbox"/>	Controlled temperature <input type="checkbox"/>	Ambient <input type="checkbox"/>	Frozen <input type="checkbox"/>	Commercial document reference	
				Date of issue	
				Country	
				Place of issue	
I.19. Container No / Seal No					
I.20. Certified as					
Artificial reproduction <input type="checkbox"/>		Ornamental use/research <input type="checkbox"/>		Pollination <input type="checkbox"/>	
Consignments according to Regulation No 999/2001 <input type="checkbox"/>		Ornamental bird food <input type="checkbox"/>		Registered equidae <input type="checkbox"/>	
Breeding and production <input type="checkbox"/>		Racing <input type="checkbox"/>		Circus exhibition <input type="checkbox"/>	
Training <input type="checkbox"/>		Pets <input type="checkbox"/>		Category 3 fish oil/fish meal for detoxification according to Regulation 2015/786 <input type="checkbox"/>	
Rodent food <input type="checkbox"/>		Technical use <input type="checkbox"/>		Further process <input type="checkbox"/>	
Sales <input type="checkbox"/>		Storage <input type="checkbox"/>		Pharmaceutical use <input type="checkbox"/>	
Relaying <input type="checkbox"/>		Other <input type="checkbox"/>		Animal Feedingsstuff <input type="checkbox"/>	
Competition <input type="checkbox"/>		Human consumption <input type="checkbox"/>		Organic fertilizers <input type="checkbox"/>	
Fattening <input type="checkbox"/>		Production of petfood <input type="checkbox"/>		Quarantine <input type="checkbox"/>	
				Pet food <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		

Part I : Details of consignment	I.28. Description of consignment				
	Commodity	Species	Quantity	Batch number	Manufacturing plant
	Cold store	Cutting plant	Date of freezing	Date of production	Date of slaughter
	Net weight	Product Description	Package count	Identification mark	

Part II: Certification	<p>II. Health information</p>																										
	<p>Section A (to be signed in Section IV by the Semen Collection Center Veterinarian);</p> <p>I, the undersigned, Center Veterinarian of the described semen collection center, hereinafter "SCC", certify that:</p>																										
	<p>II.1. The semen described in this certificate:</p>																										
	<p>II.1.1. was collected in a Member State that is recognized by the USDA to be free of African horse sickness, in accordance with the list of USDA-recognized animal health status of countries/area as listed on the internet at:</p> <p>http://www.aphis.usda.gov/import_export/animals/animal_disease_status.shtml</p> <p>and was free of this disease at the time of semen collection;</p>																										
	<p>II.1.2. was collected and processed in a semen collection center, hereinafter "SCC", that was approved by the competent authority of the EU Member State in which the semen is collected and processed in accordance with Directive 92/65/EEC and under the supervision of the center veterinarian in charge of it;</p>																										
	<p>II.1.3. was collected and processed using equipment that was new or that has been cleaned and sanitized prior to use under the supervision of the center veterinarian;</p>																										
	<p>II.1.4. was collected after the first set of specimens described under point III.1.6. was cultured negative;</p>																										
	<p>II.1.5. was processed with semen extender containing</p>																										
	<p>(1)either <input type="checkbox"/> [milk originated from a country or a region recognized at the time of semen collection by the USDA as free of foot-and-mouth disease and rinderpest as listed in 9 CFR Part 94 and other publications;]</p>																										
	<p>(1)and/or <input type="checkbox"/> [egg originated from a country or a region recognized at the time of semen collection by the USDA as free of Newcastle disease as defined in the Article II.9.3 of the OIE Terrestrial Animal Health Code, 20th Edition, 2011 and not affected with highly pathogenic influenza H5N1 as listed in 9 CFR Part 94 and other publications;]</p>																										
	<p>II.1.6. was placed in individual ampoules or straws which are permanently marked with the identification of the donor, the date of collection and the name or approval number of the SCC as recorded in Section C;</p>																										
