

UNITED KINGDOM

Animal Health certificate to the EU

Part I: Description of consignment	I.1 Consignor/Exporter		I.2 Certificate reference		I.2a	
	Name					
	Address		I.3 Central Competent Authority			
			DEPARTMENT FOR ENVIRONMENT, FOOD & RURAL AFFAIRS			
	Country		ISO country code		I.4 Local Competent Authority	
					ANIMAL AND PLANT HEALTH AGENCY	
	I.5 Consignee/Importer		I.6 Operator responsible for the consignment			
	Name		Name			
	Address		Address			
	Country		ISO country code		Country	
					ISO country code	
	I.7 Country of origin		ISO country code		I.9 Country of destination	
					ISO country code	
	I.8 Region of origin		Code		I.10 Region of destination	
					Code	
	I.11 Place of dispatch		Registration/Approval No		I.12 Place of destination	
	Name				Registration/Approval No	
	Address				Name	
					Address	
	Country		ISO country code		Country	
					ISO country code	
	I.13 Place of loading		I.14 Date and time of departure			
	I.15 Means of transport		I.16 Entry Border Control Post			
	☐ Aircraft		☐ Vessel			
	☐ Railway		☐ Road vehicle		I.17 Accompanying documents	
	Identification		Type			
			Code			
			Country			
			ISO country code			
			Commercial document reference			
	I.18 Transport conditions		☐ Ambient			
			☐ Chilled			
			☐ Frozen			
	I.19 Container number/Seal number					
	Container No			Seal No		
	I.20 Certified as or for					
	☐ Germinal products					
	I.21			I.22		
	☐ For transit			☐ For internal market		
	Third country			ISO country code		
				I.23		
	I.24 Total number of packages		I.25 Total quantity		I.26	

UNITED KINGDOM

I.27 Description of consignment

1	CN code	Species	Subspecies/Category	Identification number	Quantity
	Type	Approval or registration number of plant/establishment/centre	Identification mark	Date of collection/production	
2	CN code	Species	Subspecies/Category	Identification number	Quantity
	Type	Approval or registration number of plant/establishment/centre	Identification mark	Date of collection/production	
3	CN code	Species	Subspecies/Category	Identification number	Quantity
	Type	Approval or registration number of plant/establishment/centre	Identification mark	Date of collection/production	
4	CN code	Species	Subspecies/Category	Identification number	Quantity
	Type	Approval or registration number of plant/establishment/centre	Identification mark	Date of collection/production	
5	CN code	Species	Subspecies/Category	Identification number	Quantity
	Type	Approval or registration number of plant/establishment/centre	Identification mark	Date of collection/production	

UNITED KINGDOM

II.a Certificate reference

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II. Health information

I, the undersigned official veterinarian, hereby certify that:

