

Appendix II to Annex VI

PART A

MODEL HEALTH CERTIFICATE FOR IMPORTS OF GELATINE INTENDED FOR HUMAN CONSUMPTION

COUNTRY

Veterinary certificate to EU

Part I: Details of dispatched consignment	I.1. Consignor Name Address Postal code Tel No.		I.2. Certificate reference number	I.2.a.	
			I.3. Central Competent Authority		
			I.4. Local Competent Authority		
	I.5. Consignee Name Address Postal code Tel No.		I.6.		
	I.7. Country of origin	ISO code	I.8.		I.9. Country of destination
				ISO code	I.10.
	I.11. Place of origin Name Address		I.12. Approval number		
	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"/> Identification: Documentary references:		I.16. Entry BIP in EU		I.17.
	I.18. Description of commodity			I.19. Commodity code (HS code) 35.03	
			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		
I.23. Identification of container/Seal number			I.24. Type of packaging		
I.25. Commodities certified for Human consumption <input type="checkbox"/>					
I.26.		I.27. For import or admission into EU <input type="text"/>			
I.28. Identification of the commodities Species (Scientific name) Treatment type Approval number of establishments Manufacturing plant Number of packages Net weight					

COUNTRY

Gelatine intended for human consumption

Part II: Certification	II. Health attestation	II.a. Certificate reference number	II.b.
	<p>I, the undersigned, declare that I am aware of the relevant provisions of Regulations (EC) No 178/2002, (EC) No 852/2004 and (EC) No 853/2004 and certify that the gelatine described above was produced in accordance with those requirements, in particular that it:</p> <ul style="list-style-type: none"> — comes from (an) establishment(s) implementing a programme based on the HACCP principles in accordance with Regulation (EC) No 852/2004, — has been produced from raw material which met the requirements of Section XIV, Chapters I and II of Annex III to Regulation (EC) No 853/2004, — has been manufactured in compliance with the conditions set out in Section XIV, Chapter III of Annex III to Regulation (EC) No 853/2004, — satisfies the criteria of Section XIV, Chapter IV of Annex III to Regulation (EC) No 853/2004 and to Regulation (EC) No 2073/2005 on microbiological criteria for foodstuffs, and ⁽¹⁾ — if from ruminant origin, does not contain and is not derived from: <ul style="list-style-type: none"> either ⁽¹⁾ specified risk material as defined in Annex XI, section A, to Regulation (EC) No 999/2001 produced after 31 March 2001, or mechanically recovered meat obtained from bones of bovine, ovine or caprine animals produced after 31 March 2001. After 31 March 2001 the bovine, ovine and caprine animals, from which this product is derived, have not been slaughtered after stunning by means of gas injected into the cranial cavity or killed by the same method or slaughtered by laceration after stunning of central nervous tissue by means of an elongated rod-shaped instrument introduced into the cranial cavity. or bovine, ovine and caprine materials other than those derived from animals born, continuously reared and slaughtered in⁽²⁾ ⁽³⁾. 		
Notes			
Part I:			
— Box reference I.11: Place of origin: name and address of the dispatch establishment.			
— Box reference I.15: Registration number (railway wagons or container and lorries), flight number (aircraft) or name (ship). Separate information is to be provided in case of unloading and reloading.			
— Box reference I.23: Identification of container/seal number: only where applicable.			
— Box reference I.28: Treatment type: date of manufacture (dd/mm/yyyy).			
Part II:			
⁽¹⁾ Delete as appropriate.			
⁽²⁾ Insert the name of the country.			
⁽³⁾ As listed in point 15(b) of Annex XI to Regulation (EC) No 999/2001 as amended.			
— The colour of the stamp and signature must be different to that of the other particulars in the certificate.			
Official veterinarian			
Name (in capitals):		Qualification and title:	
Date:		Signature:	
Stamp:			