	UNITED KINGDOM				alth certificate to the EU
	I.1 Consignor/Exporter		I.2 Certificate refere	ence	I.2a
	Name				
	Address		I.3 Central Compete	· ·	
			DEPARTMENT FOR FOOD & RURAL AF		
			I.4 Local Competent	Authority	1 /
	Country	ISO country code	ANIMAL AND PLAN	NT HEALTH AGENCY	
	1.5 Consignee/Importer		I.6 Operator respons	sible for the consignment	
	Name		Name		
ta√	Address		Address		
Part I: Description of consignment	, 6				
ou	Country	ISO country code	Country		SO country code
ripti	I.7 Country of origin	ISO country code	I.9 Country of destin	nation	ISO country code
: Desc	I.8 Region of origin	Code	I.10 Region of destin	ation	Code
Part I	I.11 Place of dispatch	Registration/Approval No	I.12 Place of destinat	tion	Registration/Approval No
	Name		Name		
	Address		Address		
		/(
	Country	ISO country code	Country		SO country code
	I.13 Place of loading		I.14 Date and time of		
	I.15 Means of transport		I.16 Entry Border C	ontrol Post	
	☐ Aircraft ☐	Vessel	I.17 Accompanying of	documents	
				_ Cod	
	E D ''	Road vehicle	Туре		
	□ Railway □	Road venicie	Country	ISO	country code
			Commercial documen	nt reference	
	Identification				
	I.18 Transport conditions	☐ Ambient	☐ Chilled	☐ Froze	en
	I.19 Container number/Seal number				
	Container No		Seal No	4	
	I.20 Certified as or for				
	☐ Germinal products				
	I.21		I.22	nternal market	
	Third country	ISO country code	I.23		
	I.24 Total number of packages	I.25 Total quant	ity I.:	26	

TINITT	ED KINGDOM			II.a Certificat	e reference
I.27	Description of con	nsignment			
1	CN code	Species	Subspecies/Category	Identification number	Quantity
	Туре	Approval or registration number of plant/establishment/centre	Identification mark	Date of collection/production	
2	CN code	Species	Subspecies/Category	Identification number	Quantity
	Туре	Approval or registration number of plant/establishment/centre	Identification mark	Date of collection/production	
3	CN code	Species	Subspecies/Category	Identification number	Quantity
	Туре	Approval or registration number of plant/establishment/centre	Identification mark	Date of collection/production	
4	CN code	Species	Subspecies/Category	Identification number	Quantity
	Туре	Approval or registration number of plant/establishment/centre	Identification mark	Date of collection/production	
5	CN code	Species	Subspecies/Category	Identification number	Quantity
	Туре	Approval or registration number of plant/establishment/centre	Identification mark	Date of collection/production	