	<p>II.1.7. was maintained after processing under lock and key or in the custody of the center veterinarian in a segregated storage area in the SCC until it was placed in a shipping container that is new or that has been cleaned and disinfected and, for frozen semen, was charged only with virgin liquid nitrogen;</p>																										
	<p>III.1. The donor stallion(s):</p>																										
	<p>III.1.1. has/have been free from any quarantine or movement restrictions for not less than 60 days prior to semen collection;</p>																										
	<p>III.1.2. was/were not used for natural breeding commencing 15 days prior to diagnostic testing as described in points III.1.5. and III.1.6. throughout the time the donor stallion(s) is/are in the SCC, and during semen collection for export to the United States;</p>																										
	<p>III.1.3. has/have been isolated under the supervision of the veterinarian in charge of the SCC from equidae not certified and tested to the same standards described in points III.1.5. and III.1.6. or under any restrictions which would make them ineligible as semen donors for export to the United States;</p>																										
	<p>III.1.4. was/were inspected on the day of semen collection and was/were found free of evidence of contagious and infectious diseases;</p>																										
	<p>III.1.5. was/were tested for dourine in a laboratory approved by the competent veterinary authority of the EU Member State with negative results using a complement fixation test at a dilution of 1:5 on samples taken within 30 days after entry into isolation and at 180 day intervals if the donor(s) remain(s) isolated under the supervision of the veterinarian in charge of the SCC;</p>																										
	<table style="width: 100%; border-collapse: collapse;"> <thead> <tr> <th style="width: 15%;">Dates of sampling</th> <th style="width: 15%;">Laboratory testing method</th> <th style="width: 15%;">Results</th> <th style="width: 55%;">Name and address of the approved laboratory</th> </tr> </thead> <tbody> <tr> <td>_____</td> <td>_____</td> <td>_____</td> <td>_____</td> </tr> <tr> <td>_____</td> <td>_____</td> <td>_____</td> <td>_____</td> </tr> <tr> <td>_____</td> <td>_____</td> <td>_____</td> <td>_____</td> </tr> <tr> <td>_____</td> <td>_____</td> <td>_____</td> <td>_____</td> </tr> <tr> <td>_____</td> <td>_____</td> <td>_____</td> <td>_____</td> </tr> </tbody> </table>	Dates of sampling	Laboratory testing method	Results	Name and address of the approved laboratory	_____	_____	_____	_____	_____	_____	_____	_____	_____	_____	_____	_____	_____	_____	_____	_____	_____	_____	_____	_____		
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Part II: Certification	II. Health information																										
	III.1.6. prior to semen collection and prior to the release of the donor stallion(s) and semen from isolation, was/were scrubbed and cultured negative in laboratories approved by the competent authority to culture for contagious equine metritis (CEM) under the supervision of the veterinarian in charge of the SCC, utilising the following procedure:																										
	III.1.6.1. for 5 consecutive days, the prepuce, penis, fossa glandis and urethral sinus of the donor stallion(s) must be aseptically cleaned and washed (scrubbed) while in full erection, with a solution of not less than 2.0% chlorhexidine. The entire penile area must be then thoroughly coated (packed) with an antibiotic ointment containing nitrofurazone, silver sulfadiazine or other agent recognized by the competent authority of the EU Member State as effective against the CEM agent;																										
	III.1.6.2. beginning at least 7 days after the last consecutive day of scrubbing and packing, 3 separate sets of 4 specimens each must be collected at intervals of not less than 72 hours between collection of each set. The collections must be made from the surface of the fossa glandis, the area of the urethral process and into the urethral fossa, the distal urethra, and the penile sheath, respectively;																										
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	III.1.6.3 Semen collection may begin after the first set of specimens is cultured negative. Note: a minimum of 7 days incubation time is required before samples can be considered negative.																										
