EUROPEAN UNION

INTRA

Part I: Description of consignment

I.1. Consignor
Name
Address
Country ISO Code

I.2. IMSOC reference

I.2.a. Local reference

I.3. Central Competent Authority

I.4. Local Competent Authority

I.5. Consignee
Name
Address
Country ISO Code

I.6. Operator conducting assembly operations independently of an establishment
Name
Address
Approval Number
Country ISO Code

I.7. Country of origin ISO Code

I.8. Region of origin Code

I.9. Country of destination ISO Code

I.10. Region of destination Code

I.11. Place of dispatch
Name
Address
Approval Number
Country ISO Code

I.12. Place of destination
Name
Address
Approval Number
Country ISO Code

I.13. Place of loading
Name
Address
Approval Number
Country ISO Code

I.14. Date and time of departure

I.15. Means of Transport

Mode	International transport document	Identification

I.16. Transporter
Name
Address
Approval Number
Country ISO Code

I.17. Accompanying documents
Document Type
Accompanying document reference
Date of Issue
Country
Place of issue

I.18. Transport conditions
Ambient ☐ Chilled ☐ Frozen ☐

I.19. Container Number / Seal Number

I.20. Certified as
Other ☐ Further keeping ☐ Exhibition ☐ Confined establishment ☐
Release into the wild ☐ Travelling circus/animal act ☐ Breeding ☐ Quarantine or similar establishment ☐
Event or activity near borders ☐

I.21. For transit through a third country ☐
Third country ISO Code
Exit point BCP code
Entry point BCP code

I.22. For transit through Member State(s) ☐
Member State ISO Code

I.23. For export ☐
Third country 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
1. 01 LIVE ANIMALS

Part I: Description of consignment

0103 Live swine

#1.	Commodity	Subcategory	Sex	Identification system
Species		Identification Number	Age	Quantity

Part II: Certification

II. Health information

I, the undersigned official veterinarian, hereby certify that:

II.1. The porcine animals (1) of the consignment described in Part I meet the following requirements:

II.1.1. They are identified as provided for in Article 52 or 54(2) of Commission Delegated Regulation (EU) 2019/2035.

II.1.2. They, for at least 30 days prior to the date of departure of the consignment, or since birth, if they are younger than 30 days of age,

II.1.2.1. have been continuously resident in the establishment of origin;

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;

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 last 30 days prior to the date of departure of the consignment.

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 last 24 hours prior to the time of departure of the consignment, on (insert date dd/mm/yyyy).

(2) ☐ [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 of the European Parliament and of the Council that does not meet the requirements set out in Chapter I(A), point (j), of Annex IV to Implementing Regulation (EU) 2015/1375.]

II.2. According to official information, the animals described in Part I meet the following health requirements:

II.2.1. ○ (2) either [They come from establishments or zones not subject to movement restrictions affecting porcine animals and established for reasons of diseases listed for those species or diseases subject to emergency measures relevant for those species, and they have not been in contact with kept animals of a listed species of a lower health status for an adequate period.]

(2) ○ or [They come from establishments or zones subject to movement restrictions affecting porcine animals and established for (3), but derogations from movement restrictions have been granted, and:

(2) ☐ [they comply with the requirements set out in (4);]]

(2) ☐ [and in particular, they are (5).]]

II.2.2. They come from establishments in which infection with rabies virus in kept terrestrial animals has not been reported during the last 30 days prior to the date of departure of the consignment.

II.2.3. They come from establishments in which anthrax in ungulates has not been reported during the last 15 days prior to the date of departure of the consignment.

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 the date of departure of the consignment, and in which during at least 12 months prior to the date of departure of the consignment:

(2) ☐ either [II.2.4.1. biosecurity and risk mitigating measures set out in Article 19(1), point (f)(i), of Commission Delegated Regulation (EU) 2020/688 have been introduced;]

(2) ☐ and/or [II.2.4.2. surveillance for infection with Brucella abortus, B. melitensis and B. suis has been carried out on the porcine animals kept in the establishments in accordance with Article 19(1), point (f)(ii), of Delegated Regulation (EU) 2020/688.]

