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
Ambient ☐ Controlled temperature ☐ Chilled ☐ Frozen ☐

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

I.19. Container No / Seal No

I.20. Certified as
Human consumption ☐

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. 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

#1.	Commodity	Date of manufacture	Manufacturing plant	Package count
Species		Net weight	Date of production	

Part II: Certification

II. Health information

II.1. Public health attestation

I, the undersigned official veterinarian, hereby certify that the collagen described in this certificate was produced in accordance with the following requirements:

II.1.1. It has been produced from raw materials (either ○ bones, hides and skins of farmed ruminant animals, ○ pig skins and ○ bones, tendons and sinews, ○ poultry skin, ○ fish skin and bones) (2) which met the following requirements:

– (a)(1) the raw materials are derived from animals which have been slaughtered in a slaughterhouse and have been found fit for human consumption following ante-mortem and post-mortem inspection;

– (b) the raw materials originated from establishments under the supervision of the competent veterinary authorities.

– (c) hides and skins used have not undergone any tanning process;

☐ and(2)

– If from ruminant origin:

(2)either ○ [(d) the collagen was prepared exclusively from hides and skins];

(2)or ○ [(d) the raw materials derived from animals born, continuously reared and slaughtered in countries recognised by the OIE as of negligible BSE risk];

(2)or ○ [(d) the raw materials are derived from animals which have not been slaughtered after stunning by means of compressed air or gas injected into the cranial cavity or a pithing process; such raw materials neither contained nor were contaminated by: tonsils or distal ileum of cattle of any age; brains, eyes, spinal cord and skull from cattle aged over 12 months and vertebral column(*)from cattle over 30 months of age.]

II.1.2. It has been manufactured:

– (a) in an establishment under the supervision of the competent veterinary authorities and approved for the manufacture of collagen;

– (b) by a process to ensure that the raw material is subjected to a treatment involving washing, pH adjustment using acid or alkali followed by one or more rinses, filtration and extrusion or by an approved equivalent process. The extrusion step may not be carried out when manufacturing low molecular collagen from raw materials of non-ruminant origin;

– (c) after having been subjected to the process referred to in point(b), collagen underwent a drying process.

II.1.3. After the manufacture, the product was stored under appropriate hygiene conditions and the necessary measures were taken to avoid contamination.

II.1.4. The product meets the following microbiological limits:

Micro organisms	Limits
Salmonella	Absence in 25g

II.1.5. The packages were appropriately wrapped and labelled.

II.1.6. The label bears the words "collagen fit for human consumption" and indicates the country and establishment of origin and the date of preparation.

II.1.7. The container in which the product is dispatched complies with appropriate hygiene and sanitary requirements.

Notes

Part I:

· Box reference I.11: Place of origin: Name, address and approval number of the establishment of dispatch.

Part II: Certification

II. Health information		

· Box reference I.15: Flight number(aircraft)or name(ship). Separate information is to be provided in case of unloading and reloading.

· Box reference I.21: Identification of container/seal number: only where applicable.

· Box reference I.25: Treatment type: date of manufacture(dd/mm/yyyy).

Part II:

(1) Point II.1.1(a)is not applicable when fish skin and bones are used as raw materials.

(2) Delete as appropriate.

(*) Vertebral column excluding the vertebrae of the tail, the spinous and transverse processes of the cervical, thoracic and lumbar vertebrae and the median sacral crest and wings of the sacrum, but including the dorsal root ganglia, of animals aged over 30 months.

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