- II.1. The germinal product storage centre ⁽¹⁾ described in box I.11 at which the [semen] ⁽²⁾ [oocytes] ⁽²⁾ [in vivo derived embryos] ⁽²⁾ [in vitro produced embryos] ⁽²⁾ [micromanipulated embryos] ⁽²⁾ to be dispatched to the Union was/were stored:
- II.1.1. is located in a third country or territory, or zone thereof:
- II.1.1.1. authorised for the entry into the Union of [semen] ⁽²⁾ [oocytes] ⁽²⁾ [embryos] ⁽²⁾ of bovine animals and listed in Annex IX to Commission Implementing Regulation (EU) 2021/404;
- ⁽²⁾ either II.1.1.2. where foot and mouth disease was not reported for at least 24 months immediately prior to the date of [collection] ⁽²⁾ [production] ⁽²⁾ of the [semen] ⁽²⁾ [oocytes] ⁽²⁾ [embryos] ⁽²⁾ and until the date of its/their dispatch;]
- ⁽²⁾⁽³⁾ or II.1.1.2. 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] ⁽²⁾ [production] ⁽²⁾ of the [semen] ⁽²⁾ [oocytes] ⁽²⁾ [embryos] ⁽²⁾ and until the date of its/their dispatch;]
- II.1.1.3. where infection with rinderpest virus, infection with Rift Valley fever virus, contagious bovine pleuropneumonia and lumpy skin disease were not reported for at least 12 months immediately prior to the date of [collection] ⁽²⁾ [production] ⁽²⁾ of the [semen] ⁽²⁾ [oocytes] ⁽²⁾ [embryos] ⁽²⁾ and until the date of its/their dispatch;
- II.1.1.4. where no vaccination against infection with rinderpest virus, infection with Rift Valley fever virus and contagious bovine pleuropneumonia has been carried out for at least 12 months immediately prior to the date of [collection] ⁽²⁾ [production] ⁽²⁾ of the [semen] ⁽²⁾ [oocytes] ⁽²⁾ [embryos] ⁽²⁾ and until the date of its/their dispatch, and no vaccinated animals entered into the third country or territory, or zone thereof during that period, and
- ⁽²⁾ 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;]
- ⁽²⁾ 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.1.2. is approved and listed by the competent authority of the third country or territory;
- II.1.3. complies with requirements as regards responsibilities, operational procedures, facilities and equipment set out in Part 5 of Annex I to Commission Delegated Regulation (EU) 2020/686.
- II.2. The [semen] ⁽²⁾ [oocytes] ⁽²⁾ [embryos] ⁽²⁾ described in Part I is/are intended for artificial reproduction and:
- II.2.1. has/have been [collected] ⁽²⁾ [produced] ⁽²⁾ [processed] ⁽²⁾ [stored] ⁽²⁾ [in a semen collection centre] ⁽²⁾⁽⁴⁾ [by an embryo collection team] ⁽²⁾⁽⁴⁾ [by an embryo production team] ⁽²⁾⁽⁴⁾ [and] ⁽²⁾ [processed] ⁽²⁾ [stored] ⁽²⁾ [in a germinal product processing establishment] ⁽²⁾⁽⁴⁾ and stored in a germinal product storage centre ⁽⁴⁾ complying with requirements set out in [Part 1] ⁽²⁾ [Part 2] ⁽²⁾ [Part 3] ⁽²⁾ [Part 4] ⁽²⁾ [Part 5] ⁽²⁾ of Annex I to Delegated Regulation (EU) 2020/686, and:
- ⁽²⁾ either [located in the third country or territory of dispatch to the Union;]
- ⁽²⁾⁽⁵⁾ and/or [located in _____, and has/have been introduced into the third country or territory of dispatch to the Union under conditions at least as strict as for the entry into the Union of [semen] ⁽²⁾ [oocytes] ⁽²⁾ [in vivo derived embryos] ⁽²⁾ [in vitro produced embryos] ⁽²⁾ of bovine animals in accordance with Regulation (EU) 2016/429 and Commission Delegated Regulation (EU) 2020/692;]
- II.2.2. was/were moved to the germinal product storage centre described in box I.11 under conditions at least as strict as described in:
- ⁽²⁾ either [Model BOV-SEM-A-ENTRY ⁽⁶⁾;]
- ⁽²⁾ and/or [Model BOV-SEM-B-ENTRY ⁽⁶⁾;]
- ⁽²⁾ and/or [Model BOV-SEM-C-ENTRY ⁽⁶⁾;]
- ⁽²⁾ and/or [Model 1 in Section A of Part 1 of Annex II to Commission Implementing Decision 2011/630/EU ⁽⁶⁾;]
- ⁽²⁾ and/or [Model 2 in Section B of Part 1 of Annex II to Implementing Decision 2011/630/EU ⁽⁶⁾;]
- ⁽²⁾ and/or [Model 3 in Section C of Part 1 of Annex II to Implementing Decision 2011/630/EU ⁽⁶⁾;]
- ⁽²⁾ and/or [Model BOV-OOCYTES-EMB-A-ENTRY ⁽⁶⁾;]
- ⁽²⁾ and/or [Model BOV-in-vivo-EMB-B-ENTRY ⁽⁶⁾;]
- ⁽²⁾ and/or [Model BOV-in-vitro-EMB-C-ENTRY ⁽⁶⁾;]
- ⁽²⁾ and/or [Model BOV-in-vitro-EMB-D-ENTRY ⁽⁶⁾;]
- ⁽²⁾ and/or [Model BOV-GP-PROCESSING-ENTRY ⁽⁶⁾;]
- ⁽²⁾ and/or [Model BOV-GP-STORAGE-ENTRY ⁽⁶⁾;]
- II.2.3. has/have been collected, processed and stored in accordance with animal health requirements set out in Annex III to Delegated Regulation (EU) 2020/686;
- II.2.4. is/are 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.2.5. is/are transported in a container which:
- II.2.5.1. was sealed and numbered prior to the date of dispatch from the germinal product storage 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.2.5.2. has been cleaned and either disinfected or sterilised before use, or is single-use container;
- ⁽²⁾⁽⁷⁾ II.2.5.3. has been filled in with a cryogenic agent which has not been previously used for other products.]

