COUNTRY

Official certificate to the EU

Part I : Details of consignment

I.1. Consignor
Name
Address
Country ISO Code

I.2. Certificate reference

I.2.a. IMSOC reference

I.3. Central competent authority

I.4. Local competent authority

I.5. Consignee
Name
Address
Country ISO Code

I.6. Responsible for the consignment in EU
Name
Address
Country ISO Code

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 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 BCP
Authority
Country

I.18. Transport conditions
Ambient ☐ **Frozen** ☐ **Chilled** ☐

I.17. Accompanying documents
Type
Number

I.19. Container No / Seal No

I.20. Certified as
Slaughter ☐ **Trade samples** ☐ **Animal Feedingstuff** ☐ **Other** ☐ **Pets** ☐ **Artificial reproduction** ☐ **Canning industry** ☐ **Technical Use** ☐ **Approved Bodies** ☐ **Human Consumption** ☐ **Fattening** ☐ **Quarantine** ☐ **Pharmaceutical use** ☐ **Further process** ☐ **Registered equidae** ☐ **Relaying** ☐ **Breeding/production** ☐ **Circus/exhibition** ☐ **Game restocking** ☐

I.21. For transit ☐
Non-EU ISO Code

I.22. For internal market ☐
Definitive import ☐ **Horses re-entry** ☐ **Temporary admission horses** ☐

I.23. Total number of packages

I.24. Total net weight

I.24. Total gross weight

I.25. Description of consignment

1. 35 ALBUMINOIDAL SUBSTANCES; MODIFIED STARCHES; GLUES; ENZYMES

3504 Peptones and their derivatives; other protein substances and their derivatives, not elsewhere specified or included; hide powder, whether or not chromed

350400 Peptones and their derivatives; other protein substances and their derivatives, not elsewhere specified or included; hide powder, whether or not chromed

Commodity	Manufacturing plant	Package count	Net weight	Batch number
Nature of commodity				

COUNTRY

		tricalcium phosphate not intended for human consumption to be used as feed material or for uses outside the feed chain
Part II : Certification	II. Health information	

Part II : Certification

II. Health information

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 10 thereof, and Commission Regulation (EU) No 142/2011 (1b), and in particular Chapter I of Annex XIV thereto, and certify that the ○ hydrolysed protein/ ○ dicalcium phosphate/ ○ tricalcium phosphate (2) described above:

II.1. consists of ○ hydrolysed protein/ ○ dicalcium phosphate/ ○ tricalcium phosphate (2) that satisfy the health requirements below;

II.2. consists exclusively of ○ hydrolysed protein/ ○ dicalcium phosphate/ ○ tricalcium phosphate (2) not intended for human consumption;

II.3. has been prepared and stored in a plant approved and supervised by the competent authority in accordance with Article 24 of Regulation (EC) No 1069/2009, in order to kill pathogenic agents;

II.4. has been prepared exclusively with the following animal by-products:

(2) ○ either [in the case of dicalcium phosphate derived from defatted bones, carcases and parts of animals slaughtered or, in the case of game, bodies or parts of animals killed, and which are fit for human consumption in accordance with Union legislation, but are not intended for human consumption for commercial reasons;]

(2) ○ or [in the case of other materials:

(2) ☐ either [- carcases and parts of animals slaughtered or, in the case of game, bodies or parts of animals killed, and which are fit for human consumption in accordance with Union legislation, but are not intended for human consumption for commercial reasons;]

(2) ☐ and/or [- carcases and the following parts originating either from animals that have been slaughtered in a slaughterhouse and were considered fit for slaughter for human consumption following an ante-mortem inspection or bodies and the following parts of animals from game killed for human consumption in accordance with Union legislation:

(i) carcases or bodies and parts of animals which are rejected as unfit for human consumption in accordance with Union legislation, but which did not show any signs of disease communicable to humans or animals;

(ii) heads of poultry;

(iii) hides and skins, including trimmings and splitting thereof, horns and feet, including the phalanges and the carpus and metacarpus bones, tarsus and metatarsus bones;

(iv) pig bristles;

(v) feathers;]

(2) ☐ and/or [- blood of animals which did not show any signs of disease communicable through blood to humans or animals obtained from animals that have been slaughtered in a slaughterhouse after having been considered fit for slaughter for human consumption following an ante-mortem inspection in accordance with Union legislation;]

(2) ☐ and/or [- animal by-products arising from the production of products intended for human consumption, including degreased bone, greaves and centrifuge or separator sludge from milk processing;]

