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(1) / veterinarian responsible for the establishment of origin and approved by the competent authority(1) certify that:

II.1. the animals described in Box I.31 comply with the conditions of Article 4 of Council Directive 92/65/EEC and at the time of inspection were fit to be transported for the intended journey in accordance with the provisions of Council Regulation (EC) No 1/2005.

(1)either [II.2. ruminant(s) (1)/ suidae (1) other than that /those covered by Council Directive 64/432/EEC (1) or Council Directive 91/68/EEC (1):

(a) belong(s) to the species ;

(b) at the time of examination, do(does) not show any clinical sign of any disease to which it/they is /are susceptible;

(c) come(s) from an officially tuberculosis-free (1)/ officially brucellosis-free (1) or brucellosis-free (1) herd (1)/ holding (1) not subject to swine fever restrictions or from a holding where it/they was/were subjected with negative results to the tests laid down in Article 6(2)(b) (1)/ the test laid down in Article 6(3)(d) (1) of Council Directive 92/65/EEC.]

(1)(2)or [II.2. the birds other than those referred to in Council Directive 2009/158/EC

(a) at the time of examination do not show any clinical sign of any disease to which they are susceptible;

(b) satisfy the requirements of Article 7 of Council Directive 92/65/EEC;

(c) conform to Commission Decision 2007/598/EC and were vaccinated against avian influenza on (date) with vaccine (name) and come from a holding on which vaccination against avian influenza was carried out during the past 12 months.]

(1)or [II.2. the lagomorphs

(a) at the time of examination do not show any clinical signs of disease to which they are susceptible;

(b) satisfy the requirements of Article 9 of Council Directive 92/65/EEC.]

(1)or [II.2. the dogs

(a) at the time of examination by a veterinarian authorised by the competent authority within 48 hours prior to the time of dispatch, showed no signs of diseases;

(b) are marked in accordance with Article 17(1) of Regulation (EU) No 576/2013 of the European Parliament and of the Council;

(1)either [(c) were at least 12 weeks old at the time of vaccination against rabies and at least 21 days have elapsed since the completion of the primary anti-rabies vaccination carried out in accordance with the validity requirements set out in Annex III to Regulation (EU) No 576/2013 of the European Parliament and of the Council, and any subsequent revaccination was carried out within the period of validity of the preceding vaccination]] ;

(1)or [(c) are less than 12 weeks old and have not received an anti-rabies vaccination, or are between 12 and 16 weeks old and have received an anti-rabies vaccination, but 21 days at least have not elapsed since the completion of the primary vaccination against rabies carried out in accordance with the validity requirements set out in Annex III to Regulation (EU) No 576/2013 of the European Parliament and of the Council], and

(i) the Member State of destination has informed the public in accordance with point (b) of Article 37(2) of Regulation (EU) No 576/2013 of the European Parliament and of the Council that it authorises the movement of such animals into its territory; and they are accompanied by

(1)either [(ii) a declaration of the owner(3), attached to this certificate, stating that from birth until the time of dispatch the animals have had no contact with wild animals of species susceptible to rabies] ;

(1)or [(ii) their mother, on whom they still depend, and from the passport of their mother, it can be established that the mother received before their birth an anti-rabies vaccination which complied with the validity requirements set out in Annex III to Regulation (EU) No 576/2013 of the European Parliament and of the Council] ;] ;

(d) are accompanied by a passport drawn up in accordance with Commission Implementing Regulation (EU) No 577/2013;

(1)and [(e) due to their scheduled destination indicated in Box I.10 or in Box I.11 where regionalisation is applied, have been treated against Echinococcus multilocularis in accordance with Commission Delegated Regulation (EU) No 1152/2011]] ;

(1)or [II.2. the cats (1)/ ferrets (1)

(a) at the time of examination by a veterinarian authorised by the competent authority within 48 hours prior to the time of dispatch, showed no signs of diseases;

(b) are marked in accordance with Article 17(1) of Regulation (EU) No 576/2013 of the European Parliament and of the Council;

