United Kingdom

Health certificate for the import into the UK of ovine and caprine ova embryos - Part A from EU Member States

Part I: Details of dispatched consignment

1.1. Consignor
Name
Address
Country

1.2. Certificate reference number

1.2.a.Local reference number::

1.3. Central Competent Authority

1.2b Unique notification number:

1.4. Local Competent Authority

1.5. Consignee
Name
Address
Country

1.6. No.(s) of related original certificates

No.(s) of accompanying documents

1.7. Dealer
Name
Approval number

1.8.Country of origin 1SO code

1.9. Region of origin Code

1.10. Country of destination 1SO code

1.11. Region of destination Code

1.12. Place of origin/Place of harvest
Holding ☐ Assembly centre ☐ Dealer s premise ☐
Approved body ☐ Semen centre ☐ Approved aquaculture holding ☐
Embryo team ☐ Establishment ☐ Other ☐
Name
Approval number
Address
Postal code / Region

1.13. Place of destination
Holding ☐ Assembly centre ☐ Dealer s premise ☐
Approved body ☐ Semen centre ☐ Approved aquaculture holding ☐
Embryo team ☐ Establishment ☐ Other ☐
Name
Approval number
Address
Postal code / Region

1.14. Place of loading
Postal code / Region

1.15. Date and time of departure

1.16. Means of transport
Aeroplane ☐ Ship ☐ Railway wagon ☐
Road vehicle ☐ Other ☐
1dentification::
Number(s):

1.17. Transporter
Name
Approval number
Address
Postal code / Region
Member state

1.21 Temperature of products
Ambient ☐ Chilled ☐ Frozen ☐

1.20. Number/Quantity

1.22. Number of packages

1.23. 1dentification of container/Seal number

1.25. Animals certified for/products certified for:

1.26. Transit through 3rd country ☐
Exit point Code
Entry point B1P unit no.:

1.27. Transit through Member states ☐

1.28. Export ☐
3rd country 1SO code
Exit point Code

1.29. Estimated journey time

1.30. Route plan
Yes ☐ No ☐

1.31. 1dentification of the animals

Part II: Certification

11. Health information	11.a. Certificate reference number	11.b. Local reference number:

1, the undersigned official veterinarian, hereby certify that:

(1)either [11.1. the in vivo derived embryos(1) / in vivo derived ova(1) described above were collected, processed and stored by an embryo collection team(2) approved and supervised in accordance with Chapter 1(111)(1) of Annex D to Directive 92/65/EEC;]

(1)or [11.1. the in vitro produced embryos(1) / micromanipulated embryos(1) described above were produced, processed and stored by an embryo production team(2) approved and supervised in accordance with Chapter 1(111)(1) and (2) of Annex D to Directive 92/65/EEC;]

(1)either [11.2. the in vivo derived embryos described above meet the requirements of Chapter 111(11)(1) of Annex D to Directive 92/65/EEC;]

(1)or [11.2. the in vivo derived ova described above meet the requirements of Chapter 111(11)(2) of Annex D to Directive 92/65/EEC;]

(1)or [11.2. the in vitro produced embryos described above meet the requirements of Chapter 111(11)(3) of Annex D to Directive 92/65/EEC;]

(1)or [11.2. the micromanipulated embryos described above meet the requirements of Chapter 111(11)(4) of Annex D to Directive 92/65/EEC;]

(1) [11.3. the consignment consists of embryos of the ovine or caprine species which:

(1)either [were collected from animals which have been kept continuously since birth on a holding or holdings recognised as having a negligible or a controlled risk of classical scrapie in accordance with point 1 of Section A of Chapter A of Annex V111 to Regulation (EC) No 999/2001;]

(1)or [were collected from animals which have been kept continuously for the last three years before the collection on a holding or holdings which have complied for the last three years before collection with the requirements laid down in points (a) to (f) of point 1.3. of Section A of Chapter A of Annex V111 to Regulation (EC) No 999/2001;]

(1)or [were collected from animals which have been kept continuously since birth in a Member State or zone of a Member State with a negligible risk status for classical scrapie approved in accordance with the first subparagraph of point 2.2. of Section A of Chapter A of Annex V111 to Regulation (EC) No 999/2001;]

(1)or [were collected from ovine animals and

(1)either [are of the ARR/ARR prion protein genotype;]

(1)or [carry at least one ARR allele and were collected after the date of 1 January 2015;]]]

11.4. the ova or embryos described above come from female donors of the ovine(1) / caprine species(1) which meet the requirements of Chapter 1V(3) of Annex D to Directive 92/65/EEC;

(1)either [11.5. the embryos described above were conceived as a result of artificial insemination of the donor females with semen which was collected, processed, stored and transported under conditions which comply with the requirements of Chapters 1(1), 11(1) and 111(1) of Annex D to Directive 92/65/EEC;]

(1)or [11.5. the embryos described above were conceived as a result of in vitro fertilisation of ova complying with the conditions in Chapter 111(11)(2) of Annex D to Directive 92/65/EEC with semen which was collected, processed, stored and transported under conditions which comply with the requirements of Chapters 1(1), 11(1) and 111(1) of Annex D to Directive 92/65/EEC;]

(1)or [11.5. the ova have not been in contact with semen of the ovine and caprine species;]

11.6. the ova or embryos described above were sent to the place of loading in a sealed container in accordance with point 6 of Chapter 111(11) of Annex D to Directive 92/65/EEC and bearing the number detailed in Box 1.23.

Notes

Part 1:

Box 1.12.: Place of origin shall correspond to the embryo collection team or embryo production team of embryos collection/production.

Box 1.13.: Place of destination shall correspond to the embryo collection team, embryo production team or to the holding of ova/embryos destination.

Box 1.23.: Identification of container and seal number shall be indicated.

Box 1.31.: Category: specify if: in vivo derived embryos, in vivo derived ova, in vitro produced embryos or micromanipulated embryos.

Donor identity shall correspond to the official identification of the animal.

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

Approval number of the team shall correspond to the embryo collection team or embryo production team of ova/embryos collection/production.

Part 11:

(1) Delete as appropriate.

(2) Only approved embryo collection or production teams listed in accordance with Article 11(4) of Directive 92/65/EEC on the Commission website:

http://ec.europa.eu/food/animal/approved_establishments/establishments_vet_field_en.htm.

· The colour of the stamp and signature must be different from that of the other particulars in the certificate.

Official veterinarian or official inspector

Name (in Capital): Qualification and title:

Local Veterinary Unit: LVU N°:

Date: Signature:

Stamp