Commission Implementing Regulation (EU) 2016/759 of 28 April 2016 drawing up lists of third countries, parts of third countries and territories from which Member States are to authorise the introduction into the Union of certain products of animal origin intended for human consumption, laying down certificate requirements, amending Regulation (EC) No 2074/2005 and repealing Decision 2003/812/EC (Text with EEA relevance)

ANNEX II

Model certificates as referred to in Article 2

PART I

MODEL CERTIFICATE FOR IMPORTS OF CHILLED, FROZEN OR PREPARED FROGS' LEGS INTENDED FOR HUMAN CONSUMPTION COUNTRY: Veterinary certificate to EU

	I.1.	Consignor	1.2.	Certificate ref	erence No	l.2.a.
		Name Address	I.3. Central competent authority			
		Address	1.4.	Local compete	ent authority	
		Tel.				
ŧ	1.5.	Consignee	I.6.			
me		Name				
sigr		Address				
con						
per		Postcode				
atch		Tel.				
Part I: Details of dispatched consignment	I.7.	Country of ISO code I.8. origin	1.9.	Country of destination	ISO co	de I.10.
Detail						
Part I:	l.11.	Place of origin	I.12.			
		Name Approval number				
		Address				
		Address				
			\sim			
	I.13.	Place of loading	I.14.	Date of depar	ture	
	I.15.	Means of transport	I.16.	Entry BIP in E	U	
		Aeroplane 🛛 Ship 🖾 Railway wagon 🗖				
		Road vehicle Other	I.17.			
		Documentation references				
	I.18.	Description of commodity			I.19. Comm	odity code (HS code)
						02.08.90
						I.20. Quantity
	I.21.	Temperature of product				I.22. Number of
		Ambient Chilled		Frozen 🗖		packages
	1.23.	Seal/Container No				I.24. Type of packaging

Status: Point in time view as at 28/04/2016.

1.25.	Commodities cert	tified for:				
	Human consumpt	tion 🗖				
1.26.				I.27. For impo	ort or admission into EU	
1.28.	Identification of th	e commodities				
(so	Species Treatment type establish		val number of blishments Number of packages acturing plant		Net weight	

	COUNTRY Model FRG Frogs' legs							
	П.	Health information	II.a.	Certificate reference No	II.b.			
	II.1.	Public Health Attestation						
Part II: Certification		I, the undersigned, declare that I am aware of the relevant provisions of Regulation (EC) No 178/2002 of the European Parliament and of the Council of 28 January 2002 laying down the general principles and requirements of food law, establishing the European Food Safety Authority and laying down procedures in matters of food safety (OJ L 31, 1.2.2002, p. 1), Regulation (EC) No 852/2004 of the European Parliament and of the Council of 29 April 2004 on the hygiene of foodstuffs (OJ L 139, 30.4.2004, p. 1) and Regulation (EC) No 853/2004 of the European Parliament and of the Council of 29 April 2004 on the hygiene and of the Council of 29 April 2004 and Regulation (EC) No 853/2004 of the European Parliament and of the Council of 29 April 2004 of the European Parliament and of the Council of 29 April 2004 of the European Parliament and of the Council of 29 April 2004 and Regulation (EC) No 853/2004 of the European Parliament and of the Council of 29 April 2004 and Regulation (EC) No 853/2004 of the European Parliament and of the Council of 29 April 2004 laying down specific hygiene rules for food of animal origin (OJ L 139, 30.4.2004, p. 55) and certify that the frogs' legs described above were produced in accordance with those requirements, in particular that they:						
: Certi		 come from (an) establishme accordance with Article 5 of I 		plementing a programme based on (EC) No 852/2004;	on the HACCP principles in			
Ē		and						
Ра			stored in	n bled, prepared and, where and a hygienic manner in accordan (EC) No 853/2004.				
	Notes							
	Part I:							
		ox reference I.11: Place of origin: na	me and	address of the dispatch establishing	nent			
	— в	ox reference I.15: Registration numl ame (ship). Separate information is t	ber (railv	vay wagons or container and lorri	es), flight number (aircraft) or			
	— в	ox reference I.20: Indicate total gros	s weight	and total net weight.	-			
		ox reference I.23: Identification of co	-	-	le.			
		ox reference I.28: Treatment type: fr						
		7,						
	Part II	:						
	 The colour of the stamp and signature must be different from that of the other particulars in the certificate. 							
	Official inspector							
		Name (in capital letters):		Qu	alification and title:			
		Date:		Sig	gnature:			
		Stamp:						

COUNTRY:

PART II

MODEL CERTIFICATE FOR IMPORTS OF CHILLED, FROZEN, SHELLED, COOKED, PREPARED OR PRESERVED SNAILS INTENDED FOR HUMAN CONSUMPTION

	l.1.	Consignor	I.2. Certificate reference No I.2.a.			
		Name Address	I.3. Central competent authority			
		Address	I.4. Local competent authority			
		Tel.				
	1.5.	Consignee	1.6.			
lent		Name				
nsignn		Address				
еф со		Postcode				
patch		Tel.				
Part I: Details of dispatched consignment	1.7.	Country of ISO code I.8. origin	I.9. Country of ISO code I.10. destination			
	l.11.	Place of origin	I.12.			
		Name Approval number				
		Address				
	I.13.	Place of loading	I.14. Date of departure			
	l.15.	Means of transport	I.16. Entry BIP in EU			
		Aeroplane □ Ship □ Railway wagon □				
		Road vehicle	I.17.			
		Identification				
		Documentation references				
	l.18.	Description of commodity	I.19. Commodity code (HS code)			
			I.20. Quantity			
	I.21.	Temperature of product Ambient Chilled	Frozen			
	1.23.	Seal/Container No	I.24. Type of packaging			

Veterinary certificate to EU

I.25. Commodities certified for:						
Human consumption 🗖						
1.26.		I.27. For import or admission into EU				
I.28. Identification of the commoditie	s					
Species (scientific name)	Treatment type	Approval number of establishments				
Number of packages	Net weight					
Manufacturing plant						

	COUNTRY Model SNS Snails							
	II. Health information II.a. Certificate reference No II.b.							
	II.1.	Public Health Attestation						
Part II: Certification	I, the undersigned, declare that I am aware of the relevant provisions of Regulation (EC) No 178/2002 the European Parliament and of the Council of 28 January 2002 laying down the general principles are requirements of food law, establishing the European Food Safety Authority and laying down procedure in matters of food safety (OJ L 31, 1.2.2002, p. 1), Regulation (EC) No 852/2004 of the European Parliament and of the Council of 29 April 2004 on the hygiene of foodstuffs (OJ L 139, 30.4.2004, p. and Regulation (EC) No 853/2004 of the European Parliament and of the Council of 29 April 2004 on the hygiene of foodstuffs (OJ L 139, 30.4.2004, p. and Regulation (EC) No 853/2004 of the European Parliament and of the Council of 29 April 2004 on the hygiene of foodstuffs (OJ L 139, 30.4.2004, p. and Regulation (EC) No 853/2004 of the European Parliament and of the Council of 29 April 2004 laying down specific hygiene rules for food of animal origin (OJ L 139, 30.4.2004, p. 55) and certify that the snails described above were produced in accordance with those requirements, in particular that they:							
: Certi		 come from (an) establishme accordance with Article 5 of 		nplementing a programme based on (EC) No 852/2004;	on the HACCP principles in			
iii t		and						
Pai			nygienic	appropriate, shelled, cooked, p manner in accordance with the 2004.				
	— E	-	ber (rail\ to be pro ate Harm s weight ntainer/s	way wagons or container and lorri ovided in case of unloading and rei nonised System (HS) code under t and total net weight. Seal number: only where applicab	es), flight number (aircraft) or loading. the following headings: 03.07,			
	Part II:							
	— The colour of the stamp and signature must be different from that of the other particulars in the certificate.							
	Official inspector							
		Name (in capital letters):		Qu	alification and title:			
		Date:		Sig	gnature:			
		Stamp:			-			

PART III

MODEL CERTIFICATE FOR IMPORTS OF GELATINE INTENDED FOR HUMAN CONSUMPTION

COUNTRY:

Veterinary certificate to EU

	l.1.	Consignor	I.2. Certificate ref	ference No	l.2.a.	
		Name Address	I.3. Central competent authority			
		Address	I.4. Local compet	tent authority		
		Tel.				
	1.5.	Consignee	I.6.			
nen		Name				
ignr		Address				
suo:						
led o		Postcode				
atch		Tel.				
Part I: Details of dispatched consignment	1.7.	Country of ISO code I.8. origin	I.9. Country of destination	ISO coo	de I.10.	
Deta						
Part I:	l.11.	Place of origin	I.12.	I		
		Name Approval number				
		Address				
		Address				
	I.13.	Place of loading	I.14. Date of depart	rture		
	I.15.	Means of transport	I.16. Entry BIP in E	EU		
		Aeroplane 🛛 Ship 🖾 Railway wagon 🗆				
		Road vehicle Other	l.17.			
		Identification				
		Documentation references				
	I.18.	Description of commodity		I.19. Commo	odity code (HS code)	
					I.20. Quantity	
	I.21.	Temperature of product Ambient Chilled	Frozen 🗖		I.22. Number of packages	
	I.23.	Seal/Container No			I.24. Type of packaging	

Status: Point in time view as at 28/04/2016.

