EUROPEAN UNION **Intra trade certificate**

Part I : Details of dispatched consignment

I.1. Consignor
Name
Address
Country

I.2. Certificate reference number

I.2.a.Local reference number::

I.3. Central Competent Authority

I.4. Local Competent Authority

I.5. Consignee
Name
Address
Country

I.6. No.(s) of related original certificates

No.(s) of accompanying documents

I.7. Dealer
Name
Approval number

I.8.Country of origin ISO code

I.9. Region of origin Code

I.10. Country of destination ISO code

I.11. Region of destination Code

I.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

I.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

I.14. Place of loading
Postal code / Region

I.15. Date and time of departure

I.16. Means of transport
Aeroplane ☐ Ship ☐ Railway wagon ☐
Road vehicle ☐ Other ☐
Identification::
Number(s):

I.17. Transporter
Name
Approval number
Address
Postal code / Region Member state

I.21 Temperature of products
Ambient ☐ Chilled ☐ Frozen ☐

I.20. Number/Quantity

I.22. Number of packages

I.23. Identification of container/Seal number

I.25. Animals certified for/products certified for:

I.26. Transit through 3rd country ☐
Exit point Code
Entry point BIP unit no.:

I.27. Transit through Member states ☐

I.28. Export ☐
3rd country ISO code
Exit point Code

I.29. Estimated journey time

I.30. Route plan
Yes ☐ No ☐

I.31. Identification of the animals

Part II: Certification

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

I, the undersigned official veterinarian, hereby certify, that the animals described above meet the following requirements:

(1)either [II.1. The animals were born and have been reared since birth on Union territory.]

(1)or [II.1. The animals were imported from a third country complying with the animal health conditions laid down in Commission Regulation (EU) No 206/2010, at least 30 days prior to loading.]

II.2. The animals:

II.2.1. have been inspected today (within 24 hours prior to loading) and show no clinical sign of disease;

II.2.2. are not animals which are to be destroyed under a scheme to eradicate a contagious or infectious disease;

II.2.3. come from holdings which have been free from any official prohibition on health grounds, for the last 42 days in the case of brucellosis, for the last 30 days in the case of rabies, for the last 15 days in the case of anthrax, and, have not been in contact with animals from holdings which did not comply with these conditions;

II.2.4. do not come from a holding nor have been in contact with animals from a holding in a protection zone which has been set up under Union legislation and which animals are prohibited to leave;

II.2.5. are not the subject of animal health measures pursuant to Union legislation on foot-and-mouth disease nor have been vaccinated against foot-and-mouth disease.

II.3. Based on the written declaration made by the keeper or an examination of the holding register and movement documents kept in accordance with Council Regulation (EC) No 21/2004, in particular in Sections B and C of the Annex to that Regulation, the animals have remained on a single holding of origin for a period of at least the last 30 days, or on the holding of origin since birth where the animals are less than 30 days old, and no animal of the ovine and caprine species has been introduced into the holding of origin during the last 21 days and no biungulate animal imported from a third country has been introduced into the holding of origin during the last 30 days, unless those animals were introduced in accordance with Article 4a(1) of Council Directive 91/68/EEC.

(1) [II.4. The animals comply with the additional guarantees provided for in Articles 7 or 8 of Council Directive 91/68/EEC and laid down for the Member State of destination or part of its territory (insert Member State or part of its territory) in Commission Decision (insert number).]

II.5. The animals comply with at least one of the following conditions and therefore qualify for admission to an ovine or caprine holding which is officially brucellosis-free (B. melitensis):

(1) either [the holding of origin is situated in a Member State or part of its territory (insert name of Member State or part of its territory) which is recognised as being officially brucellosis-free in accordance with Commission Decision (insert number).]

(1) or [come from an officially brucellosis-free (B. melitensis) holding.]

(1) or [come from a brucellosis-free (B. melitensis) holding, and

(i) are identified individually,

(ii) have never been vaccinated against brucellosis or if the animals have been vaccinated were so vaccinated more than two years previously or the animals are females over two years old which were vaccinated before the age of seven months,

(iii) were isolated under official supervision on the holding of origin and, during such isolation, underwent, with negative results, two tests for brucellosis in accordance with Annex C to Directive 91/68/EEC, separated by an interval of at least six weeks.]