UNITED KINGDOM II. Health information

II.a Certificate reference

II. Health information							
I, the undersigned official veterinarian, hereby certify that:							
П.1.							
	II.1.1.	authorised for the entry into the Union of semen of porcine animals and listed in Annex XI to Commission Implementing Regulation (EU) 2021/404;					
(1) either	[11.1.2.	where foot and mouth disease was not reported for at least 24 months immediately prior to the date of collection of the semen and until the date of its dispatch;]					
(1)(2) or	[II.1.2.	where foot and mouth disease was not reported for a period starting on the date/ (dd/mm/yyyy) immediately prior to the date of collection of the semen and until the date of its dispatch;]					
(1) either	[II.1.3.	where classical swine fever was not reported for at least 12 months immediately prior to the date of collection of the semen and until the date of its dispatch;]					
(1)(2) or	[II.1.3.	where classical swine fever was not reported for a period starting on the date// (dd/mm/yyyy) immediately prior to the date of collection of the semen and until the date of its dispatch;]					
	II.1.4.	where infection with rinderpest virus and African swine fever were not reported for at least 12 months immediately prior to the date of collection of the semen and until the date of its dispatch;					
	II.1.5.	where no vaccination against infection with rinderpest virus and classical swine fever has been carried out for at least 12 months immediately prior to the date of collection of the semen and until the date of its dispatch, and no vaccinated animals entered into the third country or territory, or zone thereof during that period, and					
	⁽¹⁾ either	[no vaccination against foot and mouth disease has been carried out for the same period, and no vaccinated animals entered into the third country or territory, or zone thereof during that period.]					
	⁽¹⁾ or	[vaccination against foot and mouth disease has been carried out for the same period, or vaccinated animals entered into the third country or territory, or zone thereof during that period.]					
II.2.		escribed in Part I was obtained from donor animals which originated, prior to the date of commencement of the quarantine referred to 6, from establishments:					
	II.2.1.	situated in an area where foot and mouth disease has not been reported within a 10-km radius centred on the establishments for at least the preceding 30 days and in which foot and mouth disease has not been reported during at least the preceding 3 months,					
	(1) either	[in which they were not vaccinated against foot and mouth disease;]					
	⁽¹⁾ or	[in which they were vaccinated against foot and mouth disease during 12 months immediately prior to the date of collection of the semen but not during the last 30 days immediately prior to the date of collection of the semen, and in which 5% (with a minimum of five straws) of each quantity of semen taken from a donor animal at any time is submitted to a virus isolation test for foot and mouth disease with negative results;					
	II.2.2.	which are free from infection with <i>Brucella abortus</i> , <i>B. melitensis</i> and <i>B. suis</i> in accordance with the requirements laid down in Part 5, Chapter IV, of Annex II to Commission Delegated Regulation (EU) 2020/686;					
	II.2.3.	where no clinical, serological, virological or pathological evidence of infection with Aujeszky's disease virus had been detected during at least the preceding 12 months;					
	II.2.4.	where, during at least 3 months immediately prior to the date of entry into the quarantine accommodation, no animal was vaccinated against infection with porcine reproductive and respiratory syndrome virus and no infection with porcine reproductive and respiratory syndrome virus was detected.					
II.3.	The semen de	escribed in Part I has been collected, processed and stored, and dispatched from the semen collection centre (3) which:					
	II.3.1.	is approved and listed by the competent authority of the third country or territory;					
	II.3.2.	complies with requirements as regards responsibilities, operational procedures, facilities and equipment set out in Part 1 of Annex I to Delegated Regulation (EU) 2020/686.					
II.4.	The semen de	escribed in Part I was obtained from donor animals which:					
	II.4.1.	were not vaccinated against infection with rinderpest virus, classical swine fever and infection with poreine reproductive and respiratory syndrome virus;					
	II.4.2.	remained for at least 3 months immediately prior to the date of collection of the semen in a third country or territory or zone thereof referred to in box I.7;					
	II.4.3.	did not show symptoms or clinical signs of transmissible animal diseases on the day of their admission to a semen collection centre and on the day of collection of the semen;					
	II.4.4.	are individually identified as provided for in Article 21(1) of Delegated Regulation (EU) 2020/692;					
	II.4.5.	for at least 30 days immediately prior to the date of collection of the semen and during the collection period:					
		II.4.5.1. were kept in establishments not situated in a restricted zone established due to the occurrence of foot and mouth disease, infection with rinderpest virus, classical swine fever or African swine fever, or of an emerging disease relevant for porcine animals;					
	I, the und 11.1. (1) etther (1) (2) or (1) either (1) (2) or	I, the undersigned offici II.1. The semen d territory, or a II.1.1. (Petther II.1.2. (1) 2) or III.1.2. (1) either III.1.3. (1) (2) or III.1.3. II.1.4. II.1.5. (1) either (1) or II.2. The semen d in point II.4. II.2.1. (1) either (1) or II.2.2. II.2.3. II.2.4. II.3. The semen d II.3.1. II.3.2. II.4. The semen d II.4.1. II.4.2. II.4.3. II.4.4.					