	IV. The teaser:																										
	(1)either <input type="checkbox"/> [was not used.]																										
	(1)or <input type="checkbox"/> [was a phantom teaser mare.]																										
	(1)or <input type="checkbox"/> [was a live teaser mare provided that																										
	(1)either <input type="checkbox"/> [she has never been used for natural or artificial breeding prior to entrance into the SCC.]																										
	(1)or <input type="checkbox"/> [she has not been bred for the last 60 days and was tested negative for dourine and CEM as described in III.1.5 and III.1.6. For CEM, one specimen each must be taken with separate swabs from the clitoral sinuses and clitoral fossa. If any teaser mare tests positive for CEM, the mare must be treated for CEM in a manner approved by the competent authority of the EU Member State. The mare must be retested for CEM with negative results at least 21 days after treatment is completed.]]																										
	<table border="0" style="width: 100%;"> <tr> <td style="width: 15%;">Date and place</td> <td style="width: 15%;">Name and qualification of the Center Veterinarian</td> <td style="width: 70%;">Signature and stamp of the Center Veterinarian</td> </tr> <tr> <td>_____</td> <td>_____</td> <td>_____</td> </tr> <tr> <td>_____</td> <td>_____</td> <td>_____</td> </tr> <tr> <td>_____</td> <td>_____</td> <td>_____</td> </tr> <tr> <td>_____</td> <td>_____</td> <td>_____</td> </tr> <tr> <td>_____</td> <td>_____</td> <td>_____</td> </tr> </table>	Date and place	Name and qualification of the Center Veterinarian	Signature and stamp of the Center Veterinarian	_____	_____	_____	_____	_____	_____	_____	_____	_____	_____	_____	_____	_____	_____	_____								
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Part II: Certification	II. Health information		
Section B (to be signed at the end of the certificate by the Official Veterinarian);			
V.1. The semen collection center was approved by the competent authority of the Member State where the semen was collected;			
V.2. The Center Veterinarian that completed Section A of this certificate is authorized by the National Veterinary Service to perform this service.			
V.3. The donor animals for the semen to be exported to the United States have been part of the national flock/herd of the Member State where the semen was collected and are free from any movement or quarantine restrictions, according to point III.1.1 above;			
V.4. Health tests required for export to the United States of equine semen were performed by testing methods recognized by the Office International des Epizooties (OIE) Manual of Diagnostic Tests and Vaccines for Terrestrial Animals, as acceptable for international trade;			
V.5. The laboratory tests mentioned in _____ to _____ were carried out with negative results in a laboratory approved by the competent veterinary services;			
V.6. The EU Member State, _____, is free of African horse sickness.			
V.7. The semen to be exported to the United States was maintained under lock and key or in the custody of the SCC veterinarian, and segregated from other semen of lesser health status, until it was placed in the shipping container and sealed with official seals of the exporting Member State;			
V.8. None of the semen for export to the USA has been stored or transported in containers with semen produced under less than equivalent animal health conditions;			
V.9. The entire shipment exported under this certificate (including semen that might have been collected in more than one approved semen collection center by the same SCC veterinarian) has been maintained under continuous oversight of the Official Veterinarian until the conveyance is scheduled to depart for the United States.			
V.10. The shipping containers were sealed with an official seal of the exporting Member State, and the seal number is recorded on the health certificate;			
V.11. The semen is routed directly to the United States from the Member State in which it was collected with no stops en route other than those provided on the USDA import permit.			
Notes			
Part I:			
Box I.12.: Place of destination : this information is not compulsory.			
Box I.25.: 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 centre : shall correspond to the approval number of the semen collection centre of origin of the semen			
Quantity : shall correspond to the number of straws.			
Part II:			
(1) Delete as appropriate.			
· The signature and the stamp must be in a different colour to that of the printing.			
Certifying Officer			
Name (in capital letters)		Qualification and title	
Date of signature		Signature	
Stamp		Signature	