II.6. The animals comply with at least one of the following conditions and therefore qualify for admission to an ovine or caprine holding which is brucellosis-free (B. melitensis):

(1) either [come from an officially brucellosis-free (B. melitensis) holding.]

(1) or [come from a brucellosis-free (B. melitensis) holding.]

(1) or [until the qualifying date under eradication plans approved pursuant to Council Decision 90/242/EEC, the animals originate from a holding other than officially brucellosis-free or brucellosis-free and satisfy the following conditions:

(i) are identified individually,

(ii) originate from a holding in which all the animals of species susceptible to brucellosis (B. melitensis) have been free of clinical symptoms or any other symptoms of brucellosis for at least 12 months; and

(1) either [have not been vaccinated against brucellosis (B. melitensis) in the last two years, and were isolated under veterinary supervision on the holding of origin and, during such isolation, underwent, with negative results, two tests for brucellosis in accordance with Annex C to Directive 91/68/EEC, separated by an interval of at least six weeks.]

(1) or [were vaccinated with Rev. 1 vaccine before the age of seven months, and were not vaccinated in the 15 days before the date of emission of this health certificate.]]

(1) [II.7. The uncastrated breeding rams must:

(i) come from a holding on which no case of contagious epididymitis of rams (B. ovis) has been recorded in the past 12 months,

(ii) have been kept permanently on that holding for the 60 days preceding consignment,

(iii) have undergone, within 30 days prior to consignment, with a negative result, a test to detect contagious epididymitis of rams (B. ovis) in accordance with Annex D to Directive 91/68/EEC.]

II.8. To the best of the knowledge of the undersigned and according to the written declaration made by the owner, the animals were not obtained from a holding nor have been in contact with animals from a holding in which the following diseases have been clinically detected:

(i) within the last six months, contagious agalactia of sheep (Mycoplasma agalactiae) and contagious agalactia of goats (Mycoplasma agalactiae, M. capricolum, M. mycoides subsp. mycoides large colony),

(ii) within the last 12 months, paratuberculosis or caseous lymphadenitis,

(iii) within the last three years, pulmonary adenomatosis, maedi/visna or caprine viral arthritis/encephalitis. However, this time limit is reduced to 12 months if animals affected by maedi/visna or caprine viral arthritis/encephalitis have been slaughtered and the remaining animals have reacted negatively to two tests.

(1)either [II.9. The animals are intended for a Member State or zone of a Member State listed in point 2.3. of Section A of Chapter A of Annex VIII to Regulation (EC) No 999/2001 as having a negligible risk status for classical scrapie or for a Member State listed in point 3.2. of that Section as having an approved national scrapie control programme, and

(1)either [come from a holding situated in a Member State or zone of a Member State listed in point 2.3. of Section A of Chapter A of Annex VIII to Regulation (EC) No 999/2001 as having a negligible risk status for classical scrapie.]

(1)and/or [come from a holding recognised as having a negligible risk of classical scrapie in accordance with point 1.2. of Section A of Chapter A of Annex VIII to Regulation (EC) No 999/2001 and listed as such by the competent authority of the Member State in accordance with point 1.1. of that Section.]

(1)and/or [come from a holding not subject to the measures laid down in points 3 and 4 of Chapter B of Annex VII to Regulation (EC) No 999/2001 and the animals are of the ovine species of the ARR/ARR prion protein genotype.]

(1)and/or [come from and are destined for an approved body, institute or centre as defined in Article 2(1)(c) of Directive 92/65/EEC.]

(1)or [comply with the conditions set out in point 4.1.(d) of Section A of Chapter A of Annex VIII to Regulation (EC) No 999/2001.]]

(1)or [II.9. The animals are intended for a Member State or zone of a Member State other than those listed in point 2.3. of Section A of Chapter A of Annex VIII to Regulation (EC) No 999/2001 as having a negligible risk status for classical scrapie or other than those listed in point 3.2. of that Section as having an approved national scrapie control programme, and

(1)either [come from a holding situated in a Member State or zone of a Member State listed in point 2.3. of Section A of Chapter A of Annex VIII to Regulation (EC) No 999/2001 as having a negligible risk status for classical scrapie.]

Part II: Certification

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

(1)and/or [come from a holding recognised as having a negligible risk of classical scrapie in accordance with point 1.2. of Section A of Chapter A of Annex VIII to Regulation (EC) No 999/2001 and listed as such by the competent authority of the Member State in accordance with point 1.1. of that Section.]