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	COUN	ITRY				Model GEL Gelatine intended for human consumption					
	П.	Heal	th in	formation	II.a.	Certificate reference No	II.b.				
	II.1.	Pub	lic He	alth Attestation							
Part II: Certification		the f requ in m Parli and dowr	Europ ireme atters amen Regu n spe	ean Parliament and of ints of food law, establis of food safety (OJ L t and of the Council of lation (EC) No 853/2004 cific hygiene rules for f	the Cour shing the 31, 1.2 29 April 4 of the t food of a	a aware of the relevant provisions of Regulation (EC) No 178/2002 of e Council of 28 January 2002 laying down the general principles and ng the European Food Safety Authority and laying down procedures 1, 1.2.2002, p. 1), Regulation (EC) No 852/2004 of the European April 2004 on the hygiene of foodstuffs (OJ L 139, 30.4.2004, p. 1) f the European Parliament and of the Council of 29 April 2004 laying d of animal origin (OJ L 139, 30.4.2004, p. 55) and certify that the ced in accordance with those requirements, in particular that:					
art II: C		-		mes from (an) establish ordance with Article 5 of		implementing a programme base ion (EC) No 852/2004;	ed on the HACCP principles in				
æ		_		as been produced from tion XIV of Annex III to F		naterials that met the requirem in (EC) No 853/2004;	ents of Chapters I and II of				
		-		s been manufactured ir ex III to Regulation (EC)		ance with the conditions set out i /2004;	n Chapter III of Section XIV of				
		_	of C		ĖC) No	of Section XIV of Annex III to Reg 2073/2005 of 15 November 2005 1);					
		and	if fron	n ruminant origin, excep	t for gela	tine derived from hides and skins	of ruminants,				
		(1) ei	ther								
		_	No 9 the	999/2001 of the Europea prevention, control an	an Parlia d eradio	n classified in accordance with ament and of the Council of 22 M cation of certain transmissible ntry or region posing a negligible	lay 2001 laying down rules for spongiform encephalopathies				
		_				was derived were born, continuo and passed ante-mortem and pos					
		_	if in	the country or region the	ere have	been BSE indigenous cases:					
		 (i) it comes from animals which were born after the date from which the ban on the ruminants with meat-and-bone meal and greaves derived from ruminants had been or 									
	 the products of bovine, ovine and caprine animal origin do not contain and a from specified risk material as defined in Annex V to Regulation (EC) No mechanically separated meat obtained from bones of bovine, ovine or caprine a 						lation (EC) No 999/2001, or				
		(¹) or									
		 [it comes from a country or a region classified in accordance with Article 5(2) of Regulation (No 999/2001 as a country or region posing a controlled BSE risk; 									
		 the animals from which the gelatine was derived passed ante-mortem and post-mortem inspect 									
		_	stun slau	ning by means of gas	s injecte fter stun	destined for export was derived h ad into the cranial cavity or kil ning of central nervous tissue by the cranial cavity;	led by the same method or				
		_	Reg			not derived from specified risk ma mechanically separated meat ob					

