

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
	Country			ISO country code		ISO country code
	I.7 Country of origin			I.9 Country of destination		
ISO country code			ISO country code			
I.8 Region of origin			I.10 Region of destination			
Code			Code			
I.11 Place of dispatch			I.12 Place of destination			
Registration/Approval No			Registration/Approval No			
Name			Name			
Address			Address			
Country			ISO country code		Country	
Country			ISO country code		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			
<input type="checkbox"/> Aircraft <input type="checkbox"/> Vessel <input type="checkbox"/> Railway <input type="checkbox"/> Road vehicle			I.17			
Identification						
I.18 Transport conditions			<input type="checkbox"/> Ambient <input type="checkbox"/> Chilled <input type="checkbox"/> Frozen			
I.19 Container number/Seal number						
Container No			Seal No			
I.20 Certified as or for						
<input type="checkbox"/> Germinal products						
I.21			I.22			
<input type="checkbox"/> For transit			<input type="checkbox"/> For internal market			
Third country			ISO country code		I.23	
I.24 Total number of packages			I.25 Total quantity		I.26	

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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

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II.a Certificate reference

II. Health information

I, the undersigned official veterinarian, hereby certify, that:

- II.1. The semen⁽¹⁾/ *in vivo* derived embryos⁽¹⁾/ oocytes⁽¹⁾/ *in vitro* produced embryos⁽¹⁾/ micromanipulated embryos⁽¹⁾ described in Part I is/are intended for artificial reproduction and was/were obtained from donor animals which
- II.1.1. originate from a third country, territory or zone thereof authorised for entry into the Union of the particular species and category of animals and listed in Annex III to Commission Implementing Regulation (EU) 2021/404;
- II.1.2. originate from a confined establishment in the third country, territory or zone of origin, which is included in a list of confined establishments, established in accordance with Article 29 of Commission Delegated Regulation (EU) 2020/692, from which the entry of animals of specific species into the Union may be authorised;
- II.1.3. do not come from an establishment, nor have been in contact with animals from an establishment, situated in a restricted zone established due to the occurrence of a category A disease, referred to in the Annex to Commission Implementing Regulation (EU) 2018/1882, or of an emerging disease relevant for species of those kept terrestrial animals;
- II.1.4. come from an establishment where no category D disease, relevant for species of those kept terrestrial animals as referred to in the Annex to Implementing Regulation (EU) 2018/1882, has been reported for a period of at least the preceding 30 days;
- II.1.5. have remained in a single confined establishment of origin for a period of at least 30 days prior to the collection of the semen⁽¹⁾/ oocytes⁽¹⁾/ embryos⁽¹⁾ intended for entry into the Union;
- ⁽¹⁾⁽²⁾II.1.6. are bovine, porcine, ovine, caprine or equine animals and they are identified in accordance with Article 21 of Delegated Regulation (EU) 2020/692;] or
- ⁽¹⁾⁽³⁾II.1.6. are terrestrial animals other than bovine, porcine, ovine, caprine or equine animals and they are identified and registered in accordance with the rules of the confined establishment;]
- II.1.7. have been clinically examined by the establishment veterinarian responsible for the activities carried out at the confined establishment and showed no disease symptoms on the day of collection of the semen⁽¹⁾/ oocytes⁽¹⁾/ embryos⁽¹⁾;
- II.1.8. as much as possible, were not used for natural breeding during a period of at least 30 days prior to the date of collection of the semen⁽¹⁾/ oocytes⁽¹⁾/ embryos⁽¹⁾ and during the collection period.
- II.2. The semen⁽¹⁾/ oocytes⁽¹⁾/ embryos⁽¹⁾ described in Part I
- II.2.1. is/are placed in straws or other packages on which the mark is applied in accordance with requirements provided for in
- ⁽¹⁾⁽²⁾[Article 83(a) of Delegated Regulation (EU) 2020/692 and that mark is indicated in Box I.27;]
- ⁽¹⁾⁽³⁾[Article 119(a) of Delegated Regulation (EU) 2020/692 and that mark is indicated in Box I.27;]
- II.2.2. is/are placed in a transport container which:
- II.2.2.1. was sealed and numbered prior to the dispatch from the confined establishment by the establishment veterinarian responsible for the activities of the confined establishment and the seal bears the number as indicated in Box I.19;
- II.2.2.2. has been cleaned and either disinfected or sterilised before use, or is single-use container;
- ⁽¹⁾⁽⁴⁾II.2.2.3. has been filled in with the cryogenic agent which not have been previously used for other products.]
- ⁽¹⁾⁽²⁾⁽⁵⁾II.2.3. is/are placed in straws or other packages which are securely and hermetically sealed;
- II.2.4. is/are transported in a container where they are separated from each other by physical compartments or by being placed in secondary protective bags.]
- II.3. The consignment of semen⁽¹⁾/ oocytes⁽¹⁾/ embryos⁽¹⁾
- II.3.1. is destined to a confined establishment in the Union, which is approved in accordance with Article 95 of Regulation (EU) 2016/429;
- II.3.2. is transported directly to the confined establishment as indicated in Box I.12.

Notes

This certificate is intended for entry into the Union of semen, oocytes and embryos of terrestrial animals kept at confined establishments, 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 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 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, if assigned by the competent authority, and the name and address of the confined establishment of dispatch of the consignment of semen, oocytes or embryos.
- Box reference I.12: "Place of destination": Indicate the name, address and unique approval number of the confined establishment of destination in the Union of the consignment of semen, oocytes or embryos.
- Box reference I.27: "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.

Part II: Certification

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II.a Certificate reference

“*Identification mark*”: Indicate mark on the straw or other packages where semen, oocytes or embryos of the consignment are placed.
 “*Date of collection/production*”: Indicate the date on which semen, oocytes or embryos of the consignment were collected or produced.
 “*Approval or registration number of plant/establishment*”: Indicate the unique approval number, if assigned by the competent authority, and the name and address of the confined establishment of the collection or production of semen, oocytes or embryos of the consignment.
 “*Quantity*”: Indicate number of straws or other packages with the same mark.

Part II:

- (1) Delete if not applicable.
- (2) Applicable for the consignment of semen, oocytes or embryos of bovine, porcine, ovine, caprine or equine animals.
- (3) Applicable for the consignment of semen, oocytes or embryos of terrestrial animals other than bovine, porcine, ovine, caprine or equine animals.
- (4) Applicable for frozen semen, oocytes or embryos.
- (5) Applicable for the consignment where in one container oocytes, *in vivo* derived embryos, *in vitro* produced embryos and micromanipulated embryos of bovine, porcine, ovine, caprine or equine animals are placed and transported.

Official veterinarian

Name (in capital letters)

Date

Qualification and title

Stamp

Signature