(1)and/or [come from a holding not subject to the measures laid down in points 3 and 4 of Chapter B of Annex VII to Regulation (EC) No 999/2001 and the animals are of the ovine species of the ARR/ARR prion protein genotype.]

(1)and/or [come from a holding recognised as having a controlled risk of classical scrapie in accordance with point 1.3. of Section A of Chapter A of Annex VIII to Regulation (EC) No 999/2001 and listed as such by the competent authority of the Member State in accordance with point 1.1. of that Section.]

(1)and/or [come from and are destined for an approved body, institute or centre as defined in Article 2(1)(c) of Directive 92/65/EEC.]

(1)or [comply with the conditions set out in point 4.1.(d) of Section A of Chapter A of Annex VIII to Regulation (EC) No 999/2001.]]

II.10.1. The animals were transported using means of transport and containment which had, before-hand, been cleansed and disinfected using an officially approved disinfectant, and in such a way as to provide effective protection of the animals' health status.

II.10.2. Based on the official documentation accompanying the animals, the consignment covered by this health certificate are due to start the journey on (insert date)(2).

II.10.3. At the time of inspection the animals covered by this health certificate were fit to be transported on the intended journey in accordance with the provisions of Council Regulation (EC) No 1/2005(3).

Blue Tongue (BT): Exemption from the exit ban

- BT-1: Animals in compliance with Article 7(1) or 7(2)(a) or 7(2)(b) or 7(2)(c) or 7(2a)(a) or 7(2a)(b) or 7(2a)(c) , (indicate as appropriate) of Regulation (EC) No 1266/2007.
- BT-2: Animals , semen , ova and embryos , (indicate as appropriate) in compliance with Article 8(1)(a) or 8(1)(b) or 8(4) or 8(5a) ,(indicate as appropriate) of Regulation (EC) No1266/2007.
- BT-3: Insecticide/repellent treatment with (insert name of the product) on (insert date) in conformity with Regulation (EC) No 1266/2007.
- BT-4: Animal(s) in compliance with Article 9a(1) of Regulation (EC) No 1266/2007.

- BTA1: Animal(s) were kept until dispatch in a bluetongue seasonally-free zone during the seasonally vector-free period that started on (insert date) since birth or for at least 60 days and, if appropriate (indicate as appropriate), were then subjected to an agent identification test according to the OIE Terrestrial Manual on samples taken within seven days prior to dispatch, with negative results , in conformity with Annex III.A(1) to Regulation (EC) No 1266/2007.
- BTA2: Animal(s) in conformity with Annex III.A(2) to Regulation (EC) No 1266/2007.
- BTA3: Animal(s) in conformity with Annex III.A(3) to Regulation (EC) No 1266/2007.
- BTA4: Animal(s) in conformity with Annex III.A(4) to Regulation (EC) No 1266/2007.
- BTA5: Animal(s) vaccinated against bluetongue serotype/s (insert serotype/s) with (insert name of the vaccine) with a inactivated / modified live vaccine (indicate, as appropriate) in conformity with Annex III.A(5) to Regulation (EC) No 1266/2007.
- BTA6: Animal(s) subjected to a serological test according to the OIE Terrestrial Manual to detect antibodies against the bluetongue virus serotype (indicate serotype) in conformity with Annex III.A(6) to Regulation (EC) No 1266/2007.
- BTA7: Animal(s) subjected to a specific serological test according to the OIE Terrestrial Manual to detect antibodies against all the bluetongue virus serotypes (indicate serotypes) present or likely to be present in conformity with Annex III.A(7) to Regulation (EC) No 1266/2007.
- BTA8: "Animal(s) is (are) not pregnant" or "Animal(s) may be pregnant and complies (comply) with the condition(s) (set out in points 5 , 6 and 7 before insemination or mating, or set out in point 3 ; indicate as appropriate)".

Notes

Part I:

- Box reference I.19.: Use the appropriate CN code under the following headings: 01.04.10 or 01.04.20.
- Box reference I.23.: For containers or boxes, the container number and the seal number (if applicable) should be included.
- Box reference I.31.: Identification system: The animals must bear: An individual number which permits tracing of their premises of origin, according to Council Regulation (EC) No 21/2004.

Age: (months).

Sex: (M = male, F = female, C = castrated)

Part II:

(1) Delete where not applicable.

