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 **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

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II.a Certificate reference

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

Certificate model BOV-in-vivo-EMB-B-ENTRY

II.a Certificate reference

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Part II: Certification

II. Health information

I, the undersigned, official veterinarian of the ……..…….....……..…… certify that:

(exporting country)[1]

II.1. The embryos to be exported:

II.1.1. were collected in the exporting country, which according to official findings:

II.1.1.1. was free from rinderpest during the 12 month period immediately prior to their collection;

[2]*either*[II.1.1.2. was free from foot-and-mouth disease and lumpy skin disease during the 12 month period immediately prior to their collection and did not carry out vaccination against foot-and-mouth disease or lumpy skin disease during that period.]

[2]*or* [II.1.1.2. was not free from foot-and-mouth disease or lumpy skin disease during the 12 months immediately prior to their collection or carried out vaccination against foot-and-mouth disease or lumpy skin disease during that period, and:

- the embryos were not subjected to penetration of the zona pellucida,
- the embryos were stored under approved conditions for at least 30 days immediately after their collection,
- the donor females come from holdings on which no animal was vaccinated against foot-and-mouth disease or lumpy skin disease during the 30 days prior to collection and no animal of a susceptible species showed clinical signs of foot-and-mouth disease or lumpy skin disease during the 30 days prior to, and at least the 30 days after, the embryos were collected.]

II.1.2. were collected by the embryo collection team[3] which :

- had been approved in accordance with Chapter I of Annex A to Directive 89/556/EEC;
- which carried out the collection, processing, storing and transport of the embryos in accordance with Chapter II of Annex A to Directive 89/556/EEC;
- was subject to inspection by an official veterinarian at least twice a year.

II.1.3. were collected and processed on premises situated in an area of at least 10 km radius centred on them, on which according to official findings there was no occurrence of foot-and-mouth disease, epizootic haemorrhagic disease, vesicular stomatitis, Rift Valley fever, contagious bovine pleuropneumonia or lumpy skin disease in the 30 days immediately prior to their collection and until dispatch to the Union, in the case of fresh embryos, or during the 30 days after collection, in the case of embryos subject to a mandatory storage for at least 30 days in accordance with point II.1.1.2.

II.1.4. from the time of collection until 30 days thereafter or, in the case of fresh embryos until the day of their dispatch to the Union, they were stored on premises situated in an area of at least 10 km radius centred on them, on which according to official findings there was no occurrence of foot-and-mouth disease, vesicular stomatitis, Rift Valley fever, contagious bovine pleuropneumonia or lumpy skin disease.

II.1.5. were collected from the donor females, which:

II.1.5.1. were located, during the 30 days immediately prior to collection, on premises situated in an area of at least 10 km radius centred on them, on which, according to official findings, there was no occurrence of foot-and-mouth disease, bluetongue, epizootic haemorrhagic disease, vesicular stomatitis, Rift Valley fever, contagious bovine pleuropneumonia or lumpy skin disease;

II.1.5.2. showed no clinical signs of disease on the day of collection;

II.1.5.3. spent the six months immediately prior to collection within the territory of the exporting country in no more than two herds:

- which, according to official findings, were free from tuberculosis during that time,
- which, according to official findings, were free from brucellosis during that time,
- which were free from enzootic bovine leukosis or in which no bovine animal showed clinical signs of enzootic bovine leukosis during the previous three years,
- in which no bovine animal showed clinical signs of infectious bovine rhinotracheitis/infectious pustular vulvo-vaginitis during the previous 12 months.

II.1.6. The embryos to be exported were conceived by artificial insemination using semen coming from semen collection or storage centres approved for the collection, processing and/or storage of semen by the competent authority of a third country or part thereof listed in Annex I to Implementing Decision 2011/630/EU[4] or by the competent authority of a Member State.

Notes

This certificate is intended for entry into the Union of embryos of bovine animals, including when the Union is not the final destination of the 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 I.6: "*Operator responsible for the consignment"*: this box is to be filled in only if it is a certificate for transit commodity.

Box I.11: "*Place of dispatch"* shall correspond to the embryo collection team from which the embryos are dispatched to the Union and which is listed in accordance with Article 8(2) of Directive 89/556/EEC on the Commission website: http://ec.europa.eu/food/animal/semen_ova/bovine/ova_embryos_en.htm.

Box I.19: Identification of container and seal number shall be indicated.

Box I.21: Fill in according to whether it is a transit or an import certificate.

Box I.22: Fill in according to whether it is a transit or an import certificate.

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

Box I.27: "*Species"*: Select amongst "*Bos taurus*", "*Bison bison" or "Bubalus bubalis*" as appropriate.

"*Type"*: Select "*in vivo* derived embryos".

Identification number shall correspond to the official identification of the animal.

"Date of collection/production" shall be indicated in the following format: dd.mm.yyyy

II.a Certificate reference

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"Approval or registration number of plant/establishment/centre" shall correspond to the embryo collection team by which the embryos were collected, processed and stored; and listed in accordance with Article 8(2) of Directive 89/556/EEC on the Commission website:

http://ec.europa.eu/food/animal/semen_ova/bovine/ova_embryos_en.htm.

Part II:

(1) Only third country, territory or zone thereof listed in Annex IX to Commission Implementing Regulation (EU) 2021/404 for embryos of bovine animals.

(2) Delete as appropriate.

(3) Only embryo collection teams listed in accordance with Article 8(2) of Directive 89/556/EEC on Commission website: http://ec.europa.eu/food/animal/semen_ova/bovine/ova_embryos_en.htm.

(4) OJ L 247, 24.9.2011, p. 32.

Official veterinarian

Name (in capital letters)

Date

Qualification and title

Stamp

Signature