 No 999/2001 as a country or region with an undetermined BSE risk; the animals from which the gelatine was derived have not been fed meat-and-bone meal or greaves derived from ruminants and passed ante-mortem and post-mortem inspections; the animals from which the gelatine was 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; the gelatine is not derived from: specified risk material as defined in Annex V to Regulation (EC) No 999/2001; nervous and lymphatic tissues exposed during the deboning process; mechanically separated meat obtained from bones of bovine, ovine or caprine animals.] Notes Part I: Box reference I.11: Place of origin: name and address of the dispatch establishment. 	COUNTRY			Model GEL Gelatine intended for human consumption						
 [It comes from a country or a region classified in accordance with Article 5(2) of Regulation (EC) No 999/2001 as a country or region posing a negligible BSE risk; the gelatine is derived from animals which passed ante-mortem and post-mortem inspections; the gelatine is derived both from animals born, continuously reared and slaughtered in a country or region with negligible BSE risk in accordance with Article 5(2) Of Regulation (EC) No 999/2001, and, if there have been BSE indigenous cases in the country or region posing a controlled BSE risk, and which have not been slaughtered as a country or region posing a controlled BSE risk, and which have not been slaughtered as a country or region posing a controlled BSE risk, and which have not been slaughtered as laceration after stunning by means of as injected into the cranial cavity or reliade by the same method or slaughtered by laceration after stunning or prain posing a controlled BSE risk, and which have not been slaughtered as laceration after stunning the prain and use of the equation (EC) No 999/2001 is a country or region classified in accordance with Article 5(2) of Regulation (EC) No 999/2001 as a country or region with an undetermined BSE risk; the gelation (EC) No 999/2001 as a country or region with an undetermined BSE risk; the animals from which the gelatine was derived have not been slaughtered after stunning to means of gas injected into the cranial cavity or killed by the same method or slaughtered by laceration after stunning to mutinantia nervous tissue by means of an elongated rod-shaped instrument introduced into the cranial cavity: the animals from which the gelatine was derived have not been slaughtered after stunning to means of gas injected into the cranial cavity or killed by the same method or slaughtered by laceration after stunning to material nervous tissue by means of an elongated rod-shaped instrument introduced into the cranial cavity or killed by the sa	II. Health information				II.a.	Certificate reference No	II.b.			
No 999/2001 as a country or region posing a negligible BSE risk; — the gelatine is derived from animals which passed ante-mortem and post-mortem inspections; — the gelatine is derived both from animals born, continuously reared and slauphtered in a country or region with medligible BSE risk in accordance with Article 5(2) of Regulation (EC) No 999/2001 and a country or region posing a controled base method or slaughtered and snaminals born in a country or region posing a controled BSE risk, and which have not been slaupdifered after stuming by means of gas nigcted into the cranial cavity or killed by the same method or slaughtered by laceration after stuming of central nervous tissue by means of an elongated rod-shaped instrument introduced into the cranial cavity or wine or caprine animals.] (f) or — the gelatine does not contain and is not derived from specified risk material as defined in Annex V to Regulation (EC) No 999/2001, or mechanically separated meat obtained from bones of bovine, ovine or caprine animals.] (f) or — the animals from which the gelatine was derived have not been feature. — the animals from which the gelatine was derived have not been featured or slaughtered after stunning by means of an elongate rod-shaped antivous tissue by means of an elongate rod-shaped instrument introduced into the cranial cavity. — the animals from which the gelatine was derived have not been feat meat-and-bone meal or greaves derived from the cranial cavity. — the animals from which the gelatine was derived have not been slaughtered after stunning by lacerati		(¹) 0	r							
 the gelatine is derived both from animals born, continuously reared and slaughtered in a country or region with negligible BSE risk in accordance with Article 5(2) of Regulation (EC) No 999/2001, and, if there have been BSE indigenous cases in the country or region born after the date from which the ban on the feeding of ruminants with meat-and-borne meal and greaves derived from ruminants had been enforced, and from animals born in a country or region poing a controlled BSE risk, and which have not been slaughtered after stunning by means of gas injected into the cranial cavity or reliably on reliably on the same method or slaughtered is as a country or region poing a controlled BSE risk, and which have not been slaughtered after stunning by means of gas injected into the cranial cavity. the gelatine does not contain and is not derived from specified risk material as defined in Annex V to Regulation (EC) No 999/2001, or mechanically separated meat obtained from bones of bovine, ovine or caprine animals.] (f) or If comes from a country or a region classified in accordance with Article 5(2) of Regulation (EC) No 999/2001 as a country or region with an undetermined BSE risk; the animals from which the gelatine was derived have not been fed meat-and-bone meal or greaves derived from ruminants and passed ante-montern and post-mortern inspections; the animals from which the gelatine was 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 or slaughtered by laceration after stunning or counts and passed ante-mortern and post-mortern inspections; the animals from which the gelatine was derived have not been slaughtered after stunning or cancertal nervous tissue by means of gas injected into the cranial cavity. the gelatine is not derived from: (i) specified risk material										
 region with negligible BSE risk in accordance with Article 5(2) of Regulation (EC) No 099/2001, if there have been BSE indigenous cases in the country or region, points and for animals born in a country or region posing a controlled BSE risk, and which 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 by means of an elingated rod-shaped instrument introduced into the cranial cavity, or killed by the same method or slaughtered by laceration after stunning of central nervous tissue by means of an elingated rod-shaped instrument introduced into the cranial cavity. the gelatine does not contain and is not derived from specified risk material as defined in Annex V to Regulation (EC) No 999/2001, or mechanically separated meat obtained from bones of bovine, ovine or caprine animals.] (¹) or It comes from a country or a region classified in accordance with Article 5(2) of Regulation (EC) No 999/2001 as a country or region with an undetermined BSE risk; the animals from which the gelatine was derived have not been fautrad-bone meal or greaves derived from numinants and passed ante-mortem and post-mortem inspections; the animals from which the gelatine was 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; the gelatine is not derived from: (i) specified risk material as defined in Annex V to Regulation (EC) No 999/2001; (ii) nervous and lymphatic tissues exposed during the deboning process; (iii) mechanically separated meat obtained from bones of bovine, ovine or caprine animals.] Notes<		 the gelatine is derived from animals which passed ante-mortem and post-mortem inspections; 								
 Regulation (EC) No 999/2001, or mechanically separated meat obtained from bones of bovine, ovine or caprine animals.] (¹) or [it comes from a country or a region classified in accordance with Article 5(2) of Regulation (EC) No 999/2001 as a country or region with an undetermined BSE risk; the animals from which the gelatine was derived have not been fed meat-and-bone meal or greaves derived from ruminants and passed ante-mortem and post-mortem inspections; the animals from which the gelatine was derived have not been slaughtered after stunning by laceration after stunning of central nervous tissue by means of an elongated rod-shaped instrument introduced into the cranial cavity; the gelatine is not derived from:		region with negligible BSE risk in accordance with Article 5(2) of Regulation (EC) No 999/2001, and if there have been BSE indigenous cases in the country or region, born after the date from which the ban on the feeding of ruminants with meat-and-bone meal and greaves derived from ruminants had been enforced, and from animals born in a country or region classified in accordance with Article 5(2) of Regulation (EC) No 999/2001 as a country or region posing a controlled BSE risk and which 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								
 [it comes from a country or a region classified in accordance with Article 5(2) of Regulation (EC) No 999/2001 as a country or region with an undetermined BSE risk; the animals from which the gelatine was derived have not been fed meat-and-bone meal or greaves derived from ruminants and passed ante-mortem and post-mortem inspections; the animals from which the gelatine was 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; the gelatine is not derived from: (i) specified risk material as defined in Annex V to Regulation (EC) No 999/2001; (ii) nervous and lymphatic tissues exposed during the deboning process; (iii) mechanically separated meat obtained from bones of bovine, ovine or caprine animals.] 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.19: Use the appropriate Harmonised System (HS) code under the heading of 35.03. Box reference I.20: Indicate total gross weight and total net weight. 		_	Regulation (EC) N	lo 999/200						
 No 999/2001 as a country or region with an undetermined BSE risk; the animals from which the gelatine was derived have not been fed meat-and-bone meal or greaves derived from ruminants and passed ante-mortem and post-mortem inspections; the animals from which the gelatine was 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; the gelatine is not derived from: (i) specified risk material as defined in Annex V to Regulation (EC) No 999/2001; (ii) nervous and lymphatic tissues exposed during the deboning process; (iii) mechanically separated meat obtained from bones of bovine, ovine or caprine animals.] 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.19: Use the appropriate Harmonised System (HS) code under the heading of 35.03. Box reference I.20: Indicate total gross weight and total net weight. 		(¹) O	r							
 derived from ruminants and passed ante-mortem and post-mortem inspections; the animals from which the gelatine was 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; the gelatine is not derived from: (i) specified risk material as defined in Annex V to Regulation (EC) No 999/2001; (ii) nervous and lymphatic tissues exposed during the deboning process; (iii) mechanically separated meat obtained from bones of bovine, ovine or caprine animals.] 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.19: Use the appropriate Harmonised System (HS) code under the heading of 35.03. Box reference I.20: Indicate total gross weight and total net weight. 		-					Article 5(2) of Regulation (EC)			
 means of gas injected into the cranial cavity or killed by the same method or slaughtered by laceration after sturning of central nervous tissue by means of an elongated rod-shaped instrument introduced into the cranial cavity; the gelatine is not derived from: (i) specified risk material as defined in Annex V to Regulation (EC) No 999/2001; (ii) nervous and lymphatic tissues exposed during the deboning process; (iii) mechanically separated meat obtained from bones of bovine, ovine or caprine animals.] 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.19: Use the appropriate Harmonised System (HS) code under the heading of 35.03. Box reference I.20: Indicate total gross weight and total net weight. 		-								
 (i) specified risk material as defined in Annex V to Regulation (EC) No 999/2001; (ii) nervous and lymphatic tissues exposed during the deboning process; (iii) mechanically separated meat obtained from bones of bovine, ovine or caprine animals.] 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.19: Use the appropriate Harmonised System (HS) code under the heading of 35.03. Box reference I.20: Indicate total gross weight and total net weight.		-	means of gas inje laceration after stu	ected into nning of ce	the cra entral ne	nial cavity or killed by the sai	me method or slaughtered by			
 (ii) nervous and lymphatic tissues exposed during the deboning process; (iii) mechanically separated meat obtained from bones of bovine, ovine or caprine animals.] 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.19: Use the appropriate Harmonised System (HS) code under the heading of 35.03. Box reference I.20: Indicate total gross weight and total net weight. 		—	the gelatine is not o	derived fro	m:					
 (iii) mechanically separated meat obtained from bones of bovine, ovine or caprine animals.] 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.19: Use the appropriate Harmonised System (HS) code under the heading of 35.03. Box reference I.20: Indicate total gross weight and total net weight. 			(i) specified risk	material a	s define	d in Annex V to Regulation (EC)	No 999/2001;			
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.19: Use the appropriate Harmonised System (HS) code under the heading of 35.03. — Box reference I.20: Indicate total gross weight and total net weight.			(ii) nervous and	lymphatic	tissues e	exposed during the deboning pro	ocess;			
 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.19: Use the appropriate Harmonised System (HS) code under the heading of 35.03. Box reference I.20: Indicate total gross weight and total net weight. 			(iii) mechanically	separated	l meat o	btained from bones of bovine, or	vine or caprine animals.]			
 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.19: Use the appropriate Harmonised System (HS) code under the heading of 35.03. Box reference I.20: Indicate total gross weight and total net weight. 										
 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.19: Use the appropriate Harmonised System (HS) code under the heading of 35.03. Box reference I.20: Indicate total gross weight and total net weight. 										
 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.19: Use the appropriate Harmonised System (HS) code under the heading of 35.03. Box reference I.20: Indicate total gross weight and total net weight. 	Not	tes								
 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.19: Use the appropriate Harmonised System (HS) code under the heading of 35.03. Box reference I.20: Indicate total gross weight and total net weight. 	Par	rt I:								
 name (ship). Separate information is to be provided in case of unloading and reloading. Box reference I.19: Use the appropriate Harmonised System (HS) code under the heading of 35.03. Box reference I.20: Indicate total gross weight and total net weight. 	-	Box refe	erence I.11: Place of	origin: nai	me and	address of the dispatch establis	nment.			
 Box reference I.20: Indicate total gross weight and total net weight. 	-									
	-	Box refe	erence I.19: Use the	appropriat	e Harmo	onised System (HS) code under	the heading of 35.03.			
 Box reference I.23: Identification of container/Seal number: only where applicable. 	-	Box refe	erence I.20: Indicate	total gross	s weight	and total net weight.				
	-	Box refe	erence I.23: Identifica	ation of co	ntainer/S	Seal number: only where applica	ble.			

