

Part I: Description of consignment	I.1. Consignor		I.2. IMSOC reference		I.2.a. Local reference	
	Name				I.3. Central Competent Authority	
	Address				I.4. Local Competent Authority	
	Country		ISO Code			
	I.5. Consignee			I.6. Operator conducting assembly operations independently of an establishment		
	Name			Name		
	Address			Address		
	Country			Country		
	Approval Number			Approval Number		
	ISO Code			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			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. Transporter			
Mode	International transport document	Identification	Name			
			Address			
			Approval Number			
			Country			
			ISO Code			
			I.17. Accompanying documents			
			Commercial document reference		Date of issue	
			Country		Place of issue	
I.18. Transport conditions						
Chilled <input type="checkbox"/>		Ambient <input type="checkbox"/>		Frozen <input type="checkbox"/>		
I.19. Container No / Seal No						
I.20. Certified as						
Quarantine or similar establishment <input type="checkbox"/>		Travelling circus/animal act <input type="checkbox"/>		Slaughter <input type="checkbox"/>		
Event or activity near borders <input type="checkbox"/>		Release into the wild <input type="checkbox"/>		Further keeping <input type="checkbox"/>		
Confined establishment <input type="checkbox"/>		Exhibition <input type="checkbox"/>		Other <input type="checkbox"/>		
I.21. For transit through a third country <input type="checkbox"/>						
Third country		ISO Code				
Exit point		BCP code				
Entry point		BCP code				
I.22. For transit through Member State(s) <input type="checkbox"/>			I.23. For export <input type="checkbox"/>			
Member State			Third country			
ISO Code			ISO Code			
			Exit point			
			BCP code			
I.24. Estimated journey time			I.25. Journey Log			
I.27. Total quantity			I.28. Total gross weight			
I.30. Description of consignment						
Commodity	Species	Subcategory	Sex	Identification system		
Identification Number		Age	Quantity			

Part II: Certification	II. Health information		
	<p>I, the undersigned official veterinarian, hereby certify that:</p> <p>II.1. The porcine animals(1) of the consignment described in Part I meet the following requirements:</p> <p>II.1.1. They are identified as provided for in Article 52 or 54(2) of Commission Delegated Regulation (EU) 2019/2035.</p> <p>II.1.2. They, for at least the 30 day period prior to the departure of the consignment, or since birth, if they are younger than 30 days of age,</p> <p>II.1.2.1. have been continuously resident in the establishment of origin;</p> <p>II.1.2.2. have not been in contact with kept porcine animals of a lower health status or subject to movement restrictions for animal health reasons;</p> <p>II.1.2.3. have not been in direct or indirect contact with kept animals that have entered the Union from a third country or territory during the 30 day period prior to the departure of the animals.</p> <p>II.1.3. They have not shown clinical signs or symptoms of diseases listed for porcine animals during the clinical examination which was carried out, within the 24 hour period prior to departure of consignment, on _____ (insert date dd/mm/yyyy).</p> <p>(2) <input type="checkbox"/> II.1.4. They come from one or more holdings officially recognised as applying controlled housing conditions in accordance with Article 8 of Commission Implementing Regulation (EU) 2015/1375 and have not passed through an establishment approved for assembly operations in accordance with Article 99(3) of Regulation (EU) 2016/429 that does not meet the requirements set out in Chapter I(A)(j) of Annex IV of Implementing Regulation (EU) 2015/1375.]</p> <p>II.2. According to official information, the animals described in Part I meet the following health requirements:</p> <p>II.2.1. They do not come from establishments subject to movement restrictions affecting the species or situated in a restricted zone established for reasons of diseases listed for porcine animals.</p> <p>II.2.2. They come from establishments in which infection with rabies virus in kept terrestrial animals has not been reported during the 30 day period prior to departure.</p> <p>II.2.3. They come from establishments in which anthrax in ungulates has not been reported during the 15 day period prior to departure.</p> <p>II.2.4. They come from establishments in which infection with Brucella abortus, B. melitensis and B. suis in porcine animals has not been reported during the last 42 days prior to departure, and in which during at least the 12 month period prior to departure</p> <p>(2) either <input type="checkbox"/> biosecurity and risk mitigating measures set out in Article 19(1)(f)(i) of [II.2.4.1. Commission Delegated Regulation (EU) 2020/688 have been introduced;]</p> <p>(2) and/or <input type="checkbox"/> surveillance for infection with Brucella abortus, B. melitensis and B. suis has [II.2.4.2. been carried out on the porcine animals kept on the establishments in accordance with Article 19(1)(f)(ii) of Delegated Regulation (EU) 2020/688.]</p> <p>II.2.5. They come from establishments in which infection with Aujeszky's disease virus has not been reported during the 30 day period prior to departure of the consignment.</p> <p>(2) <input type="checkbox"/> II.2.6. They are moved to a Member State or zone thereof with the status free from infection with Aujeszky's disease virus and have not been vaccinated against infection with Aujeszky's disease virus, and</p> <p>(2) either <input type="checkbox"/> come from establishments free from infection with Aujeszky's disease virus, and [II.2.6.1.</p> <p>(2) either <input type="checkbox"/> the establishments of origin are situated in a Member State or zone [II.2.6.1.1. with the status free from infection with Aujeszky's disease virus;]]</p> <p>(2) and/or <input type="checkbox"/> the animals in the consignment have been subjected to a serological test for the detection of antibodies against whole Aujeszky's disease virus with one of the diagnostic methods provided for in Part 7 of Annex I to Delegated Regulation (EU) 2020/688(3)(4), with a negative result, on a sample taken during the 15 day period prior to departure;]]</p>		

