

Part I : Details of consignment	I.1. Consignor			I.2. IMSOC Reference		
	Name			I.2.a. Local Reference		
	Address					
	Country			ISO Code		
	I.5. Consignee			I.3. Central competent authority		
	Name			I.4. Local competent authority		
	Address					
	Country			ISO Code		
	I.7. Country of origin			I.9. Country of destination		
	ISO Code			ISO Code		
	I.8. Region of origin			<del>I.10. Region of destination</del>		
	Code					
	I.11. Place of Dispatch			I.12. Place of destination		
	Name			Name		
Address			Address			
Approval Number			Approval Number			
Country			Country			
ISO Code			ISO Code			
I.13. Place of Loading			I.14. Date and time of departure			
Name						
Address						
Approval Number						
Country						
ISO Code						
I.15. Means of Transport			I.16 Entry Point			
Mode	International transport document	Identification				
I.18. Transport conditions			I.17. Accompanying documents			
Ambient <input type="checkbox"/> Frozen <input type="checkbox"/> Controlled temperature <input type="checkbox"/> Chilled <input type="checkbox"/>			Commercial document reference			
			Date of issue			
			Country			
			Place of issue			
I.19. Container No / Seal No						
I.20. Certified as						
Animal Feedingstuff <input type="checkbox"/> Other <input type="checkbox"/>						
I.21. For transit through a third country <input type="checkbox"/>			I.22. For transit through Member State(s) <input type="checkbox"/>			
Country			Country			
ISO Code			ISO Code			
EU Exit Authority			BCP code			
EU Entry Authority			BCP code			
I.23. Total number of packages		I.25. Total net weight		I.25. Total gross weight		
I.28. Description of consignment						
<b>1. 05 PRODUCTS OF ANIMAL ORIGIN, NOT ELSEWHERE SPECIFIED OR INCLUDED</b>						
<b>0511</b> Animal products not elsewhere specified or included; dead animals of Chapter  1  or 3, unfit for human consumption						
<b>051199</b> Other						
Commodity	Manufacturing plant	Product Description	Species	Date of production range		
Package count			Net weight			

II. Health information		
<b>Part II: Certification</b>	I, the undersigned official veterinarian, after due inquiry and to the best of my knowledge, do hereby certify that the processed porcine blood described within this certificate meets all the conditions laid down in the health attestations below:	
	II.1.	The country or zone of origin is free of foot-and-mouth disease, classical swine fever, swine vesicular disease, and African swine fever. Vaccination against these diseases is prohibited in that country.
	II.2.	The products were only produced and exported to Canada by the producer and exporter identified in the Import Permit and mentioned on this certificate's first page.
	II.3.	The certified rendered products were produced:
	(2)	○ either [II.3.1. In a dedicated facility that does not receive, process or store any ruminants and things derived from ruminants, and the product has been prepared, processed, packaged, stored, shipped and otherwise handled in a manner to avoid contamination with any ruminant tissues or things derived from ruminants,]
	(2)	○ or [II.3.1. on a dedicated line from receipt of raw material to final packaging and storage with no risk of cross contamination with any ruminants and things derived from ruminants, and the product has been prepared, processed, packaged, stored, shipped and otherwise handled in a manner to avoid contamination with any ruminant tissues or things derived from ruminants (if the facility has ineligible materials on the premises, all rendered products for export to Canada must be produced on a totally dedicated line.)]
	II.4.	The blood used in the certified products was obtained only from carcasses that passed ante-mortem inspection and were subjected to post-mortem inspection in slaughterhouses approved and supervised by the competent authority.
	II.5.	None of the animals from which any of the unprocessed blood used to manufacture the rendered products are derived from premises that were under any official restrictions by the Member State competent veterinary authority for any serious epizootic disease to which the species from which the product or by-product was derived is susceptible and can be transmitted by the untreated product or by-product and none of the animals from which the animal origin raw materials are derived were under movement restrictions for or were culled or eradicated as part of a disease response for any reportable disease as defined by Canada (1).
	II.6.	The raw materials used to produce the rendered products have been transported in dedicated vehicles.
	II.7.	The porcine blood products have been processed in an establishment approved by the Canadian Food Inspection Agency for export to Canada.
	II.8.	The blood products are exclusively of porcine origin.
	II.9.	The blood products have been heat treated to at least 80° Celsius throughout their substance.
	II.10.	The blood products have a moisture content of less than 10%.
	II.11.	Each shipment to Canada has been tested negative for the presence of ruminant DNA with a PCR method, in an accredited laboratory.
	II.12.	The porcine blood products have been packed using new and secure packaging materials.
II.13.	The raw materials and additives used in the porcine blood products have been clearly indicated on the exterior packaging.	
II.14.	The porcine blood products for export have been produced, processed, stored and transported in such a manner as to prevent contamination by communicable animal disease pathogens up to the point of departure from the country of origin.	
II.15.	The product label bears the following statement "The product does NOT contain prohibited material, as defined by section 162 of the Canadian Health of Animals Regulations <sup>2</sup> ."	
Notes		
	All pages must be presented at least in English and/or French as well as at least one of the official languages of the exporting Member State of the EU. The official stamp and the signature of the official veterinarian must appear on each separate sheet, including any attached lists. The signature and the stamp must be in a different colour to that of the printing.	

II. Health information			
<b>Part II: Certification</b>	Part I:		
	Box reference I.1:	Indicate the details of the exporter.	
	Box reference I.2.:	Indicate the unique reference number.	
	Box reference I.2.a:	In case this certificate is produced via TRACES system, a unique reference number assigned by the TRACES system is indicated.	
	Box reference I.5.:	Indicate the details of the importer.	
	Box reference I.6.:	Indicate the CFIA Import permit number	
	Box reference I.11.:	Place of origin: name and address of the dispatch establishment.	
	Box reference I.15.:	Indicate the names of the ships and, if known, the flight numbers of aircraft.	
	Box reference I.19.:	Indicate total gross weight and total net weight.	
	Box reference I.21.:	For containers or boxes, the container number and the seal number should be included.	
	Box reference I.22.:	Commodities certified for must identify the end use.	
	Box reference I.25.:	Customs code and title: Use the appropriate Harmonized System (HS) code under the following headings: 0511, 3002, 3502, 2301	
		Processing plant: Indicate the establishment approval number.	
		Description of product: as per Import permit	
		Animal species: only porcine blood allowed	
	Date of production: Shall be indicated in the following format: dd/mm/yyyy		
Part II:			
(1)	The CFIA list of diseases that are reportable in Canada can be found on the CFIA website: Animal Health Status By Disease - Animals - Canadian Food Inspection Agency. The CFIA accepts the OIE classification of BSE Risk Status: List of BSE risk status: OIE - World Organisation for Animal Health.		
(2)	Where the product is destined for use in livestock feed, it is approved for use and listed in Schedule IV of the Feeds Regulations and is labelled as required by the Feed Regulations, which have been captured here: <a href="https://ec.europa.eu/food/safety/international_affairs/agreements/export-library_en">https://ec.europa.eu/food/safety/international_affairs/agreements/export-library_en</a>		
Certifying Officer			
Name (in capital letters)		Qualification and title	
Date of signature		Signature	
Stamp			