II.a Certificate reference

H.4.5.2. were kept in a single establishment whose infection with <i>Proceeding abortum</i> . <i>B. meditions</i> with relabor viture, and an authority of the context with animals from establishments with an infection with procine reproductive programment of the context with animals from establishments which do not meet the conditions referred to in point II.4.5.1; or from establishments which do not meet the conditions referred to in point II.4.5.2; and the conditions referred to in point II.4.5.2; and the same health status were present, which on the day of their arthristion to the same nealth status were present, which on the day of their arthristion to the same nealth status were present, which on the day of their arthristion to the same nealth status were present, which on the day of their arthristion to the same nealth status were present, which on the day of their arthristion to the same nealth status were present, which on the day of their arthristion to the same nealth status and present and the same health status were present, which on the day of their arthristion to the same nealth status and the referred to in point II.4.5.1; it was fareful an an area where for an adm and this excels has to been reported within a 10-km radius control on the quantities and common of the diseases referred to in point II.4.5.1 has been exported to the point and the same and to days from the date of the collection of the sense and the same and to days from	UNITED KI	NGDOM		
### referred to in point ### 14.5.1 or from establishments which do not meet the conditions referred to in point ### 14.5.2; ### 14.6.2 have been subjected for a quanardine for at least 28 days in quanardine accommodation, where only other cloven-hoofed animals with at least the sums health tains were present, which on the day of their admission to the seemes collection centre compiled with the following conditions: #### 14.6.1 in was not situated in a restricted zone established due to diseases referred to in point ### 14.5.1; #### 14.6.2 in more of the diseases referred to in point ### 14.5.2 has been reported for at least the preceding 30 days; #### 14.6.3 it was situated in an area where foot and mouth disease has not been reported within a 10-km radius centred on the quanardine accommodation for at least the preceding 30 days; ##### 14.6.2 in was free from infection with *### 37.6 has been reported for at least the preceding the date of admission of the animals to the seeme collection centre: #### 14.6.2 in was free from infection with *### 37.6 has been reported mentions and *### 37.6 has been admission of the animals to the seeme collection centre: #### 14.7.2 were gent in segion collection centres: #### 14.7.3 were gent in segion collection centres: #### 14.7.4 were from infection with *### 37.6 has been reported for at least 30 days immediately prior to the date of objection of the senior of objection of the consignment of senior to the Union.] #### 14.7.2 where more of the diseases referred to in point ### 3.5 has been reported for at least 30 days immediately prior to the date of collection of the senior objection of the consignment of senior to the Union.] #### 14.7.2 where more of the diseases referred to in point ### 3.5 has been reported within a 10-km radius centred on the seman collection entry for the consignment of senior to the Union. #### 14.7.2 where more of the diseases referred to in point ### 3.5 has been reported for at least 30 days immediately prior to the date of collect			II.4.5.2.	rabies virus, anthrax, infection with Aujeszky's disease virus and infection with porcine reproductive and
14.6. have been subjected to a quarantine for at least 28 days in quarantine accommodation, where only other cloves—booked animals with at least the same health status were present, which on the day of their admission to the semen collection center complied with the following conditions: 14.6.1 it was not situated in a restricted zone established due to diseases referred to in point II.4.5.1; 14.6.2 none of the diseases referred to in point II.4.5.2 has been reported for at least the preceding 30 days; 14.6.3 it was situated in an area where foot and mouth disease has not been reported within a 10-km radius centred on the quarantine accommodation for at least the preceding 30 days; 14.6.3 has had no outbreak of foot and mouth disease reported during at least 3 months immediately preceding the date of admission of the animatis to the semen collection centre; 14.6.5 it was free from infection with Brucella abortus, Brucella netitensis and Brucella suits for at least the preceding 3 nonths; 14.7.1 were step in a summer collection centres: 14.4.2 spiffer were not situated in a restricted zone established due to diseases referred to in point II.4.5.1; 14.7.2 viding none of the diseases referred to in point II.4.5.2 has been reported for at least 30 days immediately prior to the dust of of situation of the semen, and			II.4.5.3.	