(1)either [(c) were at least 12 weeks old at the time of vaccination against rabies and at least 21 days have elapsed since the completion of the primary anti-rabies vaccination carried out in accordance with the validity requirements set out in Annex III to Regulation (EU) No 576/2013 of the European Parliament and of the Council, and any subsequent revaccination was carried out within the period of validity of the preceding vaccination] ;

(1)or [(c) are less than 12 weeks old and have not received an anti-rabies vaccination, or are between 12 and 16 weeks old and have received an anti-rabies vaccination, but 21 days at least have not elapsed since the completion of the primary vaccination against rabies carried out in accordance with the validity requirements set out in Annex III to Regulation (EU) No 576/2013 of the European Parliament and of the Council], and

(i) the Member State of destination has informed the public in accordance with point (b) of Article 37(2) of Regulation (EU) No 576/2013 of the European Parliament and of the Council that it authorises the movement of such animals into its territory; and they are accompanied by

(1)either [(ii) a declaration of the owner(3), attached to this certificate, stating that from birth until the time of dispatch the animals have had no contact with wild animals of species susceptible to rabies] ;

(1)or [(ii) their mother, on whom they still depend, and from the passport of their mother, it can be established that the mother received before their birth an anti-rabies vaccination which complied with the validity requirements set out in Annex III to Regulation (EU) No 576/2013 of the European Parliament and of the Council] ;]

(d) are accompanied by a passport drawn up in accordance with Commission Implementing Regulation (EU) No No 577/2013.]

(1)or [II.2. the dogs (1)/ cats (1)/ ferrets (1) are destined for a body, institute or centre described in Box I.13 and approved in accordance with Annex C to Council Directive 92/65/EEC, and

(a) at the time of examination by a veterinarian authorised by the competent authority within 48 hours prior to the time of dispatch, showed no signs of diseases;

(b) are marked in accordance with Article 17(1) of Regulation (EU) No 576/2013 of the European Parliament and of the Council;

(c) are accompanied by a passport drawn up in accordance with Commission Implementing Regulation (EU) No 577/2013.]

II.3. The additional guarantees regarding diseases listed in Annex B(4) to Council Directive 92/65/EEC are as follows:(1)

Disease Decision

Disease Decision

Disease Decision

Blue Tongue (BT) : exemption from the exit ban

Part II: Certification

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

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.

Animals 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.

Insecticide/repellent treatment with (insert name of the product) on (insert date) in conformity with Regulation (EC) No 1266/2007.

Animal(s) in compliance with Article 9a(1) of Regulation (EC) No 1266/2007

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.

Animal(s) in conformity with Annex III.A(2) to Regulation (EC) No 1266/2007

Animal(s) in conformity with Annex III.A(3) to Regulation (EC) No 1266/2007

Animal(s) in conformity with Annex III.A(4) to Regulation (EC) No 1266/2007

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.

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.

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.

"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 I.6.: No(s) of accompanying documents: CITES, if applicable.

Box I.19.: Use the appropriate CN code: 01.06.19, 01.06.31, 01.06.32, 01.06.39.

Box I.31.: Identification system: individual identification must be used wherever possible but in the case of small animals, batch identification may be used. In the case of dogs, cats and ferrets, select passport.
Identification number: in the case of dogs, cats and ferrets, indicate the alphanumeric code of the tattoo or transponder.
Passport number: in the case of dogs, cats and ferrets, indicate the unique alphanumeric code of the passport.

Part II:

(1) Delete as necessary.

(2) Certification requirements only apply to birds that have been vaccinated against avian influenza under a preventive vaccination plan approved by Commission Decision 2007/598/EC.

(3) The declaration referred to in point II.2 to be attached to the certificate shall be drawn up in accordance with the model set out in Annex I to Commission Implementing Regulation ((EU) No 577/2013.

(4) As requested by a Member State benefiting from additional guarantees under Union legislation.

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

This certificate is valid for 10 days from the date of signature of the official veterinarian or of the veterinarian responsible for the holding of origin and approved by the competent authority.

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	Control post

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