EUROPEAN UNION **Export Health Certificate**

Part I : Details of consignment

I.1. Consignor
Name
Address
Country ISO Code

I.2. IMSOC Reference
I.2.a. Local Reference

I.5. Consignee
Name
Address
Country ISO Code

I.3. Central competent authority
I.4. Local competent authority

I.7. Country of origin ISO Code

I.9. Country of destination ISO Code

I.8. Region of origin Code

~~I.10. Region of destination~~

I.11. Place of Dispatch
Name
Address
Approval Number
Country ISO Code

I.12. Place of destination
Name
Address
Approval Number
Country ISO Code

I.13. Place of Loading
Name
Address
Approval Number
Country ISO Code

I.14. Date and time of departure

I.15. Means of Transport

Mode	International transport document	Identification

I.16 Entry Point

I.18. Transport conditions
Frozen ☐ Chilled ☐ Ambient ☐ Controlled temperature ☐

I.17. Accompanying documents
Commercial document reference Date of issue
Country Place of issue

I.19. Container No / Seal No

I.20. Certified as
Other ☐ Animal Feedingstuff ☐

I.21. For transit through a third country ☐
Country ISO Code
EU Exit Authority BCP code
EU Entry Authority BCP code

I.22. For transit through Member State(s) ☐
Country ISO Code

I.23. Total number of packages	I.25. Total net weight	I.25. Total gross weight

I.28. Description of consignment

1. 23 RESIDUES AND WASTE FROM THE FOOD INDUSTRIES; PREPARED ANIMAL FODDER

2301 Flours, meals and pellets, of meat or meat offal, of fish or of crustaceans, molluscs or other aquatic invertebrates, unfit for human consumption; greaves

Commodity	Manufacturing plant	Product Description	Species	Date of production range

Package count	Net weight

Part II: Certification

II. Health information

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 80º 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[2].

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.

Part II: Certification

II. Health information

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: https://ec.europa.eu/food/safety/international_affairs/agreements/export-library_en

Certifying Officer

Name (in capital letters)	Qualification and title
Date of signature	Signature
Stamp	