at least the same health status were present, which on the day of their admission to the semen collection centre complied with the following conditions: II.4.6.1. it was not situated in a restricted zone established due to diseases referred to in point II.4.5.1; II.4.6.2. none of the diseases referred to in point III.4.5.2 has been reported for at least the preceding 30 days; II.4.6.3. has had no outbrack of foot and month disease reported during at least 3 months immediately preceding the date of admission of the animals to the semen collection centre; II.4.6.3. has had no outbrack of foot and month disease reported during at least 3 months immediately preceding the date of admission of the animals to the semen collection centre; II.4.7. were sept in semen collection centres: II.4.7. works are not of the disease referred to pinotif II.4.5.2 has been reported for at least 30 days immediately prior to the due of solvegion of the semen, and III.4.7. which were not situated in a restricted zone established due to diseases referred to in point II.4.5.1; III.4.7. which were not situated in the semen, and III.4.7. which were not situated in the semen, and III.4.7. which were not situated in the semen, and III.4.7. which were not situated in the semen and III.4.5.2 has been reported within a 10-km radius centred on the semen collection enters for all east a place of the collection.] III.4.7. where no days are supported for at least 3 months immediately prior to the date of collection of the semen and 30 days from the part per preceding 30 days, and III.4.8. where no climal, stronger due to the consignment of semen to the Union and they have been kept at that semen collection enterts for a least 30 days immediately prior to the date of collection of the semen and 30 days from due to the process provided the semen and 30 days from due to the process provided the semen and 30 days			II.4.5.4.	were not used for natural breeding;
II.4.6.2. none of the diseases referred to in point II.4.5.2 has been reported for at least the preceding 30 days; II.4.6.3. it was situated in an ear where foot and mouth disease has not been reported within a 10-km radius centred on the quarantine accommodation for at least the preceding 30 days; II.4.6.4. has had no outbreak of foot and mouth disease reported during at least 3 months immediately preceding the date of admission of the animates to the seeme collection centre; II.4.8.5. it was free from infection with Brucella abortus. Brucella melitensis and Brucella suis for at least the preceding 3 mouths; III.4.7.1. were sent in season collection centres: II.4.1. II.4.7.2. whice noise of the diseases referred to in point II.4.5.1; white noise of the diseases referred to in point II.4.5.1; white noise of the diseases referred to in point II.4.5.2 has been reported for at least 30 days immediately prior to the due of splits into the semen, and Only either results flavals including the date of the collection; [until the date of dispatals of the consignment of semen to the Union;] II.4.7.3. situatella in a near where foot and mouth disease has not been reported within a 10-km radius centred on the semen collection reporting foot least ple proceding 30 days; and Only or [were free from foot and massamh disease for at least 3 months immediately prior to the date of collection of the semen and 30 days from the date of the collection; [were free from foot ambassamh disease for at least 3 months immediately prior to the date of collection of the semen and 30 days from the date of the collection; [were free from footward massamh disease for at least 30 days inmediately prior to the date of collection of the semen and 30 days from the date of the collection of semen to the Union and they have been kept at that seme collection centre foot a feet as 30 days inmediately prior to the date of collection of the semen and 30 days from the date of collection; and the prior to the date of collection of the semen.] III		II.4.6.	at least the sa	ime health status were present, which on the day of their admission to the semen collection centre complied with the
11.4.6.3. it was situated in an area where floot and mouth disease has not been reported within a 10-km radius centred on the quarantine accommodation for at least the preceding 30 days; 11.4.6.1. has had no outbreak of foot and mouth disease reported during at least 3 months immediately preceding the date of admission of the animals to the semen collection centre; 11.4.1. were for in segare collection centres: 11.4.2. where no collection centres: 11.4.3. II.4.7.2. white notes of the diseases referred to in point II.4.5.1; 11.4.7.2. white notes of the diseases referred to in point II.4.5.2 has been reported for at least 30 days immediately prior to the olle of softenion of the semen, and 10.00 either in the olle of softenion of the semen, and 10.00 either in the olle of softenion of the semen, and 10.00 either in the olle of softenion of the semen and 10.00 either in the olle of softenion of the semen of the collection; in the ollegate of softenion of the semen and 20.00 either in the ollegate of softenion of the semen and 30.00 sys from the date of dispatch of the consignment of semen to the Union.] 