

Veterinary certificate to EU

Part I : Details of consignment presented

I.1. Consignor Name Address Postal code / Region Country	I.2. Certificate reference number	I.2.a. TRACES reference number :
	I.3. Central Competent Authority	
	I.4. Local Competent Authority	
	I.6 Person responsible for load in EU	
I.5. Consignee Name Address Postal code / Region Country		I.6 Person responsible for load in EU
I.7. Country of origin, ISO code	I.8. Region of origin, Code	I.9. Country of destination ISO code I.10. Region of destination Code
I.11. Place of origin Holding <input type="checkbox"/> Semen centre <input type="checkbox"/> Establishment <input type="checkbox"/> Name Approval number Address Name Approval number Address Name Approval number Address		I.12. Place of destination
I.13 Place of loading Address Approval number		I.14 Date of departure
I.15. Means of transport Aeroplane <input type="checkbox"/> Ship <input type="checkbox"/> Railway wagon <input type="checkbox"/> Road vehicle <input type="checkbox"/> Other <input type="checkbox"/> Identification:: Document:		I.16. Entry BIP in EU Name BIP unit no.: I.17. No.(s) of CITES
I.18. Description of commodity		I.19. Commodity code (HS code)
I.21 Temperature of products		I.20. Quantity I.22. Number of packages
I.23. Identification of container/Seal number		I.24. Type of packaging
I.25. Commodity certified as: Breeding <input type="checkbox"/> Fattening <input type="checkbox"/> Slaughter <input type="checkbox"/> Approved bodies <input type="checkbox"/> Artificial reproduction <input type="checkbox"/> Quarantine <input type="checkbox"/> Game restocking <input type="checkbox"/> Registered horses <input type="checkbox"/> Pets <input type="checkbox"/> Circus <input type="checkbox"/> Relaying <input type="checkbox"/> Other <input type="checkbox"/> Humane consumption <input type="checkbox"/> Animal feedingstuff <input type="checkbox"/> Further process <input type="checkbox"/> Pharmaceutical use <input type="checkbox"/> Technical use <input type="checkbox"/>		
I.26. For transit to 3rd Country by EU		I.27. For import or admission into EU Definitive import <input type="checkbox"/> <input type="checkbox"/> Horses Re-entry <input type="checkbox"/> <input type="checkbox"/> Temporary admission horses <input type="checkbox"/> <input type="checkbox"/>
I.28. Identification of the commodity Species (scientific name) Breed/category Identification system Identification number Age Sex Quantity Test Age(dd/mm/yyyy) Age(Weeks) Age(Months)		

Part II: Certification

II. Health information	II.a. Certificat reference number	II.b. TRACES reference number
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1. VACCINATION AGAINST RABIES

Manufacturer and name of vaccine:

Batch No:

Vaccination date (3):

Valid until (3):

2. RABIES SEROLOGICAL TEST (when required – strike out when not certified)

I have seen the official record of the result of a serological test for the animal, carried out on a sample taken on (3), and tested in an EU-approved laboratory, which states that the rabies neutralizing antibody titre was equal to or greater than 0.5 IU/ml.

3. CLINICAL EXAMINATION

I declare that the animal is at present free of clinical signs and transportable.

4. TICK TREATMENT (when required – strike out when not certified)

Manufacturer and name of product:

Date (3) and time of treatment (24-hour clock):

5. ECHINOCOCCUS TREATMENT (when required – strike out when not certified)

Manufacturer and name of product:

Date (3) and time of treatment (24-hour clock):

NOTES FOR GUIDANCE

1. Identification of the animal (tattoo or microchip) must be verified before any entries are made on the certificate.
2. The rabies vaccine used must be an inactivated vaccine produced in accordance with OIE standards.
3. The certificate shall be valid for 4 months from the date of signature by the approved or official veterinarian or until the date of expiry of the vaccination shown in Part IV, whichever is earlier.
4. Animals from, or prepared in, third countries not listed in Annex II to Regulation (EC) No 998/2003, may not enter Ireland, Sweden or the United Kingdom, either directly or via another country listed in Annex II unless brought into conformity with National Rules.
5. The clinical examination (Part IX) must be done within 24 hours before movement.
6. Parts not certified must be struck out.

APPLICABLE CONDITIONS (Regulation (EC) No 998/2003)

A. ENTRY IN A MEMBER STATE OTHER THAN IRELAND, SWEDEN AND UNITED KINGDOM

- A1. from a third country listed in Annex II to Regulation (EC) No 998/2003: Parts I to VII and IX must be completed (and XI for Finland).
- A2. from a third country not listed in Annex II to Regulation (EC) No 998/2003: Parts I to IX must be completed (and XI for Finland). The sample referred to in Part VIII must have been taken more than 3 months before the date of entry.

B. ENTRY IN IRELAND, SWEDEN AND UNITED KINGDOM

- B1. from a third country listed in Annex II to Regulation (EC) No 998/2003: Parts I to XI must be completed (Parts VI, VIII, X and XI complying with national rules)
- B2. from a third country not listed in Annex II to Regulation (EC) No 998/2003: The certificate is not valid - See Note 4

- (1) Add ISO code
- (2) Delete as applicable
- (3) dd/mm/yyyy

Official veterinarian or official inspector

Name (in Capital):

Local Veterinary Unit:

Date:

Stamp

Qualification and title:

LVU N°:

Signature: