Export Health Certificate

	I.1. Consignor				I.2. IMSOC Reference			
	Name				I.2.a. Local Reference			
	Address							
	Country ISO Code							
	I.S. Consignee				I.3. Central competent authority			
nt	Name				I.4. Local competent authority			
me	Address		700.0.1					
g	Country		ISO Code					
consignment	I.7. Country of origin ISO Code				I.9. Country of destinatio	n	ISO Code	
ofc	I.8. Region of origin Code				I.10. Region of destination	n	-	
ls c	I.11. Place of Dispa	atch			I.12. Place of destination			
: Details	Name				Name			
۵	Address				Address			
	Approval Number				Approval Number	ISO Code		
Part I	Country		ISO Code		Country	150 Code		
4	I.13. Place of Load	ing			I.14. Date and time of dep	parture		
	Name							
	Address	_						
	Approval Number Country	Γ	ISO Code					
	-							
\dashv	I.15. Means of Tra Mode		T-1		I.16 Entry Point			
	Mode	International transport	Identification					
		document			_			
					_			
	I.18. Transport conditions Ambient Chilled Controlled Frozen				I.17. Accompanying documents Accompanying document			
	Ambient \square	Chilled	temperature \square	Frozen 🗆	reference Date of issue Country			
					Place of issue			
	I.19. Container No	/ Seal No			1			
	I.20. Certified as							
	Human consumpt	ion 🗆						
	I.21. For transit th	rough a third cour	trv \Box		I.22. For transit through Member State(s)			
	I.21. For transit through a third country Country ISO Code					·		
	Country EU Exit				Country	ISO Code		
	Authority		BCP code		-			
	EU Entry Authority		BCP code		-			
	I.23. Total number	of packages	I.25	. Total net weight	I.25. Total gross weight			
	I.28. Description o	f consignment						
	-	•	MODIFIED STAR	CHES; GLUES; ENZY	MES			
	1. 35 ALBUMINOIDAL SUBSTANCES; MODIFIED STARCHES; GLUES; ENZYMES 3503 Gelatin (including gelatin in rectangular (including square) sheets, whether or not surface-worked or coloured) and gelatin derivatives; isinglass; other glues of animal origin, excluding casein glues of heading 3501							
						40117441700,		
	#1. Commodity Manufacturing plant		Cold store	Final consumer				
	Species		Package count		Net weight	Batch number		

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II. Health information

II.1. Public health attestation

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 laying down specific hygiene rules for food of animal origin (OJ L 139, 30.4.2004, p. 55) and Regulation (EU) 2017/625 of the European Parliament and of the Council of 15 March 2017 on official controls and other official activities performed to ensure the application of food and feed law, rules on animal health and welfare, plant health and plant protection products, amending Regulations (EC) No 999/2001, (EC) No 396/2005, (EC) No 1069/2009, (EC) No 1107/2009, (EU) No 1151/2012, (EU) No 652/2014, (EU) 2016/429 and (EU) 2016/2031 of the European Parliament and of the Council. Council Regulations (EC) No 1/2005 and (EC) No 1099/2009 and Council Directives 98/58/EC, 1999/74/EC, 2007/43/EC, 2008/119/EC and 2008/120/EC and repealing Regulations (EC) No 854/2004 and (EC) No 882/2004 of the European Parliament and of the Council, Council Directives 89/608/EEC, 89/662/EEC, 90/425/EEC, 91/496/EEC, 96/23/EC, 96/93/EC and 97/78/EC and Council Decision 92/438/EEC (Official Controls Regulation) (OJ L 95, 7.4.2017, p. 1), and

I certify that the gelatine described above was produced in accordance with these requirements, in particular that:

- it comes from (an) establishment(s) implementing a programme based on the hazard analysis and critical control points (HACCP) principles in accordance with Article 5 of Regulation (EC) No 852/2004;
- it has been produced from raw materials that met the requirements of Chapters I and II of Section XIV of Annex III to Regulation (EC) No 853/2004;
- it has been manufactured in compliance with the conditions set out in Chapter III of Section XIV of Annex III to Regulation (EC) No 853/2004;
- it satisfies the criteria of Chapter IV of Section XIV of Annex III to Regulation (EC) No 853/2004 and of Commission Regulation (EC) No 2073/2005 of 15 November 2005 on microbiological criteria for foodstuffs (OJ L 338, 22.12.2005, P. 1);

(1) and, if of bovine, ovine and caprine animal origin, it has been derived from animals which have passed ante-mortem and post-mortem inspections,

(1) and, except for gelatine derived from hides and skins,

$(1) \circ either$

- [it comes from a country or a region classified in accordance with Commission Decision 2007/453/EC of 29 June 2007 establishing the BSE status of Member States or third countries or regions thereof according to their BSE risk (OJ L 172, 30.6.2007, p. 84) as a country or region posing a negligible BSE risk;
- the gelatine does not contain and is not derived from 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 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) (2);
- the gelatine does not contain and is not derived from mechanically separated meat obtained from the bones of bovine, ovine or caprine animals, except for gelatine derived from animals that were born, continuously reared and slaughtered in a country or region classified in accordance with Decision 2007/453fEC as a country or region posing a negligible BSE risk in which there has been no indigenous BSE cases;
- the animals, from which the gelatine is derived were not slaughtered after stunning by means of gas injected into the cranial cavity or killed by the same method or slaughtered, after stunning, by laceration of central nervous tissue by means of an elongated rod-shaped instrument introduced into the cranial cavity, except if the animals were born, continuously reared and slaughtered in a country or region classified in accordance with Decision 2007/453/EC as a

EU	ROPEAN UNION	(GB) G	(GB) GEL Gelatine intended for human consumption / GBHC109E (v2.0)			
	II. Health information					
		,	country or region posing a negligible BSE risk;			
Part II: Certification		or region region po meat-and	(1) ☐ [the animals, from which the gelatine is derived, originate from a country or region classified in accordance with Decision 2007/453/EC as a country or region posing an undetermined BSE risk, and the animals were not fed with meat-and-bone meal or greaves, as defined in the Terrestrial Animal Health Code of the World Organisation for Animal Health];			
		or region region po handled i	canimals, from which the gelatine is derived, originate from a country classified in accordance with Decision 2007/453/EC as a country or sing an undetermined BSE risk, and the gelatine was produced and n a manner which ensures that it did not contain and was not ates with nervous and lymphatic tissues exposed during the deboning			
"	(1) ○ Or					
		Decision States or	[it comes from a country or a region classified in accordance with Commission Decision 2007/453/EC of 29 June 2007 establishing the BSE status of Member States or third countries or regions thereof according to their BSE risk (OJ L 172, 30.6.2007, p. 84) as a country or region posing a controlled BSE risk;			
		by lacera instrume	e animals, from which the gelatine is derived, were not killed, after stunning, a laceration of central nervous tissue by means of an elongated rod-shaped strument introduced into the cranial cavity, or by means of gas injected into e cranial cavity;			
		defined in	the gelatine does not contain and is not derived from specified risk material as defined in point 1 of Annex V to Regulation (EC) No 999/2001, or mechanically separated meat obtained from the bones of bovine, ovine or caprine animals.]			
	(1) ○ Or					
		Decision : States or	[it comes from a country or a region classified in accordance with Commission Decision 2007/453/EC of 29 June 2007 establishing the BSE status of Member States or third countries or regions thereof according to their BSE risk (OJ L 172, 30.6.2007, p. 84) as a country or region posing an undetermined BSE risk;			
		meal or g	the animals, from which the gelatine is derived, were not fed meat-and-bone meal or greaves derived from ruminants, as defined in the Terrestrial Animal Health Code of the World Organisation for Animal Health;			
		by lacera instrume	the animals, from which the gelatine is derived, were not killed, after stunning, by laceration of 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;			
		- the gelati	ne is not derived from:			
		(i)	specified risk material as defined in point 1 of Annex V to Regulation (EC) No 999/2001;			
		(ii)	nervous and lymphatic tissues exposed during the deboning process;			
		(iii)	mechanically separated meat obtained from the bones of bovine, ovine or caprine animals			

(*) Those countries subject to the transitional import arrangements include: an EU member State; Liechtenstein; Norway; Iceland and Switzerland.

References to European Union legislation within this certificate are references to direct EU legislation which has been retained in Great Britain (retained EU law as defined in the European Union (Withdrawal) Act 2018).

References to Great Britain in this certificate include Channel Islands and Isle of Man.

See notes in Annex II of Commission Implementing Regulation (EU) 2019/628 of 8 April 2019 concerning model official certificates for certain animals and goods and amending Regulation (EC) No 2074/2005 and Implementing Regulation (EU) 2016/759 as regards these model certificates (OJ L 131, 17.5.2019, p. 101).

EUROPEAN UNION (GB) GEL Gelatine intended for human consumption / GBHC109E (v2.							
	II. Health info	rmation					
	- The colou	r of the star	mp and signature must be different	from that of the other particu	ılars in the certificate.		
	Part l:						
	raiti.	Dow	Do not use this how until the and or	f the tues sitional etering new	a d		
	-	Box reference	Do not use this box until the end of	i the transitional staging peri	oa.		
ᆸ		I.16:					
lti0	_	Box	Insert the appropriate Harmonised	d System (HS) code(s) using h	eadings such as 3503.		
lica		reference	inout the appropriate married	a o y o to iii (110) o o u o (0) u o 1116 11	04441.60 04011 40 00001		
Ï		I.25:					
ပ္	Part II:						
1 1	(1)	Delete as a	ppropriate.				
Part II: Certification	(2)		val of specified risk material is not re	amirad if the galatine is deriv	and from animals horn		
	(2)		sly reared and slaughtered in a third				
			e with Decision 2007/453/EC as posi				
	Certifying Offi	cer					
	Name (in capi	tal letters)		Qualification and title			
	Date of signati	are		Signature			
	Stamp						
	-						

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