(2) In the case where a consignment is grouped in an assembly centre and comprises animals that were loaded on different dates, the date at which the journey commenced for the whole consignment is considered to be the earliest date when any part of the consignment left the holding of origin.

(3) This statement does not exempt transporters from their obligations in accordance with Union rules in force in particular regarding the fitness of animals to be transported.

· This certificate is valid for 10 days

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

Part II: Certification

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

Official veterinarian or official inspector

Name (in Capital):

Qualification and title:

Local Veterinary Unit:

LVU N°:

Date:

Signature:

Stamp

Part III: Control

III.1. Date of the inspection

III.2. Certificate Reference Number::

III.3. Documentary Check:: No Yes

EU Standard Satisfactory Not satisfactory

Additional guarantees Satisfactory Not satisfactory

National requirements Satisfactory Not satisfactory

III.4. Identity Check:: No Yes

Satisfactory Not satisfactory

III.5. Physical Check:: No Total animals checked

Satisfactory Not satisfactory

III.6. Laboratory Tests:: No Yes

Date:

Tested for::

Random Suspicion

Results:: Pending Satisfactory Not satisfactory

III.7. Welfare check No Yes

Satisfactory Not satisfactory

III.8.Infringement of welfare regulation::

III.8.1.Transporter authorisation invalid

III.8.2.Non-compliance of the means of transport

III.8.3.Stocking density exceeded Average space

III.8.4.Travel times exceeded

III.8.5.Watering and feeding not fulfilled

III.8.6.Mishandling or negligence to the animals

III.8.7.Supplementary measures for the journeys of long duration

III.8.8.Certificate of proficiency of the driver

III.8.9.Data registered in the log book

III.8.10.Other

III.9. Infringement of health legislation

III.9.1. Absence/Invalid certificate

III.9.2. Mis-match with documents

III.9.3. Non authorised country

III.9.4. Non approved region/ zone

III.9.5. Prohibited species

III.9.6. Absence of additional guarantee

III.9.7. Non approved holding

III.9.8. Diseased or suspect animals

III.9.9. Unsatisfactory tests

III.9.10. Absence or non legal identification

III.9.11.National requirements not fulfilled

III.9.12. Address of destination invalid

III.9.13. Other

III.10. Impact of the transport on animals

Number of dead animals:: Estimation:

Number of unfit animals :: Estimation:

Number of birth or abortion:

III.11. Corrective action

III.11.1. Delayed departure

III.11.2. Transfer procedure

III.11.3. Quarantine

III.11.4. Humane killing/Euthanasia

III.11.5. Destruction of carcasses/products

III.II.6. Return of consignment

III.11.7. Treatment of products

III.11.8.7. Use of products for other purpose

Identification:

III.12. Follow-up of quarantine

III.12.1.Humanely killing/Euthanasia

III.12.2.Release

III.13. Place of inspection

Establishment Holding Assembly centre

Dealer's premise Approved body Semen centre

Port Airport Exit point

Enroute Other

III.14. Official veterinarian or official inspector

Local Veterinary Unit LVU N°

Name (in Capital):

Qualification and title

Date: Signature:

PLANNING

1.1. ORGANISER name and address (a) (b)	1.2. Name of the person in charge of the journey
	1.3. Telephone / Fax

2. TOTAL EXPECTED DURATION (hours / days)	
3.1. Place and country of DEPARTURE	4.1. Place and country of DESTINATION

3.2. Date	3.3. Time	4.2. Date	4.3. Time

5.1. Species	5.2. Number of animals	5.3. Veterinary certificate(s) number(s)

5.4. Estimated total weight of the consignment (in kg)	5.5. Total space foreseen for the consignment (in m²)

6. LIST OF FORESEEN RESTING, TRANSFER OR EXIT POINTS

6.1. Name of the places where animals are to be rested, or transferred (including exit points)	6.2. Arrival		6.3. Length (in hours)	6.4. Transporter name and authorisation N° (if different from the organiser)	6.5 identification
	Date	Time			

7. I, the organiser, hereby declare that I am responsible for the organisation of the above-mentioned journey and I have made suitable arrangements to safeguard the welfare of the animals throughout the journey in accordance with the provisions of Council Regulation 1/2005
8. Signature of the organiser

(a) Organiser: see definition laid down in Article 2(q) of Council Regulation 1/2005

(b) If the organiser is a transporter the authorisation number shall be specified