11.4.7.3. where free from format agenth diseases for at least 3 months immediately prior to the date of collection of the semen and 30.00 sys from the dates of adjust in the date of dispatch of the consignment of semen to the Union and they have been kept at that semen collection centre for fort least 30 days immediately prior to the date of collection of the semen and until the date of dispatch of the consignment of semen to the Union and they have been kept at that semen collection centre forts least 30 days immediately prior to the date of collection of the semen and until the date of dispatch of the consignment of semen to the Union and they have been kept at that semen collection centre forts least 30 days immediately prior to the date of collection of the semen. 11.4.7.4. where no clinical, serological, variogical or pathogical evidence of infection with Autorisally via disease virus has been preported		•	II.4.6.1.	it was not situated in a restricted zone established due to diseases referred to in point II.4.5.1;
II.4.6.4. As had no outbrack of foot and month disease reported during at least 3 months immediately preceding the date of admission of the animals to the semen collection centre:			II.4.6.2.	none of the diseases referred to in point II.4.5.2 has been reported for at least the preceding 30 days;
admission of the animals to the semen collection centre; it was free from infection with <i>Brucella abortus, Brucella melitensis</i> and <i>Brucella suis</i> for at least the preceding 3 months; in 4.1. in 4			II.4.6.3.	
II.4.7. were sept in senten collection centres: II.4.9. which were not situated in a restricted zone established due to diseases referred to in point II.4.5.1; II.4.7.2 which were not situated in a restricted zone established due to diseases referred to in point II.4.5.2 has been reported for at least 30 days immediately prior to the date of of origination of the semen, and Olide inter a face of which are a considerable of the consignment of semen to the Union; II.4.7.3. situated an an area where foot and mouth disease has not been reported within a 10-km radius centred on the semen collection entre for at least the preceding 30 days; and Olide inter semen and an area where foot and mouth disease has not been reported within a 10-km radius centred on the semen and 30 days froughts date of selection.] Oliver free from foot and magnith diseases for at least 3 months immediately prior to the date of collection of the semen and 30 days froughts date of selection.] II.4.7.4. where no clinical, sendaged, a vincide of consignment of semen to the Union and they have been kept at that semen collection centre foral least 30 days immediately prior to the date of collection of the semen; II.4.8. have been subjected to the following tests, carried out whim 30 days immediately prior to the date of admission and at least 30 days immediately prior to the date of admission and at least 30 days immediately prior to the date of admission and at least 30 days immediately prior to the date of admission and at least 30 days immediately prior to the date of admission and at least 30 days immediately prior to the date of admission and at least 30 days immediately prior to the date of admission and at least 30 days immediately prior to the date of admission and at least 30 days immediately prior to the date of admission and at least 30 days immediately prior to the date of admission and at least 30 days immediately prior to the date of admission and at least 30 days immediately prior to the date of admission and at least 30 day		O		
II.4. 9. II.4. 1. II.4. 2. II.4. 3. II.4. 3. II.4. 3. II.4. 3. II.4. 3. II.4. 6. II.4. 6. II.4. 6. II.4. 6. II.4. 6. II.4. 7. III.4. 8. II				months;
II.4.7.2. whose name of the diseases referred to in point II.4.5.2 has been reported for at least 30 days immediately prior to the date of collection of the semen, and (10) either (10) or [10] (10) o		11.4.7.	1	
the date of solutation of the semen, and (1)(4) either (1)(5) or (1)(4) either (1)(5) or (1)(4) either (1)(5) or (1)(4) either (1)(6) either (1)(6) either (1)(7)(7) either (2)(8) or (2)(1)(8) or (3)(1)(9) either (3)(1)(9) either (4)(1)(1)(1)(1)(1)(1)(1)(1)(1)(1)(1)(1)(1)				
[II.4.7.3. situated in an area where foot and mouth disease has not been reported within a 10-km radius centred on the semen collection reprints for at least the preceding 30 days; and [III.4.7.4] [were free from foot and mouth disease for at least 3 months immediately prior to the date of collection of the semen and 30 days from the date of collection; [III.4.7.4] [were free from foot and mash disease for at least 3 months immediately prior to the date of collection of the semen and until the date of dispatch of the consignment of semen to the Union and they have been kept at that semen collection centre for a least 30 days immediately prior to the date of collection of the semen and until the date of dispatch of the consignment of semen to the Union and they have been kept at that semen collection centre for a least 30 days immediately prior to the date of collection of the semen.] II.4.7.4. where no clinical, serological, virological or pathological evidence of infection with