EUROPEAN UNION **Export Health Certificate**

Part I : Details of consignment

I.1. Consignor
Name
Address
Country ISO Code

I.2. IMSOC Reference
I.2.a. Local Reference

I.5. Consignee
Name
Address
Country ISO Code

I.3. Central competent authority
I.4. Local competent authority

I.7. Country of origin ISO Code

I.9. Country of destination ISO Code

I.8. Region of origin Code

~~I.10. Region of destination~~

I.11. Place of Dispatch
Name
Address
Approval Number
Country ISO Code

I.12. Place of destination
Name
Address
Approval Number
Country ISO Code

I.13. Place of Loading
Name
Address
Approval Number
Country ISO Code

I.14. Date and time of departure

I.15. Means of Transport

Mode	International transport document	Identification

I.16 Entry Point

I.18. Transport conditions
Ambient ☐ Chilled ☐ Frozen ☐ Controlled temperature ☐

I.17. Accompanying documents
Commercial document reference
Date of issue
Country
Place of issue

I.19. Container No / Seal No

I.20. Certified as
Artificial reproduction ☐

I.21. For transit through a third country ☐
Country ISO Code
EU Exit Authority BCP code
EU Entry Authority BCP code

I.22. For transit through Member State(s) ☐
Country ISO Code

I.25. Total gross weight

I.28. Description of consignment

1. 05 PRODUCTS OF ANIMAL ORIGIN, NOT ELSEWHERE SPECIFIED OR INCLUDED
0511 Animal products not elsewhere specified or included; dead animals of Chapter|1|or 3, unfit for human consumption
051199 Other
05119985 Other

Commodity	Species	Identification number	Breed/Category	Date of freezing range

Date of collection/production

Part II: Certification

II. Health information

I, the undersigned State/official veterinarian, certify that:

1. Bovine oocytes or embryos exported to the Republic of Moldova were collected from donor females in premises and/or administrative territories that are officially free from the following contagious animal diseases:

a. foot-and-mouth disease;

b. brucellosis, tuberculosis and contagious bovine pleuropneumonia;

c. enzootic bovine leucosis.

2. Bovine oocytes or embryos exported to the Republic of Moldova were collected from donor female(s) that (1):

(2) ○ either [were kept in a bluetongue virus-free country or zone for at least 60 days prior to collection of the oocytes or embryos;]

(2) ○ or [were kept during a bluetongue virus seasonally free period in a seasonally free zone for at least 60 days prior to collection of the oocytes or embryos;]

(2) ○ or [were kept in a vector protected establishment for at least 60 days prior to collection of the oocytes or embryos;]

(2) ○ or [were subjected to a serological test to detect antibodies to the bluetongue virus group, carried out in accordance with the WOAH Manual of Diagnostic Tests and Vaccines for Terrestrial Animals, with negative results between 21 and 60 days after the collection for this consignment of the oocytes or embryos;]

(2) ○ or [were subject to an agent identification test for bluetongue virus, carried out in accordance with the WOAH Manual of Diagnostic Tests and Vaccines for Terrestrial Animals with negative results on blood samples taken on the day of collection for this consignment of the oocytes or embryos.]

3. Donor females spent the 60 days immediately prior to the collection of bovine oocytes or embryos within the territory of the exporting country and without contact with imported cloven-hoofed animals for the last 30 days.

4. Bovine semen used for the insemination of the donor females or fertilisation of the oocytes was collected in accordance with Commission Delegated Regulation (EU) 2020/686.

5. As regards bovine tuberculosis, brucellosis and bluetongue(3) at the time of the collection/production of the bovine oocytes or embryos:

(2) ☐ either [the establishment of origin is free from tuberculosis in accordance with Commission Delegated Regulation (EU) 2020/689 and donor females have not been tested;]

(2) ☐ and/or [the establishment of origin is free from brucellosis in accordance with Commission Delegated Regulation (EU) 2020/689 and donor females have not been tested;]

(2) ☐ and/or [the country or zone of origin is free from bluetongue in accordance with Commission Delegated Regulation (EU) 2020/689 and is listed in Part I of Annex VIII to Commission Implementing Regulation (EU) 2021/620 and donor females have not been tested;]

(2) ☐ and/or [within 30 days prior to the collection/ production of oocytes or embryos, donor females have been tested, as appropriate, in a state approved laboratory by the methods recommended by WOAH Manual of Diagnostic Tests and Vaccines for Terrestrial Animals, with negative results (indicate the name of the laboratory, if relevant, and date and the testing method) for:

i. ☐ (2) [bovine tuberculosis: ;];

ii. ☐ (2) [brucellosis: ;];

iii. ☐ (1)(2) [bluetongue - testing by serological test or agent identification test according to WOAH Manual of Diagnostic Tests and Vaccines for Terrestrial Animals (to indicate the date of investigation):]

6. The oocytes or embryos have been collected, stored and transported in accordance with Commission Delegated Regulation (EU) 2020/686.

Notes:

Part I:

Part II: Certification

II. Health information

Box I.19: Either seal- or container number or both are to be indicated in this box.

Box I.25: Indicate total gross weight and total net weight.

Box I.28: “CN code”: use the appropriate Harmonized System (HS) code of the World Customs Organisation: 0511 99

Part II:

(1) May be deleted if the oocytes or embryos are in-vivo derived and processed in accordance with Manual of the International Embryo Transfer Society (IETS).

(2) Delete as appropriate.

(3) One of the options for freedom or testing must be kept for each disease, respectively.

Signature and stamp must be different colour that in the printed certificate.

Certifying Officer

Name (in capital letters)	Qualification and title
Date of signature	Signature
Stamp	