COUN	TRY		Gelatine inten	Model GEL ded for human consumption		
н.	Health information	II.a.	Certificate reference No	II.b.		
Part II	:					
(¹) De	elete as appropriate.					
— Tř	ne colour of the stamp and signature	e must be	different from that of the other p	articulars in the certificate.		
Official	l veterinarian		N	ame (in capital letters):		
	Qualification and title:					
Date: Signature:						
	Stamp:					

PART IV

MODEL CERTIFICATE FOR IMPORTS OF COLLAGEN INTENDED FOR HUMAN CONSUMPTION

COUNTRY:

Veterinary certificate to EU

	l.1.	Consignor	1.2.	Certificate refe	erence No	l.2.a.
		Name	I.3. Central competent authority			
		Address		Local compete	ent authority	
		Tel.				
	1.5.	Consignee	I.6.			
lent		Name				
gnn		Address			/	
onsi						
o c		Postcode				
tche		Tel.				
Part I: Details of dispatched consignment	1.7.	Country of ISO code I.8.	1.9.	Country of	ISO cod	de I.10.
ls of		origin	1.0.	destination	100 000	
Detai						
÷	144		140			
Ра	1.11.	Place of origin	I.12.			
		Name Approval number				
		Address				
		Address				
			\leq			
	I.13.	Place of loading	I.14.	Date of depar	ture	
	I.15.	Means of transport	I.16.	Entry BIP in E	U	
		Aeroplane Ship Railway wagon				
		Road vehicle Other	I.17.			
		Identification				
		Documentation references				
	I.18.	Description of commodity			I.19. Commo	odity code (HS code)
				l		I.20. Quantity
	I.21.	Temperature of product				I.22. Number of
		Ambient Chilled	F	Frozen 🗖		packages
	I.23.	Seal/Container No				I.24. Type of packaging

1.25.	I.25. Commodities certified for:								
	Human consumpt	tion 🗖							
1.26.				1.27.	For impo	rt or admission into EU			
1.28.	Identification of th	e commodities							
(sc	Species Date production Approval n establish (scientific name) (dd/mm/yyyy) Manufactur		shments Number of packages			Net weight			

	COUN	TRY		Model COL Collagen intended for human consumption							
	П.	Heal	th information	II.a.	Certificate reference No	II.b.					
	II.1.	Pub	lic Health Attestation								
Part II: Certification		the E requ in m Parli and dowr	European Parliament and of t irements of food law, establis latters of food safety (OJ L ament and of the Council of Regulation (EC) No 853/2004 n specific hygiene rules for fi	he Coun shing the 31, 1.2. 29 April I of the E ood of a	e of the relevant provisions of Re icil of 28 January 2002 laying dox European Food Safety Authority 2002, p. 1), Regulation (EC) No 2004 on the hygiene of foodstuffs European Parliament and of the C nimal origin (OJ L 139, 30.4.200 accordance with those requirement	wn the general principles and and laying down procedures 5 852/2004 of the European s (OJ L 139, 30.4.2004, p. 1) council of 29 April 2004 laying 14, p. 55) and certify that the					
art II: C		-	it comes from (an) establishment(s) implementing a programme based on the HACCP principles in accordance with Article 5 of Regulation (EC) No 852/2004;								
æ		-	it has been produced from Section XV of Annex III to Re		naterials that met the requireme n (EC) No 853/2004;	nts of Chapters I and II of					
		-	it has been manufactured in Annex III to Regulation (EC)		ance with the conditions set out in 2004;	h Chapter III of Section XV of					
		 it satisfies the criteria of Chapter IV of Section XV of Annex III to Regulation (EC) No 853/2004 a of Commission Regulation (EC) No 2073/2005 of 15 November 2005 on microbiological criteria foodstuffs (OJ L 338, 22.12.2005, p. 1); 									
		and	if from ruminant origin, except	for colla	gen derived from hides and skins	of ruminants,					
		(¹) either									
		 [it comes from a country or a region classified in accordance with Article 5(2) of Regulation (E No 999/2001 of the European Parliament and of the Council of 22 May 2001 laying down rules the prevention, control and eradication of certain transmissible spongiform encephalopath (OJ L 147, 31.5.2001, p. 1) as a country or region posing a negligible BSE risk; 									
		—		•	was derived were born, continuou and passed ante-mortem and post	,					
		_	if in the country or region the	re have	been BSE indigenous cases:						
					vere born after the date from whi meal and greaves derived from r						
		(ii) the products of bovine, ovine and caprine animal origin do not contain and are not de from specified risk material as defined in Annex V to Regulation (EC) No 999/200 mechanically separated meat obtained from bones of bovine, ovine or caprine animals.]									
		(¹) or									
		 [it comes from a country or a region classified in accordance with Article 5(2) of Regulation No 999/2001 as a country or region posing a controlled BSE risk; 									
		 the animals from which the collagen was derived passed ante-mortem and post- inspections; 									
		_	after stunning by means of	gas inje fter stun	n destined for export were derive ected into the cranial cavity or k ning of central nervous tissue by e cranial cavity;	illed by the same method or					
		-			not derived from specified risk m mechanically separated meat ob						

со	UNTRY			Model COL Collagen intended for human consumption								
П.	Hea	lth inf	formation	II.a.	Certificate reference No	II.b.						
	(¹) 0	r										
	—				n classified in accordance with A posing a negligible BSE risk;	rticle 5(2) of Regulation (EC)						
	_	the c	collagen is derived from	animals	which passed ante-mortem and po	ost-mortem inspections;						
	_	regio if the the t had Artic and or ki	on with negligible BSE ri- ere have been BSE ind ban on the feeding of ru- been enforced, and fro- ble 5(2) of Regulation (B which have not been sha lled by the same method	isk in acc igenous iminants om anima EC) No 9 aughtere d or slaug	mals born, continuously reared ar ordance with Article 5(2) of Regul cases in the country or region, b with meat-and-bone meal and gre als born in a country or region of 199/2001 as a country or region of d after stunning by means of gas ghtered by laceration after stunnin instrument introduced into the cran	ation (EC) No 999/2001, and, orn after the date from which eaves derived from ruminants classified in accordance with posing a controlled BSE risk, injected into the cranial cavity g of central nervous tissue by						
	_	to R			not derived from specified risk m mechanically separated meat ob							
	(¹) 0	(¹) or										
	_	[it comes from a country or a region classified in accordance with Article 5(2) of Regulation (EC) No 999/2001 as a country or region with an undetermined BSE risk;										
	_		the animals from which the collagen was derived have not been fed meat-and-bone meal or greaves derived from ruminants and passed ante-mortem and post-mortem inspections;									
	_	the animals from which the collagen was 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;										
	_	the c	collagen was not derived	d from:								
		(i)	specified risk material	as define	d in Annex V to Regulation (EC) N	No 999/2001;						
		(ii)	nervous and lymphatic	tissues e	exposed during the deboning proc	ess;						
		(iii)	mechanically separate	d meat o	btained from bones of bovine, ovi	ne or caprine animals.]						
Not	tes											
Par												
_	Box refe	erence	e I.11: Place of origin: na	ame and	address of the dispatch establishr	nent.						
-					vay wagons or container and lorri vided in case of unloading and rel							
-	Box refe	erence	e I.18: This certificate ma	ay also b	e used for import of collagen casir	ngs.						
-	Box refe	erence	e I.19: Use the appropria	ate Harm	onised System (HS) code under th	ne heading of 35.04 or 39.17.						
-	Box refe	erence	e I.20: Indicate total gros	s weight	and total net weight.							
-	Box refe	erence	e I.23: Identification of co	ontainer/s	Seal number: only where applicab	le.						

со	UNTRY	Model COL Collagen intended for human consumption								
П.	Health information	II.a.	Certificate reference No	II.b.						
Part II:										
(¹)	Delete as appropriate.									
-	— The colour of the stamp and signature must be different from that of the other particulars in the certificate.									
Offi	cial veterinarian									
	Name (in capital letters):			Qualification and title:						
Date: Signature:										
	Stamp:									

PART V

MODEL CERTIFICATE FOR IMPORTS OF RAW MATERIALS FOR THE PRODUCTION OF GELATINE/ COLLAGEN INTENDED FOR HUMAN CONSUMPTION⁽¹⁾

cou	INTRY	:				Veterinary certificate to EU			
	l.1.	Consignor				1.2.	Certificate re	eference No	I.2.a.
		Name				1.3.	Central com	petent author	ity
		Address				I.4.	Local comp	etent authority	/
		Tel.							
	1.5.	Consignee				I.6.			
lent		Name							
ignn		Address							
suo:									
ped o		Postcode							
patcl		Tel.							
Part I: Details of dispatched consignment	1.7.	Country of origin	ISO code	I.8. Region of origin	Code	1.9.	Country of destination	ISO code	l.10.
Detail									
art I:	l.11.	Place of or	igin			I.12.			
•		Name Approval number							
				nber					
		Address							
						_			
	I.13.	Place of loa	ading			I.14.	Date of depa	arture	
	l.15.	Means of t	ransport			I.16. Entry BIP in EU			
		Aeroplane		Ship 🗖 🛛 Ri	ailway				
		wagon 🗖				I.17.			
		Road vehic		Other 🗖					
			ation referer	ices					
	140					_		140 0	a ditu aa da (UQ aa da)
	1.18.	Description	n of commoo	лту				1.19. Comm	odity code (HS code)
							l		I.20. Quantity
	I.21.	Temperatu	re of produc	ct					I.22. Number of
		Ambient 🗆	1]		Frozen 🗖		packages
	1.23.	Seal/Conta	iner No						I.24. Type of packaging

Status: Point in time view as at 28/04/2016.