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		(2)	☐ and/or	[-		products of animal origin, or foodstuffs containing products of animal origin, which are no longer intended for human consumption for commercial reasons or due to problems of manufacturing or packaging defects or other defects from which no risk to public or animal health arise;]
		(2)	☐ and/or	[-		petfood and feedingstuffs of animal origin, or feedingstuffs containing animal by-products or derived products, which are no longer intended for feeding for commercial reasons or due to problems of manufacturing or packaging defects or other defects from which no risk to public or animal health arises;]
		(2)	☐ and/or	[-		blood, placenta, wool, feathers, hair, horns, hoof cuts and raw milk originating from live animals that did not show signs of any disease communicable through that product to humans or animals;]
		(2)	☐ and/or	[-		aquatic animals, and parts of such animals, except sea mammals, which did not show any signs of diseases communicable to humans or animals;]
		(2)	☐ and/or	[-		animal by-products from aquatic animals originating from plants or establishments manufacturing products for human consumption;]
		(2)	☐ and/or	[-		the following material originating from animals which did not show any signs of disease communicable through that material to humans or animals:
					(i)	shells from shellfish with soft tissue or flesh;
					(ii)	the following originating from terrestrial animals:
						- hatchery by-products,
						- eggs,
						- egg by-products, including egg shells;
					(iii)	day-old chicks killed for commercial reasons;]]
II.5.	the ○ hydrolysed protein/ ○ dicalcium phosphate/ ○ tricalcium phosphate (2):					
				(a)		was wrapped and packaged in packaging which bear labels indicating ‘NOT FOR HUMAN CONSUMPTION’ and was stored and transported under satisfactory hygiene conditions, and in particular the wrapping and packaging took place in a dedicated room, and only preservatives permitted under Union legislation were used; and
		(2)	○ either	[(b)		in the case of hydrolysed protein, was produced by a process involving appropriate measures to minimise contamination of raw Category 3 material. In the case of hydrolysed proteins entirely or partly derived from ruminants hides and skins, was produced in a processing plant dedicated only to hydrolysed proteins production, using a process involving the preparation of the raw Category 3 material by brining, liming and intensive washing followed by:
					(i)	the exposure of the material to a pH of more than 11 for more than 3 hours at a temperature of more than 80 °C and subsequently by heat treatment at a temperature of more than 140 °C for 30 minutes at more than 3,6 bar ; or
					(ii)	the exposure of the material to a pH of 1 to 2, followed by a pH of more than 11, followed by a heat treatment at a temperature of more than 140 °C for 30 minutes at 3 bar.]
		(2)	○ or	[(b)		in the case of dicalcium phosphate, was produced by a process that:

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(i) ensures that all Category 3 bonematerial is finely crushed and degreased with hot water and treated with dilute hydrochloric acid (at a minimum concentration of 4 % and a pH of less than 1,5) over a period of at least two days,

(ii) followed by a treatment of the obtained phosphoric liquor with lime, resulting in a precipitate of dicalcium phosphate at pH 4 to 7, and

(iii) finally air-dries this precipitate, with an inlet temperature of 65 °C to 325 °C and an end temperature of between 30 °C and 65 °C.]

(2) ○ or [(b) in the case of tricalcium phosphate, was produced by a process ensuring:

(i) that all Category 3 bone-material is finely crushed and degreased in counter-flow with hot water (bone chips less than 14 mm),

(ii) the continuous cooking with steam at 145 °C during 30 minutes at 4 bars,

(iii) the separation of the protein broth from the hydroxyapatite (tricalcium phosphate) by centrifugation, and

(iv) the granulation of the tricalcium phosphate after drying in a fluidised bed with air at 200 °C.]

(2) ☐ [II.6. the ○ hydrolysed protein/ ○ dicalcium phosphate/ ○ tricalcium phosphate (2) described above

(2) ○ either [is derived from other ruminants than bovine, ovine or caprine animals.]

(2) ○ or [is derived from bovine, ovine or caprine animals and does not contain and is not derived from:

(2) ○ either [bovine, ovine and caprine materials other than those derived from animals born, continuously reared and slaughtered in a country or region classified as posing a negligible BSE risk in accordance with Decision 2007/453/EC.]

