CHAPTER 4 (D)

Health certificate

For treated blood products, excluding of equidae, for the manufacture of technical products, intended for dispatch to or for transit through (2) the European Community

COUNTRY

Veterinary certificate to EU

Part I: Details of dispatched consignment

I.1. Consignor Name Address Tel. No		I.2. Certificate reference number	I.2.a.
		I.3. Central Competent Authority	
		I.4. Local Competent Authority	
I.5. Consignee Name Address Postal code Tel. No		I.6. Person responsible for the consignment in EU Name Address Postal code Tel. No	
I.7. Country of origin ISO code	I.8. Region of origin Code	I.9. Country of destination ISO code	I.10.
I.11. Place of origin Name Approval number Address		I.12. Place of destination Custom warehouse ☐ Name Approval number Address Postal code	
I.13. Place of loading		I.14. Date of departure	
I.15. Means of transport Aeroplane ☐ Ship ☐ Railway wagon ☐ Road vehicle ☐ Other ☐ Identification: Documentary references:		I.16. Entry BIP in EU I.17.	
I.18. Description of commodity		I.19. Commodity code (HS code) **30.02**	
			I.20. Quantity
I.21. Temperature of product Ambient ☐ Chilled ☐ Frozen ☐			I.22. Number of packages
I.23. Identification of container/Seal number			I.24. Type of packaging
I.25. Commodities certified for: Technical use ☐			
I.26. For transit to 3rd Country vis-à-vis EU ☐ third country ISO code		I.27. For import or admission into EU ☐	

I.28. Identification of the commodities

Species (Scientific name)	Nature of commodity	Approval number of establishments Manufacturing plant	Batch number

Part II: Certification

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

II.	**Health attestation**
	I, the undersigned official veterinarian, declare that I have read and understood Regulation (EC) No 1774/2002 ([1]) and in particular Article 4(1)(c), Article 6 and Chapter IV of Annex VIII thereof and certify that:
II.1.	the blood products described above consist of blood products that satisfy the requirements below;
II.2.	they consist exclusively of blood products not intended for human or animal consumption;
II.3.	they have been prepared and stored in a plant approved, validated and supervised by the competent authority in accordance with Article 18 or in the establishment of collection and where appropriate Article 11 of Regulation (EC) No 1774/2002 ([2]), exclusively with the following animal by-products:
([2]) *either*	[— blood of slaughtered animals, which is fit for human consumption in accordance with Community legislation, but is not intended for human consumption for commercial reasons;]
([2]) *and/or*	[— blood of slaughtered animals, which is rejected as unfit for human consumption but is not affected by any signs of diseases communicable to humans or animals, derived from carcasses that are fit for human consumption in accordance with Community legislation;]
([2]) *and/or*	[— blood obtained from animals other than ruminants that are slaughtered in a slaughterhouse, after undergoing ante-mortem inspection, and were fit, as a result of such inspection, for slaughter for human consumption in accordance with Community legislation;]
([2]) *and/or*	[— blood and blood products originating from live animals that did not show clinical signs of any disease communicable through these products to humans or animals.]
II.4.	the blood from which such products are manufactured has been collected:
([2]) *either*	[in slaughterhouses approved in accordance with Community legislation,]
([2]) *or*	[in slaughterhouses approved and supervised by the competent authority of the third country,]
([2]) *or*	[from live animals in facilities approved and supervised by the competent authority of the third country.]
([2]) II.5.	In case of blood products derived from *taxa Artiodactyla, Perissodactyla* and *Proboscidea* including their crossbreeds, other than *Suidae* and *Tayassuidae,* the products have undergone one of the following treatments, guaranteeing the absence of pathogens of foot-and-mouth disease, vesicular stomatitis, rinderpest, *peste des petits* ruminants, Rift Valley fever and blue-tongue:
([2]) *either*	[heat treatment at a temperature of 65 °C for at least three hours, followed by an effectiveness check,]
([2]) *or*	[irradiation at 25 kGy by gamma rays, followed by an effectiveness check,]
([2]) *or*	[change in pH to pH 5 for two hours, followed by an effectiveness check,]
([2]) *or*	[heat treatment of at least 80 °C throughout their substance, followed by an effectiveness check]
([2]) II.6.	In the case of blood products derived from *Suidae, Tayassuidae,* poultry and other avian species, the products have undergone one of the following treatments guaranteeing the absence of pathogens of the following diseases: foot-and-mouth disease, vesicular stomatitis, swine vesicular disease, classical swine fever, African swine fever, Newcastle disease and highly pathogenic avian influenza as appropriate to the species;
([2]) *either*	[heat treatment at a temperature of 65 °C for at least three hours, followed by an effectiveness check,]
([2]) *or*	[irradiation at 25 kGy by gamma rays, followed by an effectiveness check,]
([2]) *or*	[heat treatment of at least 80 °C for *Suidae/Tayassuidae* ([2]) and at least 70 °C for poultry and other avian species ([2]) throughout their substance, followed by an effectiveness check]
([2]) [II.7.	In the case of blood products derived from species other than listed under II.5 or II.6 the products have undergone of the following treatment (please specify): …]
II.8.	The products were:
([2]) *either*	[packed in new or sterilised bags or bottles,]
([2]) *or*	[transported in bulk in containers or other means of transport that were thoroughly cleaned and disinfected with a disinfectant approved by the competent authority before use,]
	the outer packaging or containers bear labels indicating "NOT FOR HUMAN OR ANIMAL CONSUMPTION";

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

II.9. the products were stored in enclosed storage;

II.10. the products have undergone all precautions to avoid contamination with pathogenic agents after treatment.

Notes

Part I:

— Box reference I.6: Person responsible for the consignment in the European Community: this box is to be filled in only if it is a certificate for transit commodity; it may be filled in if the certificate is for import commodity.

— Box reference I.12: Place of destination: this box is to be filled in only if it is a certificate for transit commodity. The products in transit can only be stored in free zones, free warehouses and custom warehouses.

— Box reference I.15: Registration number (railway wagons or container and lorries), flight number (aircraft) or name (ship); information is to be provided in case of unloading and reloading.

— Box reference I.23: for bulk containers, the container number and the seal number (if applicable) should be included.

— Box reference I.26 and I.27: fill in according to whether it is a transit or an import certificate.

Part II:

([1]) OJ L 273, 10.10.2002, p. 1.

([2]) Delete as appropriate.

— The signature and the stamp must be in a different colour to that of the printing.

— Note for the person responsible for the consignment in the European Community: this certificate is only for veterinary purposes and has to accompany the consignment until it reaches the border inspection post.

Official veterinarian

Name (in capital letters): Qualification and title:

Date: Signature:

Stamp:'