# MODEL ANIMAL HEALTH CERTIFICATE FOR THE MOVEMENT BETWEEN MEMBER STATES OF PORCINE ANIMALS NOT INTENDED FOR SLAUGHTER (MODEL 'POR-INTRA-X')

EURC	PEAN UN	ION				INTRA		
	I.1	Consignor			IMSOC reference			
		Name		I.2a	Local reference			
Part I: Description of consignment		Address		I.3	Central Competent Authority	QR CODE		
		Country	ISO country code	I.4	<b>Local Competent Authority</b>			
	1.5	Consignee		I.6	6 Operator conducting assembly operations independent establishment			
		Name			Name	Registration No		
		Address			Address			
ption		Country	ISO country code		Country	ISO country code		
scri	I.7	Country of origin	ISO country code	1.9	Country of destination	ISO country code		
Ŏ.	1.8	Region of origin	Code	I.10	Region of destination	Code		
Part I:	I.11	Place of dispatch		I.12	Place of destination			
		Name	Registration/Approval No		Name	Registration/Approval No		
		Address			Address			
		Country	ISO country code		Country	ISO country code		
	I.13	Place of loading			Date and time of departure			
	I.15	Means of transport		I.16	Transporter			
		□ Vessel	□ Aircraft		Name	Registration/Authorisation No		
					Address			
		□ Railway	□ Road vehicle		Country	ISO country code		
				I.17	Accompanying documents			
		Identification	□ Other		Type	Code		
		Document			Country Commercial document reference	ISO country code		
	I.18	Transport condition	s   Ambient	'		Frozen		
	I.19	Container number/						
		Container No	S	Seal No				

I.20	Certified as or	for							
□ Further keeping		□ Slaughter	□ Slaughter		□ Confined establishment		☐ Germinal products		
□ Regi	stered equine animal	□ Travelling circus/anim	al act	□ Exhibition		□ Event or activity near borders		ear borders	
□ Relea	ase into the wild	□ Dispatch centre		□ Relaying area/purification centre		☐ Ornamental aquaculture		ulture	
□ Furth	ner processing	□ Organic fertilizers and	soil	□ Technical use		□ Quarantii	ne or simi	ilar	
		improvers	improvers				establishment		
□ Prod	ucts for human	□ Pollination		□ Live aquati	c animals for	□ Other			
consun	nption			human consu	mption				
I.21	□ For transit	through a third country							
	Third country			ISO cou	ntry code				
	Exit point			BCP coo	le				
	Entry point			BCP code					
I.22	□ For transit throu	gh Member State(s)		I.23 □ For	export				
	Member State ISO country code			Third country ISO country code			y code		
Member State		ISO country	ISO country code		Exit point		BCP code		
	Member State	ISO country	code						
I.24	Estimated journey	time		I.25 Jou	urney log	□ yes		□ no	
I.26	Total number of pa	ckages		I.27 To	tal quantity				
I.28	Total net weight/gr	oss weight (kg)		I.29 Total space foreseen for the consignment					
I.30	Description of cons	ignment							
CN coo	de Species	Subspecies/Category Sex	Ident syste	tification	Identification	number	Age	Quantity	
			sysic	Ш				Type	
Region of origin		Cold store	Identification mark		Type of packa	ging		Net weight	
Slaughterhouse		Treatment type		re of modity	Number of pac	ckages		Batch No	
		Date of collection/production	Man plant	ufacturing	Approval or re number of plant/establish		Test		

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Leno	PEAN UN					Certificat	te model POR-INTRA-A		
	II. Heal	th information	l		II.a Certificate reference	II.b	IMSOC reference		
	I, the ι	ındersigned	official veterinarian, hereby certify that:						
	II.1.	The porcir	ne animals(1)	of the consignment descri	bed in Part I meet the follow	ing requi	irements:		
		II.1.1.	Regulation	n (EU) 2019/2035.	for in Article 52 or 54(2)				
		II.1.2.	they are yo	ounger than 30 days of age					
			II.1.2.1.	have been continuously	resident in the establishment	of origin	n;		
			II.1.2.2.		ct with kept porcine animals of a lower health status or strictions for animal health reasons;				
			II.1.2.3.		or indirect contact with kept untry or territory during the				
ū		II.1.3.	the clinica		or symptoms of diseases listed carried out, within the 24 housert date dd/mm/yyyy).				
Part II: Certification		<sup>(2)</sup> [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.]						
	II.2.	According	g to official i	nformation, the animals de	escribed in Part I meet the fo	llowing l	health requirements:		
		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.						
	II.2.2. They come from establishments in whas not been reported during the 30 II.2.3. They come from establishments in the 15 day period prior to departure.						ept terrestrial animals		
					which anthrax in ungulates	has not l	been reported during		
	II.2.4. They come from establishments in <i>B. suis</i> in porcine animals has not be in which during at least the 12 mont			en reported during the last 4					
		<sup>(2)</sup> either	[II.2.4.1.		mitigating measures set of Regulation (EU) 2020/688 h				
		<sup>(2)</sup> and/or	[II.2.4.2.	surveillance for infection been carried out on the	on with <i>Brucella abortus, E</i> porcine animals kept on the of Delegated Regulation (EV	3. <i>melite</i> establish	nsis and B. suis has aments in accordance		
		II.2.5.	•		which infection with Aujeszk ior to departure of the consig	•	se virus has not been		

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<sup>(2)</sup> [11.2.6.	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					
<sup>(2)</sup> either	[II.2.6.1.	come from esta	blishments free from infection with Aujeszky's disease virus, and			
	<sup>(2)</sup> either		establishments of origin are situated in a Member State or zone in the status free from infection with Aujeszky's disease virus;]]			
	<sup>(2)</sup> and/or	test viru Ann	animals in the consignment have been subjected to a serological for the detection of antibodies against whole Aujeszky's disease is with one of the diagnostic methods provided for in Part 7 of nex I to Delegated Regulation (EU) 2020/688 <sup>(3)(4)</sup> , with a negative alt, on a sample taken during the 15 day period prior to			

- departure;]]]

  (2) and/or [II.2.6.2. come from establishments not free from infection with Aujeszky's disease virus,
  - have been kept in an approved quarantine establishment for a period of at least 30 days; and

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- 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
- [II.2.6.1. come from establishments free from infection with Aujeszky's disease virus, and [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;]]
  - [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<sup>(4)</sup>, with a negative result, on a sample taken during the 15 day period prior to departure;

EUROPEAN UNION Certificate model POR-INTRA-X (2) and/or [II.2.6.2. come from an establishment not free from infection with Aujeszky's disease 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.]] 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. According to official information and as declared by the operator, they are semen donor animals, and <sup>(2)</sup>[II.4. 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 (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] they were subjected, with negative results, to all tests referred to in point 1(b) and (c) of (2) or [II.4.2. 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] the prior consent of the centre veterinarian of the semen collection centre of destination has II.4.3. been obtained by the operator; and II.4.4. the means of transport used have been cleansed and disinfected before use.] II.5. Arrangements are made to transport the consignment in accordance with Article 4 of Delegated Regulation (EU) 2020/688. 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. (2)(5)[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 <sup>(2)</sup>either [they come from their establishments of origin.]] (2)or

approved establishment.]]

approved establishments.]]

 $^{(2)}or$ 

[at least one of the animals of the consignment has undergone one assembly operation on an

[at least one of the animals of the consignment has undergone two assembly operations on

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#### Animal welfare attestation

#### 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 European Union in this certificate include the United Kingdom in respect of Northern Ireland.

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.

### 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, must be indicated.

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.

## Part II:

There can be one or more animals in the consignment.

(2) Delete if not applicable.

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.

The number of porcine animals tested must allow at least for the detection of 10% seroprevalence of the consignment with 95% confidence.

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Applicable in case the consignment is dispatched from the establishment approved for assembly operations.

- 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.
- This statement does not exempt transporters from their obligation in accordance with Union rules in force in particular regarding the fitness to be transported.

Official	veterina	ırian
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Name (in capital letters) Qualification and title

Local Control Unit name Local Control Unit code

Date

Stamp Signature