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.24. Total quantity

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	Breed/Category	Identification number	Date of collection/production	Quantity

Part II: Certification

II. Health information

I, the undersigned official veterinarian, certify the following:

II.1. (insert EU Member State of origin or a region thereof)

is officially free from Rift Valley fever, contagious caprine pleuropneumonia, peste des petits ruminants.

II.2. The embryo collection or production team

II.2.1. is authorised and supervised in accordance with Annex D, Chapter I(III) to Directive 92/65/EEC, and meets the requirement laid down by the Terrestrial Animal Health Code of OIE.

II.3. The donor females:

II.3.1. come from holdings free from brucellosis (B. melitensis) and scrapie in accordance with the procedure laid down by the OIE Terrestrial Animal Health Code;

(2)either ○ [II.3.2. come from a country(2) or a zone(2) of origin free from bluetongue;]

(2)or ○ [II.3.2. come from holdings under surveillance and/or control programmes for bluetongue;]

II.3.3. come from holdings that have not been subject to health restrictions for infectious-contagious diseases notifiable for these species during the last 24 months.

II.4. The semen used in the artificial insemination:

II.4.1. was collected, processed and stored in accordance with the provisions of the applicable OIE recommendations;

II.4.2. in the case the semen comes from a donor that has died or which health condition regarding reproductive or infectious diseases was unknown at the moment of collection of semen, the donor females should be subject to further examination after the collection of embryos, to check that there would not have been transmitted infectious diseases (B. melitensis, scrapie and bluetongue);

II.4.3. in case of natural service or use of fresh semen, donors must satisfy the health conditions described in the applicable OIE recommendations.

II.5. The ova/embryos:

II.5.1. were collected from donor females which showed no clinical signs of disease and were controlled by a team veterinarian authorised by the Competent Authority;

II.5.2. were collected and processed by the embryo collection or production team which has not carried out activities in an infected area or in an area subject to restrictions for animal health reasons affecting these species;

II.5.3. were collected and processed in accordance with the recommendations of the Manual of the International Embryo Transfer Society (IETS).

II.6. The storage and transport:

II.6.1. the ova/embryos were stored in safe premises approved for that purpose by the Competent Authority, under the supervision of the team veterinarian, who is responsible for the performance of all operations and attestations referred to in these health requirements;

II.6.2. while the ova/embryos for export to Chile are handled and until after storage, ova/embryos with a different health status have not been processed;

II.6.3. have been stored only in sterilised vials with fresh nitrogen not used for any other purpose.

Notes

Part I:

· Box I.11.: Place of origin: shall correspond to the embryo collection or production teams approved for export to Chile, from which the embryos are dispatched.

· Box I.20.: Number of packages: shall correspond to the number of containers.

· Box I.21.: Identification of container and seal number shall be indicated.

· Box I.25.: Customs code and title: Use the appropriate Harmonized System (HS) code under the following heading: 0511.

Species: select amongst “Ovis aries” or “Capra hircus” as appropriate.

Donor identity: shall correspond to the official identification of the animal (eartag number).

Part II: Certification

II. Health information

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

Approval number of the team shall correspond to the approval number of the embryo collection or production team indicated in Box I.11.

Part II:

(1) Only embryo collection and production teams, approved as being in compliance with Article 11(3) of Directive 92/65/EEC, and listed in accordance with Article 11(4) of that Directive on the Commission website:

http://ec.europa.eu/food/animal/approved_establishments/establishments_vet_field_en.htm and listed in the Register of Authorized establishments for export to Chile:

http://www.sag.gob.cl/opendocs/asp/pagDefault.asp?boton=Doc51&argInstanciaId=51&argCarpetaId=1394&argTreeNodosAbiertos=(1394)(-51)&argTreeNodoActual=1394&argTreeNodoSel=8

(2) Delete as necessary.

- The signature and the stamp must be in a different colour to that of the printing.
- The certificate must be issued in Spanish and in the language of the EU Member State of origin.

Certifying Officer

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