I.25.	I.25. Commodities certified for:								
	Production of gelatine / collagen for human consumption \square								
I.26.				I.27. For impo	rt or admission into EU				
1.28.	Identification of the	commodities							
(so	Species Nature of Approval n establish (scientific name) commodity Manufactur		nments	Number of packages	Net weight				

COUNTRY Raw materials for the productio collagen intended for human												
	н.	Heal	th in	formation	II.a.	Certificate reference No	II.b.					
	II.1.	Pub	lic He	ealth Attestation								
Part II: Certification		the E requ in m Parli Regu spec No 8 for t (OJ	Europ ireme atters amer ulation 54/20 he or L 139	ean Parliament and of t ents of food law, establis s of food safety (OJ L at and of the Council of 2 n (EC) No 853/2004 of th ygiene rules for food of 004 of the European Par ganisation of official co	he Cound hing the 31, 1.2.2 29 April 2 he Europe of animal fliament a ntrols on	e of the relevant provisions of Re cil of 28 January 2002 laying do European Food Safety Authority 2002, p. 1), Regulation (EC) N 2004 on the hygiene of foodstuffs ean Parliament and of the Counc I origin (OJ L 139, 30.4.2004, and of the Council of 29 April 20 products of animal origin inter that the raw materials describ	wn the general principles and and laying down procedures b 852/2004 of the European s (OJ L 139, 30.4.2004, p. 1), I of 29 April 2004 laying down p. 55) and Regulation (EC) 04 laying down specific rules ded for human consumption					
-		_	and and	sinews described above	derive fr	ic and farmed ruminant animals, om animals which have been sla en found fit for human consump	ughtered in a slaughterhouse					
		and/or										
		 (1) [wild game hides, skins and bones described above derive from killed animals whose carcase have been found fit for human consumption following post-mortem inspection,] 										
			and	for								
		—		fish skins and bones de an consumption authoris		above derive from plants manu port,]	facturing fishery products for					
		(1) ar	nd									
		[if fro	om ru	minant origin, except for	hides an	d skins of ruminants,						
		(1) ei	ther:									
		_	No sthe	999/2001 of the Europea prevention, control and	an Parliar d eradica	n classified in accordance with A nent and of the Council of 22 M ation of certain transmissible s try or region posing a negligible B	ay 2001 laying down rules for spongiform encephalopathies					
		_	were		ed and s	erials of bovine, ovine and caprii claughtered in the country with ne tions;	ç					
		_	if in	the country or region the	re have t	been BSE indigenous cases:						
		 the animals were born after the date from which the ban on the feeding of ruminants meat-and-bone meal and greaves derived from ruminants had been enforced; or 										
	 the raw materials of bovine, ovine and caprine animal origin do not contain and are derived from specified risk material as defined in Annex V to Regulation (EC) No 999/200 mechanically separated meat obtained from bones of bovine, ovine or caprine animals;] 											
		(1) or	:									
		_	[the No §	y come from a country o 999/2001 as a country or	r a regio region p	n classified in accordance with A osing a controlled BSE risk;	rticle 5(2) of Regulation (EC)					
		_		animals from which the sed ante-mortem and pos		erials of bovine, ovine and capri n inspections;	ne animal origin were derived					

COUNTRY

II.	Heal	th information	II.a.	Certificate reference No	II.b.						
	_	were derived have not been cavity or killed by the same	n slaughte method o	of bovine, ovine and caprine anir ered after stunning by means of or slaughtered by laceration after shaped instrument introduced into	gas injected into the cranial stunning of central nervous						
	_	specified risk material as de	efined in	caprine animal origin do not con Annex V to Regulation (EC) No of bovine, ovine or caprine animal	999/2001, or mechanically						
	(1) or										
	_	[they come from a country on No 999/2001 as a country or		n classified in accordance with Ar osing a negligible BSE risk;	ticle 5(2) of Regulation (EC)						
	-	the animals from which the passed ante-mortem and pos		rials of bovine, ovine and caprine inspections;	e animal origin were derived						
	-	— the raw materials of bovine, ovine and caprine animal origin intended for export are derived both from animals which were born, continuously reared and slaughtered in a country or region with negligible BSE risk in accordance with Article 5(2) of Regulation (EC) No 999/2001, and, if there have been BSE indigenous cases in the country or region, born after the date from which the ban on the feeding of ruminants with meat-and-bone meal and greaves derived from ruminants had been enforced, and from animals born in a country or region classified in accordance with Article 5(2) of Regulation (EC) No 999/2001 as a country or region posing a controlled BSE risk, and which 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;									
	_	specified risk material as d	efined in	caprine animal origin do not com Annex V to Regulation (EC) No of bovine, ovine or caprine animal	999/2001, or mechanically						
	(1) or										
	_			n classified in accordance with Ar th an undetermined BSE risk;	ticle 5(2) of Regulation (EC)						
	_		nd-bone i	rials of bovine, ovine and caprine neal or greaves derived from r							
	-	have not been slaughtered a the same method or slaught	raw materials of bovine, ovine and caprine animal origin were derived fter stunning by means of gas injected into the cranial cavity or killed by ered by laceration after stunning of central nervous tissue by means of trument introduced into the cranial cavity;								
	_	the raw materials of bovine,	ovine and	caprine animal origin are not der	ived from:						
		(i) specified risk material a	as defined in Annex V to Regulation (EC) No 999/2001;								
		(ii) nervous and lymphatic	tissues e	xposed during the de-boning proc	ess;						
		(iii) mechanically separated	d meat ob	meat obtained from bones of bovine, ovine or caprine animals.]]							

COUNTRY

II.	Health in	formation	II.a.	Certificate referer	nce No	II.b.				
(¹) [II.2 .	Animal H	lealth Attestation								
	I, the und	lersigned official veter	inarian,	certify that the raw ma	aterials descri	ibed above:				
II.2.1.	consist of	f animal products that	satisfy t	the animal health requ	uirements belo	ow;				
II.2.2.	have bee from:	have been obtained in the territory of (1) either [:] (1) or [] (2) (3) $(4) from:$								
(1) either	[II.2.2.1	[II.2.2.1 animals that come from holdings and have remained in that territory since birth or for at least the last 3 months before slaughter; and								
	(1) either [(i) that are of the species referred to in Commission Regulation (EU) No 206/2010 of 12 March 2010 laying down lists of third countries, territories or parts thereor authorised for the introduction into the European Union of certain animals and fresh meat and the veterinary certification requirements (OJ L 73, 20.3.2010, p. 1), fulfilling all the relevant animal health import requirements laid down in that Regulation, and that were slaughtered for human consumption on a date for which import into the European Union of fresh meat from animals of those species was authorised from the country or territory thereof in accordance with Column 8 of Part 1 of Annex II to the Regulation;]									
	(1) or [(ii) that are of the species referred to in Commission Regulation (EC) No 119/2009 9 February 2009 laying down a list of third countries or parts thereof, for imports int or transit through, the Community of meat of wild leporidae, of certain wild lay mammals and of farmed rabbits and the veterinary certification requirements (OJ L 3 10.2.2009, p. 12), fulfilling all the relevant animal health import requirements laid dow in that Regulation.]]									
(¹) or	[II.2.2.1 poultry that have remained in that territory since hatching or have been imported as day-o chicks or slaughter poultry from (a) third country(ies) listed for that commodity in Part 1 Annex I to Commission Regulation (EC) No 798/2008 of 8 August 2008 laying down a list third countries, territories, zones or compartments from which poultry and poultry produc may be imported into and transit through the Community and the veterinary certificatio requirements (OJ L 226, 23.8.2008, p. 1) under conditions at least equivalent to those in the Regulation, and consist of species referred to in that Regulation, fulfilling all the releval animal health import requirements laid down in that Regulation, and were slaughtered for human consumption on a date for which import into the European Union of meat from animals of those species was authorised from the country or territory thereof in accordance with Column 6 B of Part 1 to Annex I to that Regulation.]									
(1) or	[II.2.2.1	animals that have t area:	een kill	ed in the wild in that	t territory ⁽⁵⁾ ; a	nd captured and killed in an				
		diseases for w Newcastle dise	in 25 km there has been no case/outbreak of any of the followin which the animals are susceptible: foot and mouth disease, rinderpes ease or highly pathogenic avian influenza during the prior 30 days, nor frican swine fever during the prior 40 days and							
		territory of a	at a distance that exceeds 20 km from the borders separating another country or part thereof, which is not authorised at these dates for e raw materials to the European Union, and							
			re and i			hours for chilling either to a establishment, or directly to a				