Part II: Certification	II. Health information			
	(2)	and/or <input type="checkbox"/> [II.2.6.2.]	<p>come from establishments not free from infection with Aujeszky's disease virus, and</p> <ul style="list-style-type: none"> - have been kept in an approved quarantine establishment for a period of at least 30 days; and - have been subjected to a serological test for the detection of antibodies against whole Aujeszky's disease virus with the diagnostic method provided for in Part 7 of Annex I to Delegated Regulation (EU) 2020/688, with a negative result, carried out on samples taken on two occasions at an interval of not less than 30 days, the last sample taken during the 15 day period prior to departure.]] 	
	(2)	○ [II.2.6.]	They are moved to a Member State or zone thereof with an approved eradication programme for infection with Aujeszky's disease virus, and	
	(2)	either <input type="checkbox"/> [II.2.6.1.]	come from establishments free from infection with Aujeszky's disease virus, and	
	(2)		either <input type="checkbox"/> [II.2.6.1.1.]	the establishments of origin are situated in a Member State or zone thereof with the status free from infection with Aujeszky's disease virus;]]
	(2)		and/or <input type="checkbox"/> [II.2.6.1.2.]	the establishments of origin are situated in a Member State or zone thereof with an approved eradication programme for infection with Aujeszky's disease virus;]]
	(2)		and/or <input type="checkbox"/> [II.2.6.1.3.]	the animals in the consignment have been subjected to a serological test for the detection of antibodies against whole Aujeszky's disease virus or antibodies against Aujeszky's disease virus-gE protein, where applicable, with one of the diagnostic methods provided for in Part 7 of Annex I to Delegated Regulation (EU) 2020/688(4), with a negative result, on a sample taken during the 15 day period prior to departure;]]
	(2)	and/or <input type="checkbox"/> [II.2.6.2.]	<p>come from an establishment not free from infection with Aujeszky's disease virus, and</p> <ul style="list-style-type: none"> - have been kept in an approved quarantine establishment for a period of at least 30 days; and - have been subjected to a serological test for the detection of antibodies against whole Aujeszky's disease virus with the diagnostic method provided for in Part 7 of Annex I to Delegated Regulation (EU) 2020/688, with a negative result, carried out on samples taken on two occasions at an interval of not less than 30 days, the last sample taken during the last 15 days prior to departure.]] 	
	II.3.	To the best of my knowledge and as declared by the operator, the animals come from establishments where there were no abnormal mortalities with an undetermined cause.		
	(2) <input type="checkbox"/> [II.4.]	<p>According to official information and as declared by the operator, they are semen donor animals, and</p> <p>II.4.1. they come from a semen collection centre and will be transported directly to another semen collection centre in accordance with Article 19 of Commission Delegated Regulation (EU) 2020/686; and</p> <p>(2) either ○ [II.4.2.] they were continuously resident since the date of their admission at the semen collection centre and were subjected, with negative results, to all compulsory routine tests referred to in point 2 of Chapter I of Part 2 of Annex II to Delegated Regulation (EU) 2020/686 in the period of the preceding 12 months prior to date of that movement; and]</p> <p>(2) or ○ [II.4.2.] they were subjected, with negative results, to all tests referred to in point 1(b) and (c) of Chapter I of Part 2 of Annex II to Delegated Regulation (EU) 2020/686, required before admission to a semen collection centre carried out during the period immediately preceding quarantine and during the quarantine period; and]</p> <p>II.4.3. the prior consent of the centre veterinarian of the semen collection centre of destination has been obtained by the operator; and</p> <p>II.4.4. the means of transport used have been cleansed and disinfected before use.]</p>		

