

CHAPTER 4(C)

Health certificate

For untreated blood products, excluding of equidae, for the manufacture of derived products for purposes outside the feed chain for farmed animals, intended for ~~dispatch to or~~ for transit through⁽²⁾ the European Union

COUNTRY: UNITED STATES

Veterinary certificate to EU

Part I : Details of dispatched consignment	I.1. Consignor Name Address		I.2. Certificate reference No		I.2.a.	
	Tel.		I.3. Central competent authority APHIS-VS			
			I.4. Local competent authority			
	I.5. Consignee Border inspection post through which consignment is intended to leave the EU Name Address Postal code Tel.			I.6. Person responsible for the load in EU Name Address Postal code Tel.		
	I.7. Country of origin US	I.8. ISO code US-0	I.8. Region of origin	I.9. Country of destination	I.10. ISO code	I.10. Region of destination
	I.11. Place of origin Name Approval number Address Name Approval number Address			I.12. Place of destination Name Address Postal code Custom warehouse <input type="checkbox"/> Approval number		
	I.13. Place of loading			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 Documentation references			I.16. Entry BIP in EU		
				I.17.		
	I.18. Description of commodity			I.19. Commodity code (HS code)		
			I.20. Quantity			
I.21. Temperature of product Ambient <input type="checkbox"/> Chilled <input type="checkbox"/> Frozen <input type="checkbox"/>			I.22. Number of packages			

COUNTRY: UNITED STATES

Untreated blood products, excluding of equidae, for the manufacture of derived products for purposes outside the feed chain for farmed animals

Part II: Certification

II. Health information	II.a. Certificate reference No	II.b.
<p>I, the undersigned official veterinarian, declare that I have read and understood Regulation (EC) No 1069/2009 of the European Parliament and of the Council^(1a), and in particular Article 8(c) and Article 8(d) and Article 10 thereof, and Commission Regulation (EU) No 142/2011^(1b), and in particular Annex XIV, Chapter II thereof, and certify that:</p> <p>II.1. the blood products described above consist of blood products that satisfy the health requirements below;</p> <p>II.2. they consist exclusively of blood products not intended for human or animal consumption;</p> <p>II.3. they have been prepared and stored in a plant supervised by the competent authority or in the establishment of collection, exclusively with the following animal by-products:</p> <p>⁽²⁾ <i>either</i> [- blood of slaughtered animals, which is fit for human consumption in accordance with Union legislation, but is not intended for human consumption for commercial reasons;]</p> <p>⁽²⁾ <i>and/or</i> [- blood of slaughtered animals, which is rejected as unfit for human consumption in accordance with Union legislation, but which did not show any signs of diseases communicable to humans or animals, derived from carcasses that have been slaughtered in a slaughterhouse and were considered fit for human consumption following an ante-mortem inspection in accordance with Union legislation;]</p> <p>⁽²⁾ <i>and/or</i> [- blood of slaughtered animals, which did not show any signs of diseases communicable to humans or animals, obtained from animals that have been slaughtered in a slaughterhouse after having been considered fit for human consumption following an ante-mortem inspection in accordance with Union legislation;]</p> <p>⁽²⁾ <i>and/or</i> [- blood and blood products derived from the production of products intended for human consumption;]</p> <p>⁽²⁾ <i>and/or</i> [- blood and blood products originating from live animals that did not show signs of any disease communicable through that product to humans or animals;]</p> <p>⁽²⁾ <i>and/or</i> [- animal by-products derived from animals which have been submitted to illegal treatment as defined in Article 1(2)(d) of Directive 96/22/EC or Article 2(b) of Directive 96/23/EC;]</p> <p>⁽²⁾ <i>and/or</i> [- animal by-products containing residues of other substances and environmental contaminants listed in Group B(3) of Annex I to Directive 96/23/EC, if such residues exceed the permitted level laid down in Union legislation or, in the absence thereof, in national legislation;]</p> <p>II.4. the blood from which such products are manufactured has been collected in slaughterhouses approved in accordance with Union legislation, in slaughterhouses approved and supervised by the competent authority of the country of collection or from live animals in facilities approved and supervised by the competent authority of the country of collection.</p> <p>⁽²⁾II.5. in the case of blood products derived from animals belonging to the taxa Artiodactyla, Perissodactyla and Proboscidea, including their crossbreds, the products come:</p> <p>II.5.1. from a country where no case of rinderpest, peste des petits ruminants and Rift Valley fever has been recorded for 12 months and in which vaccination has not been carried out against those diseases for at least 12 months;</p> <p>⁽²⁾II.5.2. <i>either</i> [from the third countries, territories or parts thereof (ISO code in case of country or codes for territories or parts thereof)⁽³⁾ where no case of foot-and-mouth disease has been recorded for 12 months and in which vaccination has not been carried out against this disease for at least 12 months;]</p> <p><i>or</i> [from the countries, territories or parts thereof (ISO code in case of country or codes for territories or parts thereof)⁽³⁾ where no case of foot-and-mouth disease has been recorded for 12 months and in which vaccination programmes against foot-and-mouth disease are being officially carried out and controlled in domestic ruminant animals for at least 12 months⁽⁴⁾.;]</p> <p>⁽²⁾II.5.3. In addition, in case of animals other than Suidae and Tayassuidae:</p> <p>⁽²⁾<i>either</i> [in the country or region of origin no case of vesicular stomatitis and bluetongue⁽²⁾ (including the presence of seropositive animals) has been recorded for 12 months and in which vaccination has not been carried out against those diseases for at least 12 months;]</p> <p>⁽²⁾<i>or</i> [in the country or region of origin vesicular stomatitis and bluetongue⁽²⁾ seropositive animals are present⁽⁴⁾.;]</p> <p>⁽²⁾II.5.4. In addition, in case of Suidae and Tayassuidae:</p> <p>II.5.4.1. in the country or region of origin no case of swine vesicular disease, classical swine fever and African swine fever has been recorded for at least 12 months and vaccination has not been carried out against those diseases for at least 12 months in the susceptible species and</p> <p>⁽²⁾II.5.4.2. <i>either</i> [in the country or region of origin no case of vesicular stomatitis (including the presence of seropositive animals) has been recorded for 12 months and in which vaccination has not been carried out against this disease for at least 12 months;]</p> <p><i>or</i> [in the country or region of origin vesicular stomatitis seropositive animals are present⁽⁴⁾.;]</p> <p>⁽²⁾II.6. in the case of blood products derived from poultry or other avian species the animals and the products come from the territory of the country or region with code⁽⁵⁾ which has been free from Newcastle disease and highly pathogenic avian influenza as defined in the Terrestrial Animal Health Code of the OIE, which for at least 12 months has not carried out vaccination against avian influenza, where the animals from which the products derive have not been vaccinated against Newcastle disease with vaccines</p>		