Aujeszky's disease virus had been reported for a period comprising at least 30 days immediately prior to the date of collection of the semen.] II.4.8. have been subjected to the following tests, carried out whim 30 days impediately prior to the date of collection of the semen.] III.4.8. have been subjected to the following tests, carried out whim 30 days impediately prior to the date of collection of understance with Part 2, Chapter 1, point 1(b), of Annex II to Delegated Regulation (EU) 2020/686: III.4.8.1. as regards infection with Brucella abortus, B. melitensis and B. sub, a buffered Brucella antigen test (rose Bengal test), a competitive ELISA or an indirect ELISA for the detection of antibodies to smooth Brucella species; and season of the sement of the quisexply sides as virus or to glycoprotein B (ADV-gB) or glycoprotein B (ADV-gB) or glycoprotein G (ADV-gB) or glyc				the date of collection of the semen, and
II.4.7.3. situated in an area where foot and mouth disease has not been reported within a 10-km radius centred on the semen collection entres for at least the preceding 30 days; and [were free from foot and missing disease for at least 3 months immediately prior to the date of collection of the semen and 30 days from the date of collection.] [Were free from foot and missing disease for at least 3 months immediately prior to the date of collection of the semen and until the date of dispatch of the consignment of semen to the Union and they have been kept at that semen collection centre for at least 30 days immediately prior to the date of collection of the semen; and until the date of dispatch of the consignment of semen to the Union and they have been kept at that semen collection centre for at least 30 days immediately prior to the date of collection of the semen; II.4.7.4. where no clinical, serological, virological or pathological evidence of infection with Aujeszky's disease virus had been reported for a period comprising at least 30 days immediately prior to the date of admission and at least 30 days immediately prior to the date of collection of the semen; II.4.8. have been subjected to the following tests, carried out within 30 aways imprediately prior to the date of commencement of the quarantine referred to in point II.4.6, with negative results, separed in accordance with Part 2, Chapter I, point 1(b), of Annex II to Delegated Regulation (EU) 2020/686: II.4.8.1. as regards infection with Brucella abortus, B. melitensis and B. susy, a buffered Brucella antigen test (rose Bengal test), a competitive ELISA or an indirect ELISA for the detection of antibodies to smooth Brucella species; are regards infection with Aujeszky's disease virus. [In the case of non-vaccinated animals, an ELISA to detect antibodies to bits whole Aujeszky's disease virus or to glycoprotein B (ADV-gB) or glycoprotein D (ADV-gD) of the virus or a serum neutralisation test; in the case of animals vaccinated with a gE deleted vacc				
collection entres for at least the preceding 30 days; and (i)(4) either (ii)(6) either (iii)(7) or [were free from foot and thought disease for at least 3 months immediately prior to the date of collection of the semen and 30 days from the date of collection;] (iii)(8) or [were free from foot and month disease for at least 3 months immediately prior to the date of collection of the semen and until the date of dispatch of the consignment of semen to the Union and they have been kept at that seme collection centre for at least 30 days immediately prior to the date of collection of the semen;] II.4.7.4. where no clinical, serological, virological or pathological evidence of infection with Aujeszky's disease virus had been reported for a period comprising at least 30 days immediately prior to the date of admission and at least 30 days immediately prior to the date of admission and at least 30 days immediately prior to the date of admission and at least 30 days immediately prior to the date of commencement of the quarantine referred to in point II.4.6, with negative results, required in accordance with Part 2, Chapter I, point 1(b), of Annex II to Delegated Regulation (EU) 2020/686: II.4.8.1. as regards infection with Brucella abortus, B. meltrensis and B. sufs. abuffered Brucella antigen test (rose Bengal test), a competitive ELISA or an indirect ELISA for the detection of annibodies to smooth Brucella species; II.4.8.2. as regards infection with Aujeszky's disease virus. (i) either (ii) or [iii the case of non-vaccinated animals, an ELISA to detect antibodies to the wild; Aujeszky's disease virus or to glycoprotein B (ADV-gB) of glycoprotein D (ADV-gD) of the virus or of serum neutralisation test.] (iii) Iii the case of animals vaccinated with a gE deleted vaccine, an ELISA to detect antibodies to glycoprotein E (ADV-gB) of Aujeszky's disease virus;] (iv) III.4.8.1 as regards infection with porcine reproductive and respiratory syndrome virus, a serological test (the immunoperoxidase monologyer assay				
semen and 30 days from the date of collection;] (iver free from foot and mouth disease for at least 3 months immediately prior to the date of collection of the semen and until the date of dispatch of the consignment of semen to the Union and they have been kept at that semen collection centre for at least 30 days immediately prior to the date of collection of the semen;] II.4.7.4. where no clinical, serological, virological or pathological evidence of infection with Aujeszky's disease virus had been reported for a period comprising at least 30 days immediately prior to the date of admission and at least 30 days immediately prior to the date of admission and at least 30 days immediately prior to the date of collection of the semen; II.4.8. have been subjected to the following tests, carried out within 30 days immediately prior to the date of commencement of the quarantine referred to in point II.4.6, with negative results, required in accordance with Part 2, Chapter I, point I(b), of Annex II to Delegated Regulation (EU) 2020/686: II.4.8.1. as regards infection with Brucella abortus, B. melituenists and B. suts, a buffered Brucella antigen test (rose Bengal test), a competitive ELISA or an indirect ELISA for the detection of ambodies to smooth Brucella species; II.4.8.2. as regards infection with Aujeszky's disease virus, (i) either (in the case of non-vaccinated animals, an ELISA to detect ambodies to the whole Aujeszky's disease virus or to glycoprotein B (ADV-gB) or glycoprotein D (ADV-gD) of the virus or a serum neutralisation test; (ii) fine case of animals vaccinated with a gE deleted vaccine, an ELISA to detect ambodies to glycoprotein E (ADV-gB) of Aujeszky's disease virus; (ii) as regards classical swine fever, an antibody ELISA or serum neutralisation test; in Jia case of animals coming from a third country or territory, or zone thereof where classical swine fever has been reported or vaccination against this disease has been practised for the preceding 12 months; II.4.9. have been subjected t				collection centres for at least the preceding 30 days; and
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glycoprotein B (ADV-gB) or glycoprotein D (ADV-gD) of the virus or a serum neutralisation test;] (1) or [in the case of animals vaccinated with a gE deleted vaccine, an ELISA to detect antibodies to glycoprotein E (ADV-gE) of Aujeszky's disease virus;] (1) [II.4.8.3. as regards classical swine fever, an antibody ELISA or serum neutralisation test, in the case of animals coming from a third country or territory, or zone thereof where classical swine fever has been reported or vaccination against this disease has been practised for the preceding 12 months;] II.4.8.4. as regards infection with porcine reproductive and respiratory syndrome virus, a serological test (the immunoperoxidase monolayer assay (IPMA), immunofluorescence assay (IFA), or ELISA); II.4.9. have been subjected to the following tests, carried out on samples taken at least 21 days after the commencement of the quarantine referred to in point II.4.6, with negative results, required in accordance with Part 2, Chapter I, point 1(c), of Annex II to Delegated Regulation (EU) 2020/686: II.4.9.1. as regards infection with Brucella abortus, B. melitensis and B. suis, a buffered Brucella antigen test (rose Bengal test), a competitive ELISA or an indirect ELISA for the detection of antibodies to smooth Brucella species; II.4.9.2. as regards infection with Aujeszky's disease virus (1) either [in the case of non-vaccinated animals, an ELISA to detect antibodies to the whole Aujeszky's disease virus or to glycoprotein B (ADV-gB) or glycoprotein D (ADV-gD) of the virus or a serum neutralisation test;]				
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referred to in point II.4.6, with negative results, required in accordance with Part 2, Chapter I, point 1(c), of Annex II to Delegated Regulation (EU) 2020/686: II.4.9.1. as regards infection with Brucella abortus, B. melitensis and B. suis, a buffered Brucella antigen test (rose Bengal test), a competitive ELISA or an indirect ELISA for the detection of antibodies to smooth Brucella species; II.4.9.2. as regards infection with Aujeszky's disease virus (i) either [in the case of non-vaccinated animals, an ELISA to detect antibodies to the whole Aujeszky's disease virus or to glycoprotein B (ADV-gB) or glycoprotein D (ADV-gD) of the virus or a serum neutralisation test;] (i) or [in the case of animals vaccinated with a gE deleted vaccine, an ELISA to detect antibodies to glycoprotein E			II.4.8.4.	
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[in the case of non-vaccinated animals, an ELISA to detect antibodies to the whole Aujeszky's disease virus or to glycoprotein B (ADV-gB) or glycoprotein D (ADV-gD) of the virus or a serum neutralisation test;] [in the case of animals vaccinated with a gE deleted vaccine, an ELISA to detect antibodies to glycoprotein E			II.4.9.1.	test), a competitive ELISA or an indirect ELISA for the detection of antibodies to smooth Brucella species;
glycoprotein B (ADV-gB) or glycoprotein D (ADV-gD) of the virus or a serum neutralisation test;] [in the case of animals vaccinated with a gE deleted vaccine, an ELISA to detect antibodies to glycoprotein E				
			⁽¹⁾ or	