COUNTRY

		1									
II.	Health information	II.a.	Certificate reference No	II.b.							
II.2.3	2.3. have been obtained in an establishment around which, within a radius of 10 km, there has been no case/outbreak of the following diseases for which the animals are susceptible: foot and mouth disease, rinderpest, Newcastle disease or highly pathogenic avian influenza, classical or African swine fever during the prior 30 days or, in the event of a case of one of those diseases, the preparation of raw materials for exportation to the European Union has been authorised only after the removal of all meat and the total cleaning and disinfection of the establishment under the control of an official veterinarian; and:										
11.2.4				aterials not complying with the I contamination with pathogenic							
II.2.	b. have been transported in clean and sealed containers or lorries.]										
Note	25										
Part	:1:										
— Box reference I.8: Provide the code of territory as appearing in Annex II to Commission Decision 2006/766/EC of 6 November 2006 establishing the lists of third countries and territories from which imports of bivalve molluscs, echinoderms, tunicates, marine gastropods and fishery products are permitted (OJ L 320, 18.11.2006, p. 53) and/or Part 1 of Annex I to Regulation (EC) 798/2008 and/or in Part 1 of Annex I to Regulation (EC) No 119/2009 and/or Part 1 of Annex II to Regulation (EU) No 206/2010.											
	Box reference I.11: Place of number as appropriate.	origin: name and	address of the dispatch estab	lishment; registration or approval							
_			vay wagons or container and I vided in case of unloading and	orries), flight number (aircraft) or reloading.							
	Box reference I.19: Use the a 03.05, 05.05, 05.06, 05.11.91			er the following headings: 02.08,							
_	Box reference I.20: Indicate t	otal gross weight	and total net weight.								
_	Box reference I.23: Identifica	tion of container/s	Seal number: only where applic	cable.							
_	Box reference I.28: Nature	of commodity: hi	des, skins, bones, tendons and	d sinews;							
	Approv	al number of esta	ablishments: registration or app	proval number as appropriate;							
			cludes slaughterhouse, facto and processing plant.	ory vessel, cutting plant, game							
Part	: 11:										
(1)	Delete as appropriate. In ca deleted.	se of products de	erived from fishery products, t	the whole section II.2 should be							
(²)	The name and ISO code nun	nber of the export	ing country or territory or zone	as laid down in:							
	 the Annexes to Decision 	1 2006/766/EC;									
	- Anney I to Regulation (C) 798/2008·									

COUNTRY

П.	Health information	II.a. Certifi	cate reference No	II.b.				
	— Part 1 of Annex II to Regulation	EC) No 119/200	9;					
	— Part 1 of Annex II to Regulation	EU) No 206/201	0.					
(³)) If parts of the materials were derived from animals originating from an(other) third country(ies) listed in Annex II to Regulation (EU) No 206/2010 for import of that commodity into the Union, then the code(s) of country(ies) or territory(ies) and of the third country slaughtering the animals shall be indicated (the material cannot come from a country or territory that has supplementary guarantees A or F as indicated in column 5 of that Annex).							
(4)	If the materials were derived from slaughter poultry originating from an(other) third country(ies) listed in Part 1 of Annex I to Regulation (EC) No 798/2008 for imports of that commodity into the Union, then the code(s) of country(ies) or territory(ies) and of the third country slaughtering the poultry shall be indicated.							
(5)	Only for countries from where game meat intended for human consumption of the same animal species is authorised for importation into the European Union.							
_	The signature and the stamp must be	in a different col	our to that of the printing.					
NB	Note for the person responsible for the and has to accompany the consignment transported directly to the manufacture	nt until it reaches	s the border inspection po					
Offi	cial veterinarian							
	Name (in capital letters):		Qu	alification and title:				
	Date:		Sig	inature:				
	Stamp:							

COUNTRY:

Status: Point in time view as at 28/04/2016. Changes to legislation: There are outstanding changes not yet made to Commission Implementing Regulation (EU) 2016/759. Any changes that have already been made to the legislation appear in the content and are referenced with annotations. (See end of Document for details)

PART VI

MODEL CERTIFICATE FOR IMPORTS OF TREATED RAW MATERIALS FOR THE PRODUCTION OF GELATINE/ COLLAGEN INTENDED FOR HUMAN CONSUMPTION

	I.1. Consignor					I.2.	Certificate r	eference No	I.2.a.	
		Name Address				1.3.	Central com	petent author	ity	
		, luar coo				1.4.	Local comp	etent authorit	/	
		Tel.								
	1.5.	Consignee				I.6.				
men		Name								
nsign		Address								
d co		Postcode					/			
tche		Tel.								
ispat						\leq				
s of d	1.7.		ISO code	I.8. Region of origin	Code	1.9.	Country of destination	ISO code	I.10.	
Part I: Details of dispatched consignment										
Part I:	I.11.	I.11. Place of origin				I.12.				
_										
	Name Approval number Address									
						\geq				
	I.13.	Place of load	ding			I.14.	Date of dep	arture		
	I.15.	Means of tra	insport			I.16.	Entry BIP in	EU		
		Aeroplane 🗖 wagon 🗖]	Ship 🗖 🛛 R	ailway					
		Road vehicle	eΠ	Other 🗖		I.17.				
		Identification	ı							
		Documentati	ion referen	ces						
	l.18.	B. Description of commodity					I.19. Commodity code (HS code)			
									I.20. Quantity	
	I.21.	Temperature	e of produc	t					I.22. Number of	
		Ambient 🗖		Chilled]		Frozen 🗖		packages	
	1.23.	Seal/Contain	ner No						I.24. Type of packaging	

Veterinary certificate to EU

I.25.	25. Commodities certified for:							
	Production of gelatine / collagen for human consumption \square							
1.26.	.26. I.27. For import or admission into EL							
1.28.	Identification of the	commodities						
(sc	Species cientific name)	Nature of commodity	Approval n establish Manufactu	nments	Number of packages	Net weight		

	COUNTR	Y		Model TCG Treated raw materials for the production of gelatine and collagen					
	н.	Health in	formation	II.a.	Certificate reference No	II.b.			
	II.1.	Public H	ealth Attestation						
		l, the un requirem		the trea	ted raw materials described abo	ve comply with the following			
c			ey have been derived ithority	from est	ablishments under the control of	and listed by the competent			
catio		and,							
Part II: Certification		ab	(¹) [bones, hides and skins of domestic and farmed ruminant animals, pigs and poultry described above are derived from animals which have been slaughtered in a slaughterhouse and the carcasses of which have been found fit for human consumption following ante- and post-mortem inspection,]						
a,		(1)	and/or						
					nes described above are derive human consumption following po				
		(1) and/or							
			sh skins and bones de r human consumption a		above are derived from plants m d for export,]	anufacturing fishery products			
		and							
	(1) either [they are dried bones of species from bovine, ovine, caprine, porcine and equine animals farmed and wild animals, poultry including ratites and feathered game for the production of gelatine, they derived from healthy animals slaughtered in a slaughterhouse, and they is treated as follows:								
			either [crushed to pieces of approximately 15 mm and degreased with hot water at a temperature of minimum 70 °C for at least 30 minutes, minimum 80 °C for at least 15 minutes, or minimum 90 °C for at least 10 minutes, and then separated and subsequently washed and dried for at least 20 minutes in a stream of hot air with an initial temperature of minimum 350 °C, or for 15 minutes in a stream of hot air with an initial temperature of more than 700 °C.]						
		(1) or	[sun-dried for a minin	num of 42	2 days at an average temperature	of at least 20 °C.]			
		(¹) or	[acid treatment such before drying.]]	that the p	oH is maintained at less than 6 to	the core for at least one hour			
	(1) or		hides and skins of far by derived from healthy		inant animals, pig skins, poultry a and they:	skins or wild game hides and			
		(¹) either	[have undergone an for at least 7 days]	alkali tre	atment which ensures a PH>12 to	o the core followed by salting			
		(¹) or	[were dried for at leas	st 42 day	s at a temperature of at least 20 °	C.]			
		(¹) or [have undergone an acid treatment that provides at least a pH of less than 5 to the co minimum of one hour.]							
		(1) or) or [have undergone an alkali treatment which ensures a pH > 12 to the core for at least 8 hours.]]						
	(¹) or	game hic of Annex that com	les and skins from thir I to this Regulation the	d countri nat have registere	d ruminant animals, pig skins, pou es, parts of third countries and te undergone any other treatment ed or approved pursuant to Regul 3/2004	rritories referred to in Part IV than those listed above, and			
		and							
		(1) [if from ruminant origin, except for hides and skins of ruminants,							