II.2.5. They come from establishments in which infection with Aujeszky's disease virus has not been reported during the last 30 days prior to the date of departure of the consignment.

(2) ☐ [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:

(2) ☐ either [II.2.6.1. come from establishments free from infection with Aujeszky's disease virus, and:

(2) ☐ [II.2.6.1.1. the establishments of origin are situated in a Member State or zone thereof with the status

Part II: Certification

II. Health information				
either		free from infection with Aujeszky's disease virus;]]		
(2) ☐ and/or	[II.2.6.1.2.	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 (6) (7), with a negative result, on a sample taken during the last 15 days prior to the date of departure of the consignment;]]]		
(2) ☐ and/or	[II.2.6.2.	come from establishments not free from infection with Aujeszky's disease virus, and:		
–		have been kept in an approved quarantine establishment for at least 30 days prior to the date of departure of the consignment; 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 the date of departure of the consignment.]]		
(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	[II.2.6.1.	come from establishments free from infection with Aujeszky's disease virus, and		
(2) ☐ either	[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	[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	[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 (7), with a negative result, on a sample taken during the last 15 days prior to the date of departure of the consignment;]]]		
(2) ☐ and/or	[II.2.6.2.	come from an establishment not free from infection with Aujeszky's disease virus, and:		
–		have been kept in an approved quarantine establishment for at least 30 days prior to the date of departure of the consignment; 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 the date of departure of the consignment.]]		
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.		
II.4.		Arrangements are made to transport the consignment in accordance with Article 4 of Delegated Regulation (EU) 2020/688.		
II.5.		This animal health 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.		
(2) (8)	☐ [II.6.	Since the date of departure from their establishments of origin and prior to the date of arrival to this establishment approved for assembly operations, none of the animals of the consignment has undergone more than two assembly operations, and:		
(2) ○ either	[they come from their establishments of origin.]]			
(2) ○ or	[at least one of the animals of the consignment has undergone one assembly operation in an approved establishment.]]			
(2) ○ or	[at least one of the animals of the consignment has undergone two assembly operations in the approved			

Part II: Certification

II. Health information		

establishments.]]

Animal welfare attestation

At the time of inspection, the animals covered by this animal 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) (9) (10).

Notes:

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 the Union in this animal health certificate include the United Kingdom in respect of Northern Ireland.

This animal health certificate shall be completed in accordance with the notes for the completion of certificates provided for in Chapter 2 of Annex I to Commission Implementing Regulation (EU) 2020/2235.

Part I:

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.
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.
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. 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, shall be indicated.
Box reference I.30:	“Identification number”: Indicate identification codes of the animals in the consignment identified in accordance with Article 52 or Article 54(2) of Delegated Regulation (EU) 2019/2035.

Part II:

(1)	There may be one or more animals in the consignment.
(2)	Delete if not applicable.
(3)	Insert the name of the disease(s).
(4)	Insert the specific reference to the article(s), title, and number of the relevant legal act(s) adopted by the Commission providing for those requirements.
(5)	Insert the specific attestation(s) provided for in and required by the relevant legal act(s) adopted by the Commission, as referred to in Article 126(1), points (b)(ii) and (iii), of Regulation (EU) 2016/429.
(6)	For porcine animals less than 4 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.
(7)	The number of porcine animals tested shall allow at least for the detection of 10 % seroprevalence of the consignment with 95 % confidence.
(8)	Applicable in case the consignment is dispatched from the establishment approved for assembly operations.
(9)	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.
(10)	This statement does not exempt transporters from their obligation in accordance with Union rules in force in particular regarding the fitness to be transported.

Part II: Certification

II. Health information		
Certifying Officer/Official veterinarian		
Name (in capital letters)	Qualification and title	
Date	Signature	
Stamp		