

CHAPTER 4(B)

Health certificate

For blood products not intended for human consumption that could be used as feed material, intended for dispatch to or for transit through² 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 Name Address Postal code Tel.		I.6. Person responsible for the load in EU Name Address Postal code Tel.			
	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 Name Approval number Address		I.12. Place of destination Name Address Postal code Custom warehouse <input type="checkbox"/> Approval number			
	Name Approval number Address					
	Name Approval number Address					
	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"/>		I.16. Entry BIP in EU				
Identification		I.17.				
Documentation references						
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

Blood products not intended for human consumption that could be used as feed material

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 Commission Regulation (EU) No 142/2011^(1b) and certify that the blood products described above:</p>			
II.1. consist of blood products that satisfy the health requirements below;				
II.2. consist exclusively of blood products not intended for human consumption;				
II.3. have been prepared and stored in a plant, approved, validated and supervised by the competent authority in accordance with Article 24 of Regulation (EC) No 1069/2009;				
II.4. have been prepared exclusively with the following animal by-products:				
<p>⁽²⁾either [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⁾and/or [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>				
II.5. have been submitted				
<p>⁽²⁾either [to processing in accordance with processing method⁽³⁾ as set out in Chapter III of Annex IV to Regulation (EU) No 142/2011]</p>				
<p>⁽²⁾or [to a method and parameters which ensure that the product complies with the microbiological standards set in Chapter I of Annex X to Regulation (EU) No 142/2011,] in order to kill pathogenic agents;</p>				
II.6. have been examined under the responsibility of the competent authority taking a random sample immediately prior to dispatch and found it to comply with the following standards ⁽⁴⁾ :				
<p>Salmonella: absence in 25g: n = 5, c = 0, m = 0, M = 0, Enterobacteriaceae: n = 5, c = 2, m = 10, M = 300 in 1 gram;</p>				
II.7. the end product was:				
<p>⁽²⁾either [packed in new or sterilised bags;]</p>				
<p>⁽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,] and which bear labels indicating 'NOT FOR HUMAN CONSUMPTION';</p>				
II.8. the end product was stored in enclosed storage;				
II.9. the product has undergone all precautions to avoid contamination with pathogenic agents after treatment;				
I.10.				
<p>⁽²⁾either [the product does not contain and is not derived from specified risk material as defined in Annex V to Regulation (EC) No 999/2001 of the European Parliament and of the Council⁽⁵⁾ or mechanically separated meat obtained from bones of bovine, ovine or caprine animals; and the animals from which this product is derived have not been slaughtered after stunning by</p>				

