UNITED KINGDOM **Animal health certificate to the EU**

Part I: Description of consignment

I.1 Consignor/Exporter
Name
Address
Country ISO country code

I.2 Certificate reference
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I.2a

I.3 Central Competent Authority
DEPARTMENT FOR ENVIRONMENT, FOOD & RURAL AFFAIRS

I.4 Local Competent Authority
ANIMAL AND PLANT HEALTH AGENCY

I.5 Consignee/Importer
Name
Address
Country ISO country code

I.6 Operator responsible for the consignment
Name
Address
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
Name
Address
Country ISO country code

I.12 Place of destination Registration/Approval No
Name
Address
Country ISO country code

I.13 Place of loading

I.14 Date and time of departure

I.15 Means of transport
☐ Aircraft ☐ Vessel
☐ Railway ☐ Road vehicle
Identification

I.16 Entry Border Control Post

I.17

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 ☐ **For transit**
Third country ISO country code

I.22 ☐ **For internal market**

I.23

I.24 Total number of packages	**I.25 Total quantity**	**I.26**

UNITED KINGDOM

II.a Certificate reference
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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

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Part II: Certification

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 or territory, or zone thereof authorised for the entry into the Union of the particular species and category of animals and listed in Annexes II to VII to Commission Implementing Regulation (EU) 2021/404, or authorised pursuant to Article 230(2) of Regulation (EU) 2016/429 by the Member State of destination, depending on the species in question;

II.1.2. originate from a confined establishment in the third country or territory, or zone thereof of origin, which is included in a list of confined establishments, established by the Member State of destination in accordance with Article 117, point (c), 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 at least 30 days immediately prior to the date of collection of the [semen] (1) [oocytes] (1) [embryos] (1) intended for entry into the Union;

II.1.5. have remained in a single confined establishment of origin for at least 30 days immediately prior to the date of collection of the [semen] (1) [oocytes] (1) [embryos] (1) intended for entry into the Union;

(1) (2) *either* [II.1.6. are bovine, porcine, ovine, caprine or equine animals and are identified in accordance with Article 21 of Delegated Regulation (EU) 2020/692;]

(1) (3) *or* [II.1.6. are terrestrial animals other than bovine, porcine, ovine, caprine or equine animals and 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 date of collection of the [semen] (1) [oocytes] (1) [embryos] (1);

II.1.8. as much as possible, were not used for natural breeding during at least 30 days immediately 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, point (a), of Delegated Regulation (EU) 2020/692 and that mark is indicated in box I.27;]

(1) (3) [Article 119, point (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 date of 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 a single-use container;

(1) (4) [II.2.2.3. has been filled in with a cryogenic agent which has not 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 the different types 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 animal health certificate is intended for the 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 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.

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II.a Certificate reference

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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.

"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 was/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 consignments of semen, oocytes or embryos of bovine, porcine, ovine, caprine or equine animals.

(3) Applicable for consignments 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 consignments where oocytes, *in vivo* derived embryos, *in vitro* produced embryos and micromanipulated embryos of bovine, porcine, ovine, caprine or equine animals are placed and transported in one container.

Official veterinarian

Name (in capital letters)

Date

Qualification and title

Stamp

Signature