COUNTRY

Model TCG Treated raw materials for the production of gelatine and collagen

II.	Health information		II.a.	Certificate reference No	II.b.				
	(¹) e	ither:							
	 [they come from a country or a region classified in accordance with Article 5(2) of Regulation (EC) No 999/2001 of the European Parliament and of the Council of 22 May 2001 laying down rules for the prevention, control and eradication of certain transmissible spongiform encephalopathies (OJ L 147, 31.5.2001, p. 1) as a country or region posing a negligible BSE risk; 								
	_	 the animals from which the treated raw materials of bovine, ovine and caprine animal origin were derived were born, continuously reared and slaughtered in the country with negligible BSE risk and passed ante-mortem and post-mortem inspections; 							
	_	if in the country or region th	nere have	been BSE indigenous cases:					
				e date from which the ban on the ban on the derived from ruminants had b					
		not derived from s	pecified r	vine, ovine and caprine animal o isk material as defined in An separated meat obtained from	inex V to Regulation (EC)				
	(¹) 01	r:							
	-			n classified in accordance with A posing a controlled BSE risk;	rticle 5(2) of Regulation (EC)				
	_	 the animals from which the treated raw materials of bovine, ovine and caprine animal origin wer derived passed ante-mortem and post-mortem inspections; 							
	_	 animals from which the treated raw materials of bovine, ovine and caprine animal origin destined for export were 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 o central nervous tissue by means of an elongated rod-shaped instrument introduced into the cranial cavity; 							
	_	derived from specified risk	material	ovine and caprine animal origin as defined in Annex V to Regu ad from bones of bovine, ovine o	lation (EC) No 999/2001, or				
		(1) or							
	_			n classified in accordance with A posing a negligible BSE risk;	rticle 5(2) of Regulation (EC)				
	-	the animals from which the derived passed ante-morte		aw materials of bovine, ovine an st-mortem inspections;	d caprine animal origin were				
	-	derived both from animals region with negligible BSE and, if there have been BS which the ban on the feed ruminants had been enfo accordance with Article 5(controlled BSE risk, and wi into the cranial cavity or ki	which we risk in ac SE indiger ling of rur prced, and (2) of Reg hich have lled by the	e, ovine and caprine animal or re born, continuously reared and coordance with Article 5(2) of Re- nous cases in the country or reg ninants with meat-and-bone me d from animals born in a cou- gulation (EC) No 999/2001 as a not been slaughtered after stunn e same method or slaughtered b of an elongated rod-shaped ins	d slaughtered in a country or egulation (EC) No 999/2001, ion, born after the date from al and greaves derived from intry or region classified in a country or region posing a ing by means of gas injected y laceration after stunning of				

COUNTRY

Model TCG Treated raw materials for the production of gelatine and collagen

II.	Health information		Certificate reference No	II.b.				
	derived from specified ris	k material	ovine and caprine animal origir as defined in Annex V to Regu d from bones of bovine, ovine o	lation (EC) No 999/2001, or				
	(1) or							
			n classified in accordance with A vith an undetermined BSE risk;	rticle 5(2) of Regulation (EC)				
	 the animals from which the treated raw materials of bovine, ovine and caprine animal origin were derived have not been fed meat-and-bone meal or greaves derived from ruminants and passed ante-mortem and post-mortem inspections; 							
	derived have not been sla or killed by the same met	aughtered a hod or slaug	aw materials of bovine, ovine ar fter stunning by means of gas in ghtered by laceration after stunr d instrument introduced into the	njected into the cranial cavity ning of central nervous tissue				
	 the treated raw materials 	of bovine, o	vine and caprine animal origin a	re not derived from:				
	(i) specified risk materi	al as define	d in Annex V to Regulation (EC)	No 999/2001;				
	(ii) nervous and lympha	itic tissues e	exposed during the de-boning pr	ocess;				
	(iii) mechanically separa	ated meat of	btained from bones of bovine, ov	vine or caprine animals.]]]				
(¹) [II.2.	Animal Health Attestation							
	I, the undersigned official veter	inarian, cer	tify that the treated raw materials	s described above:				
II.2.1.	consist of animal products that	satisfy the	satisfy the animal health requirements below;					
II.2.2.	have been obtained in the terri	tory(ies) of:	(¹) [] (¹) or [.] (²) (³)				
II.2.3.			nout contact with other materi en handled so as to avoid co					
II.2.4.	have been transported in clear	and sealed	d containers or lorries.]					
Notes								
Part I:								
— Box reference I.8: Provide the code of territory as appearing in Annex II to Commission Decision 2006/766/EC of 6 November 2006 establishing the lists of third countries and territories from which imports of bivalve molluscs, echinoderms, tunicates, marine gastropods and fishery products are permitted (OJ L 320, 18.11.2006, p. 53) or in Part 1 of Annex I to Commission Regulation (EC) 798/2008 of 8 August 2008 laying down a list of third countries, territories, zones or compartments from which poultry and poultry products may be imported into and transit through the Community and the veterinary certification requirements (OJ L 226, 23.8.2008, p. 1) or in Part 1 of Annex I to Commission Regulation (EC) No 119/2009 of 9 February 2009 laying down a list of third countries or parts thereof, for imports into, or transit through, the Community of meat of wild leporidae, of certain wild land mammals and of farmed rabbits and the veterinary certification requirements (OJ L 39, 10.2.2009, p. 12) or in Part 1 of Annex I to Commission Regulation (EU) No 206/2010 of 12 March 2010 laying down lists of third countries, territories, territories or parts thereof authorised for the introduction into the European Union of certain animals and fresh meat and the veterinary certification requirements (OJ L 73, 20.3.2010, p. 1).								
	 Box reference I.11: Place of origin: name and address of the dispatch establishment and approval number or competent authority identification number as appropriate. 							

 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.

COUNTRY

Model TCG Treated raw materials for the production of gelatine and collagen

				of genatine and conagen					
П.	Health information	II.a.	Certificate reference No	II.b.					
—	 Box reference I.19: Use the appropriate Harmonised System (HS) code under the following headings: 03.05, 05.05, 05.06, 05.11.91, 05.11.99, 41.01, 41.02, 41.03. 								
_	Box reference I.20: Indicate total	gross weight	and total net weight.						
—	Box reference I.23: Identification	of container/	Seal number: only where applicab	le.					
_	Box reference I.28: Nature o	f commodity:	hides, skins, bones, tendons and	sinews;					
			^f establishments: approval num as appropriate;	ber or competent authority					
			includes slaughterhouse, factory nt and processing plant;	vessel, cutting plant, game					
	Approva	<i>number:</i> who	en applicable.						
Par	t II:								
(¹)	Delete as appropriate. In case of deleted.	of products d	erived from fishery products, the	whole section II.2 should be					
(²)	The name and ISO code number	of the export	ing country or territory or zone as	laid down in:					
	 Part 1 of Annex II to Regula 	tion (EU) No	206/2010;						
	— Annex I to Regulation (EC)	798/2008;							
	 Part 1 of Annex II to Regula 	tion (EC) No	119/2009.						
(³)	⁽³⁾ If parts of the materials were derived from animals originating from an(other) third country(ies) listed in Annex I to Commission Implementing Regulation (EU) 2016/759 of 28 April 2016 drawing up lists of third countries, parts of third countries and territories from which Member States are to authorise the introduction into the Union of certain products of animal origin intended for human consumption, laying down certificate requirements, amending Regulation (EC) No 2074/2005 and repealing Decision 2003/812/EC (OJ L 126, 14.5.2016, p. 13), the code(s) of country(ies) or territory(ies) shall be indicated.								
_	The signature and the stamp mus	st be in a diffe	erent colour to that of the printing.						
NB	Note for the person responsible f has to accompany the consignn transported directly to the manufa	nent until it r	eaches the border inspection pos						
—	The time of transportation may be	e included in	the duration of treatment.						
Offi	Official veterinarian								
	Name (in capital letters):		(Qualification and title:					
	Date:		:	Signature:					
	Stamp:								

PART VII

MODEL CERTIFICATE FOR IMPORTS OF HONEY, ROYAL JELLY AND OTHER APICULTURE PRODUCTS INTENDED FOR HUMAN CONSUMPTION COUNTRY: Veterinary certificate to EU

	l.1.	Consignor	1.2.	Certificate ref	erence No	l.2.a.
	Name Address		1.3.	Central comp	etent authority	
		Address	I.4.	Local compete	ent authority	
		Tel.				
	1.5.	Consignee	I.6.			
nent		Name				
ignn		Address				
ons						
ed c		Postcode				
atch		Tel.				
Part I: Details of dispatched consignment	I.7.	Country of ISO code I.8.	1.9.	Country of destination	ISO co	de I.10.
ï	1 1 1	Place of origin	I.12.			
Ра	1. 1 1.		1.12.			
		Name Approval number				
		Address				
		, dai ooo				
			\leq			
	I.13.	Place of loading	I.14.	Date of depar	ture	
	l.15.	Means of transport	I.16.	Entry BIP in E	U	
		Aeroplane 🛛 Ship 🖾 Railway wagon 🗖				
		Road vehicle C Other	l.17.			
		Identification				
		Documentation references			-	
	I.18.	Description of commodity			I.19. Comm	odity code (HS code)
						I.20. Quantity
	I.21.	Temperature of product Ambient Chilled		Frozen 🗖		I.22. Number of packages
	I.23.	Seal/Container No				I.24. Type of packaging