(2) ○ or [(a) specified risk material as defined in point 1 of Annex V to Regulation (EC) No 999/2001 of the European Parliament and of the Council (3);

(b) mechanically separated meat obtained from bones of bovine, ovine or caprine animals, except from those animals that were born, continuously reared and slaughtered in a country or region classified as posing a negligible BSE risk in accordance with Commission Decision 2007/453/EC (4), in which there has been no indigenous BSE case,

(c) animal by-product or derived product obtained from bovine, ovine or caprine animals which have been killed, after stunning, by laceration of the central nervous tissue by means of an elongated rod-shaped instrument introduced into the cranial cavity, or by means of gas injected into the cranial cavity, except for those animals that were born, continuously reared and slaughtered in a country or region classified as posing a negligible BSE risk in accordance with Decision 2007/453/EC.]]]

II.7. the ○ hydrolysed protein/ ○ dicalcium phosphate/ ○ tricalcium phosphate (2) described above:

(2) ○ either [does not contain milk or milk products of ovine or caprine animal origin or is not intended for feed for farmed animals, other than fur animals.]

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(2) ○ or [contains milk or milk products of ovine or caprine animal origin and is intended for feed for farmed animals, other than fur animals, and the milk or milk products:

(a) are derived from ovine and caprine animals which have been kept continuously since birth in a country where the following conditions are fulfilled:

(i) classical scrapie is compulsorily notifiable;

(ii) an awareness, surveillance and monitoring system is in place for classical scrapie;

(iii) official restrictions apply to holdings of ovine or caprine animals in the case of a suspicion of TSE or the confirmation of classical scrapie;

(iv) ovine and caprine animals affected with classical scrapie are killed and destroyed;

(v) the feeding to ovine and caprine animals of meat-and-bone meal or greaves, as defined in the Terrestrial Animal Health Code of the World Organisation for Animal Health (OIE), of ruminant origin has been banned and effectively enforced in the whole country for a period of at least the preceding seven years;

(b) originate from holdings where no official restrictions are imposed due to a suspicion of TSE;

(c) originate from holdings where no case of classical scrapie has been diagnosed during the period of the preceding seven years or, following the confirmation of a case of classical scrapie:

(2) ○ either [all ovine and caprine animals on the holding have been killed and destroyed or slaughtered, except for breeding rams of the ARR/ARR genotype, breeding ewes carrying at least one ARR allele and no VRQ allele and other ovine animals carrying at least one ARR allele;]

(2) ○ or [all animals in which classical scrapie was confirmed have been killed and destroyed, and the holding has been subjected for a period of at least two years since the date of confirmation of the last classical scrapie case to intensified TSE monitoring, including testing with negative results for the presence of TSE in accordance with the laboratory methods set out in point 3.2 of Chapter C of Annex X to Regulation (EC) No 999/2001, of all of the following animals which are over the age of 18 months, except ovine animals of the ARR/ARR genotype:

– animals which have been slaughtered for human consumption; and

– animals which have died or been killed on the holding but which were not killed in the framework of a disease eradication campaign.]]

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II. Health information

Notes

Part I:

- Box reference I.6: Person responsible for the consignment in the European Union: this box is required to be filled in only if it is a certificate for a commodity to be transited through the European Union; it may be filled in if the certificate is for a commodity to be imported into the European Union.
- Box reference I.12: Place of destination: this box is to be filled in only if it is a certificate for a transit commodity. 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.19: use the appropriate HS code: 05.08, 28.35.25; 28.35.26, 29.22; 35.02; 35.03 or 35.04.
- Box reference I.23: for bulk containers, the container number and the seal number (if applicable) must be included.
- Box reference I.25: technical use: any use other than feeding of farmed animals, other than fur animals, and the production or manufacturing of pet food.
- Box reference I.26 and I.27: fill in according to whether it is a transit or an import certificate.
- Box reference I.28:
  - Species: select from the following: Aves, Ruminantia, Suidae, Mammalia other than Ruminantia or Suidae, Pesca, Mollusca, Crustacea, invertebrates other than Mollusca and Crustacea.
  - Nature of commodity: specify if hydrolysed protein, dicalcium phosphate or tricalcium phosphate.
  - Manufacturing plant: provide the registration number of treatment/processing establishment.

Part II:

(1a) OJ L 300, 14.11.2009, p. 1.

(1b) OJ L 54, 26.2.2011, p. 1.

(2) Delete as appropriate.

(3) OJ L 147, 31.5.2001, p. 1.

(4) OJ L 94, 1.4.2006, p. 28.

- 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 Union: this certificate is only for veterinary purposes and must accompany the consignment until it reaches the border inspection post of the point of entry into the European Union.

Official veterinarian or Official inspector

Name (in capital letters) | Qualification and title

Date of signature | Signature

Stamp