Part II: Certification	II. Health information		
	<p>II.5. Arrangements are made to transport the consignment in accordance with Article 4 of Delegated Regulation (EU) 2020/688.</p> <p>II.6. This certificate is valid for 10 days from the date of issuing. In the case of transport by waterway/sea of animals, the period of validity of the certificate may be extended by the duration of the journey by waterway/sea.</p> <p>(2)(5) <input type="checkbox"/> II.7. Since leaving their establishments of origin and before arriving to this establishment approved for assembly operations, none of the animals of the consignment has undergone more than two assembly operations, and</p> <p>(2) either <input type="checkbox"/> [they come from their establishments of origin.]</p> <p>(2) or <input type="checkbox"/> [at least one of the animals of the consignment has undergone one assembly operation on an approved establishment.]</p> <p>(2) or <input type="checkbox"/> [at least one of the animals of the consignment has undergone two assembly operations on approved establishments.]</p> <p>(1) <input type="checkbox"/> II.8. The animals meet the additional guarantees for:</p> <p>(1) either <input type="checkbox"/> II.8.1. Porcine animals kept in restricted zone I in compliance with the special control measures relating to African swine fever laid down in Commission Implementing Regulation (EU) 2021/605(*)]</p> <p>(1) or <input type="checkbox"/> II.8.2. Porcine animals kept in restricted zone II in compliance with the special control measures relating to African swine fever laid down in Commission Implementing Regulation (EU) 2021/605(*)]</p> <p>(1) or <input type="checkbox"/> II.8.3. Porcine animals kept in restricted zone III in compliance with the special control measures relating to African swine fever laid down in Commission Implementing Regulation (EU) 2021/605(*)]</p>		
<p>Animal welfare attestation</p> <p>At the time of inspection, the animals covered by this health certificate were fit to be transported in accordance with the provisions of Council Regulation (EC) No 1/2005 on the intended journey due to start on _____ (insert date) (6)(7).</p>			

Part II: Certification	II. Health information							
	<p>Notes:</p> <p>In accordance with the Agreement on the withdrawal of the United Kingdom of Great Britain and Northern Ireland from the European Union and the European Atomic Energy Community, and in particular Article 5(4) of the Protocol on Ireland / Northern Ireland in conjunction with Annex 2 to that Protocol, references to European Union in this certificate include the United Kingdom in respect of Northern Ireland.</p> <p>This animal health certificate shall be completed according to the notes for the completion of certificates provided for in Chapter 2 of Annex I to Commission Implementing Regulation (EU) 2020/2235.</p> <p>Part I:</p> <p>Box reference I.11: “Place of dispatch”: Indicate an establishment of the origin of the animals in the consignment or an establishment approved for assembly operations in accordance with Articles 97 and 99 of Regulation (EU) 2016/429.</p> <p>Box reference I.12: “Place of destination”: Indicate an establishment of the final destination of the consignment or an establishment approved for assembly operations in accordance with Articles 97 and 99 of Regulation (EU) 2016/429.</p> <p>Box reference I.17: “Accompanying documents”: In case the animals are dispatched from an establishment approved for assembly operations in the Member State of origin, the reference number(s) of the official document(s), based on which the animal health certificate for this consignment is issued in this establishment approved for assembly operations, may be indicated.</p> <p style="padding-left: 40px;">In case the animals are dispatched from an establishment approved for assembly operations in the Member State of passage, the reference number(s) of the certificate(s), based on which the animal health certificate for this consignment is issued in this establishment approved for assembly operations, must be indicated.</p> <p>Box reference I.30: “Identification number”: Indicate identification codes of the animals in the consignment identified in accordance with Article 52 or 54(2) of Delegated Regulation (EU) 2019/2035.</p> <p>Part II:</p> <p>(1) There can be one or more animals in the consignment.</p> <p>(2) Delete if not applicable.</p> <p>(3) For porcine animals less than four months old born to dams vaccinated with a gE-deleted vaccine, the diagnostic method for the detection of antibodies against Aujeszky’s disease virus gE protein provided for in Part 7 of Annex I to Delegated Regulation (EU) 2020/688 may be used.</p> <p>(4) The number of porcine animals tested must allow at least for the detection of 10% seroprevalence of the consignment with 95% confidence.</p> <p>(5) Applicable in case the consignment is dispatched from the establishment approved for assembly operations.</p> <p>(6) In the case where a consignment is grouped in an establishment approved for assembly operations and comprises animals that were loaded on different dates, the date which the journey commenced for the whole consignment is considered to be the earliest date when any part of the consignment left the establishment of origin.</p> <p>(7) This statement does not exempt transporters from their obligation in accordance with Union rules in force in particular regarding the fitness to be transported.</p>							
<p>Certifying Officer/Official veterinarian</p> <table border="0" style="width: 100%;"> <tr> <td style="width: 50%;">Name (in capital letters)</td> <td style="width: 50%;">Qualification and title</td> </tr> <tr> <td>Date of signature</td> <td>Signature</td> </tr> <tr> <td>Stamp</td> <td></td> </tr> </table>			Name (in capital letters)	Qualification and title	Date of signature	Signature	Stamp	
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