I.25.	Commodities cert	ified for:					
	Human consumpt	ion 🗖					
1.26.				1.27.	For impo	rt or admission into EU	
1.28.	Identification of th	e commodities					
(so	Species ientific name)	Treatment type	Approval n establish Manufactur			Number of packages	Net weight

	COUN	ITRY			Honey, royal jelly a	Model HON nd other apiculture products				
	П.	Health information	ı	II.a.	Certificate reference No	II.b.				
	II.1.	Public Health Atte	station							
Part II: Certification		I, the undersigned, declare that I am aware of the relevant provisions of Regulation (EC) No 178/2002 of the European Parliament and of the Council of 28 January 2002 laying down the general principles and requirements of food law, establishing the European Food Safety Authority and laying down procedures in matters of food safety (OJ L 31, 1.2.2002, p. 1), Regulation (EC) No 852/2004 of the European Parliament and of the Council of 29 April 2004 or the hygiene of foodstuffs (OJ L 139, 30.4.2004, p. 1) and Regulation (EC) No 853/2004 of the European Parliament and of the Council of 29 April 2004 or the hygiene of foodstuffs (OJ L 139, 30.4.2004, p. 1) and Regulation (EC) No 853/2004 of the European Parliament and of the Council of 29 April 2004 of the European Parliament and of the Council of 29 April 2004 of the Hygiene of foodstuffs (OJ L 139, 30.4.2004, p. 1) and Regulation (EC) No 853/2004 of the European Parliament and of the Council of 29 April 2004 laying down specific hygiene rules for food of animal origin (OJ L 139, 30.4.2004, p. 55) and certify that honey, royal jelly and other apiculture products described above were produced in accordance with those requirements, in particular that they:								
II: Ce			n) establishme vith Regulation		plementing a programme base 852/2004;	d on the HACCP principles in				
Part		accordance w			opriate, prepared, packaged and Annex II to Regulation (EC) No					
		in accordance substances a	e with Council and residues th	Directiv nereof ir	and products thereof provided e 96/23/EC of 29 April 1996 of live animals and animal proc	measures to monitor certain lucts and repealing Directives				
					Decisions 89/187/EEC and 91/60 reof, are fulfilled.	64/EEC (OJ L 125, 23.5.1996,				
	Notes									
	Part I:	:								
		ox reference I.11: P leans registration nur		name a	nd address of the dispatch es	tablishment. Approval number				
					vay wagons or container and lor vided in case of unloading and r					
		ox reference I.19: Us 4.10.	e the appropria	te Harm	onised System (HS) code under	the following headings: 04.09,				
	— Во	ox reference I.20: Ind	icate total gross	s weight	and total net weight.					
	— Во	ox reference I.23: Ide	ntification of co	ntainer/s	Seal number: only where applica	ble.				
	— В	ox reference I.28:	Treatment in pasteurisation		ndicate 'ultrasonication', 'ho ermal treatment'.	mogenisation', ultrafiltration',				
					f establishments: approval nu as appropriate	mber or competent authority				
	Part II:									
	 The colour of the stamp and signature must be different from that of the other particulars in the certificate. 									
	Official inspector									
	Name (in capital letters): Qualification and title:									
		Date:			S	ignature:				
		Stamp:								

PART VIII

MODEL CERTIFICATE FOR IMPORTS OF HIGHLY REFINED CHONDROITIN SULPHATE, HYALURONIC ACID, OTHER HYDROLYSED CARTILAGE

Status: Point in time view as at 28/04/2016. Changes to legislation: There are outstanding changes not yet made to Commission Implementing Regulation (EU) 2016/759. Any changes that have already been made to the legislation

appear in the content and are referenced with annotations. (See end of Document for details)

AND AMINO ACIDS INTENDED FOR HUMAN CONSUMPTION COUNTRY: Veterinary certificate to EU Consignor 1.2. I.1. Certificate reference No I.2.a. Name 1.3. Central competent authority Address 1.4. Local competent authority Tel. 1.6. 1.5. Consignee Part I: Details of dispatched consignment Name Address Postcode Tel. Country of ISO code 1.8. I.10. 1.7. 1.9. Country of ISO code destination origin I.11. Place of origin I.12. Approval number Name Address I.13. Place of loading I.14. Date of departure I.15. Means of transport I.16. Entry BIP in EU Ship 🗖 Aeroplane 🛛 Railway wagon 🛛 I.17. Road vehicle Other 🗖 Identification Documentation references I.18. Description of commodity I.19. Commodity code (HS code) I.20. Quantity I.21. Temperature of product I.22. Number of packages Ambient 🗖 Chilled D Frozen 🗖 I.23. Seal/Container No I.24. Type of packaging

PRODUCTS, CHITOSAN, GLUCOSAMINE, RENNET, ISINGLASS

1.25.	Commodities cert	ified for:					
	Human consumpt	tion 🗖					
1.26.				1.27.	For impo	rt or admission into EU	
1.28.	Identification of th	e commodities					
(sc	Species cientific name)	Date production (dd/mm/yyyy)	Approval n establish Manufactu			Number of packages	Net weight

COL	INIT	DV
	ואוכ	IN I

Model HRP Highly refined chondroitin sulphate, hyaluronic acid, other hydrolysed cartilage products, chitosan, glucosamine, rennet, isinglass and amino acids for human consumption

	П.	Health information		II.a.	Certificate reference No	II.b.				
	II.1.	Public Health Attestation								
ification		I, the undersigned, declare that I am aware of the relevant provisions of Regulation (EC) No 178/2002 of the European Parliament and of the Council of 28 January 2002 laying down the general principles and requirements of food law, establishing the European Food Safety Authority and laying down proceduress in matters of food safety (OJ L 31, 1.2.2002, p. 1), Regulation (EC) No 852/2004 of the European Parliament and of the Council of 29 April 2004 on the hygiene of foodstuffs (OJ L 139, 30.4.2004, p. 1) and Regulation (EC) No 853/2004 of the European Parliament and of the Council of 29 April 2004 on the hygiene of foodstuffs (OJ L 139, 30.4.2004, p. 1) and Regulation (EC) No 853/2004 of the European Parliament and of the Council of 29 April 2004 alying down specific hygiene rules for food of animal origin (OJ L 139, 30.4.2004, p. 55) and certify that the highly refined products described above were produced in accordance with those requirements, in particular:								
 down specific hygiene rules for food of animal origin (OJ L 139, 30.4.2004, p. 55) and cer highly refined products described above were produced in accordance with those requi particular: — that they come from (an) establishment(s) implementing a programme based on t principles in accordance with Regulation (EC) No 852/2004; — that they have been handled and, where appropriate, prepared, packaged and stored ir manner in accordance with the requirements of Annex II to Regulation (EC) No 852/2004; 										
		— and	that	they comply with the req	uirement	s of Section XVI of Annex III to	Regulation (EC) No 853/2004;			
		_	(1) [if	amino acids, that						
			(i)	human hair was not us	ed as a s	ource for their manufacture; an	t			
		 they comply with Regulation (EC) No 1333/2008 of the European Parliament and of th Council of 16 December 2008 on food additives ((OJ L 354, 31.12.2008, p. 16)] 								
	Notes									
	Part I:									
	— Вс	ox refe	rence	I.11: Place of origin: na	me and a	ddress of the dispatch establis	nment.			
						ay wagons or container and lo rided in case of unloading and r	ries), flight number (aircraft) or eloading.			
				e I.19: Use the appropr 29.32, 35.07, 35.03 or 3		monised System (HS) code u	nder the heading of 21.06.90,			
	— Во	ox refe	rence	I.20: Indicate total gross	s weight a	and total net weight.				
	— Во	ox refe	rence	I.23: Identification of co	ntainer/S	eal number: only where applica	ble.			
	Part II									
	(1) De	elete a	s app	ropriate.						
	— Tł	 The colour of the stamp and signature must be different from that of the other particulars in the certificate. 								
	Officia	veter	inaria	n		١	lame (in capital letters):			
	Qualification and title:									
		Date				S	Signature:			
	Stamp:									

(1) Unless covered by Part VI.

Status:

Point in time view as at 28/04/2016.

Changes to legislation:

There are outstanding changes not yet made to Commission Implementing Regulation (EU) 2016/759. Any changes that have already been made to the legislation appear in the content and are referenced with annotations.