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 Accompanying documents
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 ☐ **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**

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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Part II: Certification

II. Health information

I, the undersigned official veterinarian, hereby certify that:

II.1. The germinal product processing establishment (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 processed and 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) *or* [II.1.1.2. where foot and mouth disease was not reported for a period starting on the date (3) ……. (*insert 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 4 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 [processed] (2) [stored] (2) in a germinal product processing establishment (4) [and stored in a germinal product storage centre] (2) (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 into the Union;]

(2) *and/or* [located in ……. (5), 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) [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 processing establishment 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 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 processing establishment 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.]

(2) (8) [II.2.6. is/are placed in straws or other packages which are securely and hermetically sealed;

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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 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.

Part I:

Box reference I.11: "Place of dispatch": Indicate the unique approval number and the name and address of the germinal product processing establishment of dispatch of the consignment of semen, oocytes and/or embryos. Only germinal product processing 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/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 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 processing establishment described in box I.11. The original(s) of those document(s) or those 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 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 processing 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/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 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 processing establishment of dispatch of the semen, oocytes and/or embryos described in box I.11 shall be attached to this animal health certificate.

(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.

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Official veterinarian

Name (in capital letters)

Date

Qualification and title

Stamp

Signature