Part II: Certification

UNITED KINGDOM

II.a Certificate reference

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- (2) (8) [II.2.6. is/are placed in straws or other packages which are securely and hermetically sealed;
II.2.7. is/are transported in a container where the different types are separated from each other by physical compartments or by being placed in secondary protective bags.]

Notes

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

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 I 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 germinal product storage centre of dispatch of the consignment of semen, oocytes and/or embryos. Only germinal product storage centre listed in accordance with Article 233(3) of Regulation (EU) 2016/429 on the Commission website: http://ec.europa.eu/food/animal/semen_ova/bovine/ova_embryos_en.htm.
- Box reference I.12: "Place of destination": Indicate the address and unique registration or approval number of the establishment of destination of the consignment of semen, oocytes and/or embryos.
- Box reference I.17: "Accompanying documents": Number(s) of related original animal health certificate(s) shall correspond to the serial number of the individual official document(s) or animal health certificate(s) that accompanied the semen, oocytes and/or embryos described in Part I from the semen collection centre, where the semen was collected, and/or from the embryo collection team and/or the embryo production team by which the oocytes and/or embryos were collected or produced, and/or from the germinal product processing establishment where the semen, oocytes or embryos were processed and stored, and/or from the germinal product storage centre where the semen, oocytes or embryos were stored, to the germinal product storage centre described in box I.11. The original(s) of those document(s) or those animal health certificate(s) or the officially endorsed copies thereof shall be attached to this animal health certificate.
- Box reference I.19: Seal number shall be indicated.
- Box reference I.24: Total number of packages shall correspond to the number of containers.
- Box reference I.27: "Species": Select amongst "*Bos taurus*", "*Bison bison*" or "*Bubalus bubalis*" as appropriate.
"Type": Specify if semen, *in vivo* derived embryos, *in vivo* derived oocytes, *in vitro* produced embryos or micromanipulated embryos.
"Identification number": Indicate identification number of each donor animal.
"Identification mark": Indicate mark on the straw or other packages where semen, oocytes and/or embryos of the consignment are placed.
"Date of collection/production": Indicate the date on which semen, oocytes and/or embryos of the consignment was/were collected or produced.
"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, and/or of the embryo collection team and/or the embryo production team by which oocytes or embryos of the consignment were collected or produced.
"Quantity": Indicate number of straws or other packages with the same mark.

Part II:

- (1) Only germinal product storage centres listed in accordance with Article 233(3) of Regulation (EU) 2016/429 on the Commission website: http://ec.europa.eu/food/animal/semen_ova/bovine/ova_embryos_en.htm.
- (2) Delete if not applicable.
- (3) 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.
- (4) Only approved germinal product establishments listed in accordance with Article 233(3) of Regulation (EU) 2016/429 on the Commission website: http://ec.europa.eu/food/animal/semen_ova/bovine/index_en.htm.
- (5) Only a third country or territory, or zone thereof listed in Annex IX to Implementing Regulation (EU) 2021/404 and the Member States.
- (6) The original(s) of the document(s) or the animal health certificate(s) or the officially endorsed copies of thereof that accompanied the semen, oocytes or embryos described in Part I from the semen collection centre, where the semen was collected, and/or from the embryo collection team or the embryo production team by which the oocytes and/or embryos were collected or produced, and/or from the germinal product processing establishment where the semen, oocytes or embryos were processed and stored, and/or from the germinal product storage centre where the semen, oocytes or embryos were stored, to the germinal product storage centre of dispatch of the semen, oocytes and/or embryos described in box I.11 shall be attached to this animal health certificate.

II.a Certificate reference

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UNITED KINGDOM

(7) Applicable for frozen semen, oocytes or embryos.

(8) Applicable for consignments where semen, oocytes, *in vivo* derived embryos, *in vitro* produced embryos and micromanipulated embryos of bovine animals are placed and transported in one container.

Official veterinarian

Name (in capital letters)

Date

Qualification and title

Stamp

Signature