				11.a	Certificate reference
U	NITED KINGDOM				
		11.402	1-i-f4:i-di14:11	1	

	11.4.9.3.	as regards infection with potenic reproductive and respiratory syndrome virus, a serological test (IFMA, IFA, or
		ELISA) and a test for virus genome (reverse-transcription polymerase chain reaction (RT-PCR), nested set RT-
		PCR, real-time RT-PCR);
II 4 10	harra haam and	i - 4 - 1 - 4

- II.4.10. have been subjected, at semen collection centre, to the following compulsory routine tests, required in accordance with Part 2, Chapter I, point 2(a), of Annex II to Delegated Regulation (EU) 2020/686:
 - II.4.10.1. as regards infection with *Brucella abortus*, *B. melitensis* and *B. suis*, a buffered Brucella antigen test (rose Bengal test), a competitive ELISA or an indirect ELISA for the detection of antibodies to smooth *Brucella* species;
 - II.4.10.2. as regards infection with Aujeszky's disease virus
 - (i) either [in the case of non-vaccinated animals, an ELISA to detect antibodies to the whole Aujeszky's disease virus or to glycoprotein B (ADV-gB) or glycoprotein D (ADV-gD) of the virus or a serum neutralisation test;]
 - (1) or [in the case of animals vaccinated with a gE deleted vaccine, an ELISA to detect antibodies to glycoprotein E (ADV-gE) of Aujeszky's disease virus;]
 - as regards classical swine fever, an antibody ELISA or serum neutralisation test, in the case of animals coming from a third country or territory, or zone thereof where classical swine fever has been reported or vaccination against this disease has been practised for the preceding 12 months;]
 - II.4.10.4. as regards infection with porcine reproductive and respiratory syndrome virus, a serological test (IPMA, IFA, or ELISA);
- II.4.11. have been subjected to the tests referred to in point II.4.10 carried out, in accordance with Part 2, Chapter I, point 2(b), of Annex II to Delegated Regulation (EU) 2020/686, on samples taken from
 - (1) either [all animals immediately prior to the date of dispatch from the semen collection centre, or upon the date of arrival at the slaughterhouse, and in no case later than 12 months from the date of admission to the semen collection centre.]
 - [at least 25% of the animals in the semen collection centre every 3 months to test for infection with *Brucella abortus*, *Brucella melitensis* and *Brucella surs*, infection with Aujeszky's disease virus and classical swine fever, and at least 10% of the animals in the semen collection centre every month to test for infection with porcine reproductive and respiratory syndrome virus.]
 - (1) or [at least 10% of the animals in the semen collection centre every month to test for infection with *Brucella abortus*, *Brucella melitensis* and *Brucella suis*, infection with Abjeszky's disease virus, classical swine fever and infection with porcine reproductive and respiratory syndrome virus.]
- II.5. The semen described in Part I:
 - II.5.1. has been collected, processed and stored in accordance with animal health requirements set out in Part 1, points 1 and 2, of Annex III to Delegated Regulation (EU) 2020/686.
 - II.5.2. is placed in straws or other packages on which the mark is applied in accordance with requirements provided for in Article 83, point (a), of Delegated Regulation (EU) 2020/692 and that mark is indicated in box I.27;
 - II.5.3. is transported in a container which:
 - II.5.3.1. was sealed and numbered prior to the date of dispatch from the semen collection centre under responsibility of the centre veterinarian, or by an official veterinarian, and the seal bears the number as indicated in box I.19;
 - II.5.3.2. has been cleaned and either disinfected or sterilised before use, or is single-use container;
 - (1)(4) [II.5.3.3. has been filled in with a cryogenic agent which has not been previously used for other products.]
- (1) [II.6. Where an antibiotic or a mixture of antibiotics was added to the semen:
 - II.6.1. The following antibiotic or mixture of antibiotics has been added to the semen after final dilution, or is contained in the used semen diluents: _______; (6)
 - II.6.2. Immediately after the addition of the antibiotic(s), and before any possible freezing, the diluted semen was kept at a temperature of at least 15°C for not less than 45 minutes, or under a time-temperature regime with a documented equivalent bactericidal activity.]

Notes

"Porcine animal" means a porcine animal as defined in Article 2, point (4), of Delegated Regulation (EU) 2020/686.

This animal health certificate is intended for the entry into the Union of semen of porcine animals, including when the Union is not the final destination of the semen.

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 Windsor Framework (see Joint Declaration No 1/2023 of the Union and the United Kingdom in the Joint Committee established by 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 of 24 March 2023, OJ L 102, 17.4.2023, p. 87) in conjunction with Annex 2 to that Framework, 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 4 of Annex 1 to Commission Implementing Regulation (EU) 2020/2235.

Part I:

Box reference I.11: "Place of dispatch": Indicate the unique approval number and the name and address of the semen collection centre of dispatch

of the consignment of semen. Only semen collection centres listed in accordance with Article 233(3) of Regulation (EU) 2016/429 on the Commission website:

https://ec.europa.eu/food/animals/semen/porcine_en.

Box reference I.12: "Place of destination": Indicate the address and unique registration or approval number of the establishment of destination of

Box reference I.19: Seal number shall be indicated.

Box reference I.24: Total number of packages shall correspond to the number of containers.

the consignment of semen.

Box reference I.27: "Type": indicate semen.

"Identification number": Indicate identification number of each donor animal.

"Identification mark": Indicate mark on the straw or other packages where semen of the consignment is placed.

"Date of collection/production": Indicate the date on which semen of the consignment was collected.

"Approval or registration number of plant/establishment/centre": Indicate the unique approval number of the semen collection

centre where the semen of the consignment was collected.

"Quantity": Indicate number of straws or other packages with the same mark.

Part II

Delete if not applicable

Only for a third country or territory, or zone thereof with an opening date in accordance with column 9 of the table in Part 1 of Annex II to Implementing Regulation (EU) 2021/404.

Only semen collection centres listed in accordance with Article 233(3) of Regulation (EU) 2016/429 on the Commission website: https://ec.europa.eu/food/animals/semen/porcine_en.

(4) Applicable for frozen semen.

(5) Applicable for fresh and chilled semen.

(6) Insert the name(s) of the antibiotic(s) added and its (their) concentration or the commercial name of the semen diluent containing antibiotic(s).

Official veterinarian

Name (in capital letters)

Date Qualification and title

Stamp Signature