

Part I : Details of consignment	I.1. Consignor Name Address Country		ISO Code		I.2. IMSOC Reference <b>Specimen not to be used for exports from EU</b>		I.2.a. Local Reference	
	I.5. Consignee Name Address Country		ISO Code		I.3. Central competent authority		I.4. Local competent authority	
	I.7. Country of origin		ISO Code		I.9. Country of destination		ISO Code	
	I.8. Region of origin		Code		I.10. Region of destination		Code	
	I.11. Place of Dispatch Name Address Approval Number Country		ISO Code		I.12. Place of destination Name Address Approval Number Country		ISO Code	
	I.13. Place of Loading Name Address Approval Number Country		ISO Code		I.14. Date and time of departure			
	I.15. Means of Transport				I.16 Entry Point			
	Mode	International transport document	Identification					
	I.18. Transport conditions Ambient <input type="checkbox"/> Controlled temperature <input type="checkbox"/> Frozen <input type="checkbox"/> Chilled <input type="checkbox"/>				I.17. Accompanying documents Commercial document reference Date of issue Country Place of issue			
	I.19. Container No / Seal No							
I.20. Certified as Transhumance <input type="checkbox"/> Pollination <input type="checkbox"/> Animal Feedingstuff <input type="checkbox"/> Pharmaceutical use <input type="checkbox"/> Game Restocking <input type="checkbox"/> Fattening <input type="checkbox"/> Registered equidae <input type="checkbox"/> Quarantine <input type="checkbox"/> Artificial reproduction <input type="checkbox"/> Training <input type="checkbox"/> Further process <input type="checkbox"/> Racing <input type="checkbox"/> Production of petfood <input type="checkbox"/> Breeding and production <input type="checkbox"/> Consignments according to Regulation No 999/2001 <input type="checkbox"/> Competition <input type="checkbox"/> Storage <input type="checkbox"/> Breeding <input type="checkbox"/> Ornamental use/research <input type="checkbox"/> Laboratory <input type="checkbox"/> Rodent food <input type="checkbox"/> Pets <input type="checkbox"/> Other <input type="checkbox"/> Production <input type="checkbox"/> Technical use <input type="checkbox"/> Ornamental bird food <input type="checkbox"/> Relaying <input type="checkbox"/> Slaughter <input type="checkbox"/> Approved Bodies <input type="checkbox"/> Human consumption <input type="checkbox"/> Pet food <input type="checkbox"/> Sales <input type="checkbox"/> Organic fertilizers <input type="checkbox"/> Unregistered equidae <input type="checkbox"/> Circus exhibition <input type="checkbox"/> Category 3 fish oil/fish meal for detoxification according to Regulation 2015/786 <input type="checkbox"/>								
I.21. For transit through a third country <input type="checkbox"/> Country _____ ISO Code _____ EU Exit Authority _____ BCP code _____ EU Entry Authority _____ BCP code _____				I.22. For transit through Member State(s) <input type="checkbox"/> Country _____ ISO Code _____				
I.23. Total number of packages		I.24. Total quantity		I.25. Total net weight		I.25. Total gross weight		

Part I : Details of consignment	I.28. Description of consignment			
	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			
	Commodity	Species	Quantity	Batch number
Cold store	Cutting plant	Date of freezing	Date of production	Date of slaughter
Net weight	Product Description	Package count	Identification mark	

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II. Health information		
Part II: Certification	I, the undersigned official, declare that I have read and understood Regulation (EC) No 1069/2009 of the European Parliament and of the Council and in particular Articles 8 and 10 thereof, and Commission Regulation (EU) No 142/2011, and in particular Annex XIV, Chapter II thereof, and certify that the photographic gelatine described above:	
	<ul style="list-style-type: none"><li>II.1. consists exclusively of photographic gelatine for photographic uses and is not intended for any other purpose;</li><li>II.2. has been prepared and stored in a plant registered and supervised by the competent authority in accordance with Article 23 of Regulation (EC) No 1069/2009, which does not produce gelatine for food, feed or other uses intended for dispatch to Great Britain, Channel Islands or Isle of Man;</li><li>II.3. has been prepared with Category 3 animal by-products and/or bovine vertebral column classified as Category 1 material;</li><li>II.4. has been wrapped, packaged in new containers, stored and transported in sealed, leak-proof labelled containers in a vehicle under satisfactory hygiene conditions;</li><li>II.5. has been produced by a process ensuring that the raw material is:<ul style="list-style-type: none"><li>(3) either ◦ [treated by pressure sterilisation as referred to in definition No 19 of Article 3 of Regulation (EC) No 1069/2009 (2);]</li><li>(3) or ◦ [subjected to:<ul style="list-style-type: none"><li>(i) treatment with acid for at least two days, washing with water and treatment with an alkaline solution for at least 20 days; the pH must be adjusted and material purified by means of filtration and sterilised at 138-140°C for 4 seconds; or</li><li>(ii) treatment with alkali for at least two days, washing with water and treatment with an acid solution for 10-12 hours; the pH must be adjusted and the material purified by means of filtration and sterilised at 138-140 °C for 4 seconds.]</li></ul></li></ul></li><li>II.6. has been wrapped and packaged in wrappings and packages carrying the words 'PHOTOGRAPHIC GELATINE FOR THE PHOTOGRAPHIC INDUSTRY ONLY'.</li></ul>	

Part II: Certification	II. Health information		
	Notes		
	(*) 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.		
	Part I:		
	-	Box reference I.5:	The intended destination of the photographic gelatine can only be Great Britain, Channel Islands or Isle of Man.
	-	Box reference I.9:	Country of destination: only applicable to Great Britain, Channel Islands or Isle of Man.
	-	Box reference I.11 and I.12:	Approval number: the registration number of the establishment or plant, which has been issued by the competent authority.
	-	Box reference I.15:	Registration number (railway wagons or container and lorries), flight number (aircraft) or name (ship); information is to be provided in the event of unloading and reloading.
-	Box reference I.16:	Do not use this box until the end of the transitional staging period.	
-	Box reference I.23:	Identification of container/seal number: only where applicable.	
-	Box reference I.25:	technical use: any use other than for animal consumption.	
Part II:			
(2)	Pressure sterilisation (method 1) is also referred to in Chapter III of Annex IV to Regulation (EU) No 142/2011 as follows:		
	‘Reduction		
	1.	If the particle size of the animal by-products to be processed is more than 50 millimetres, the animal by-products must be reduced in size using appropriate equipment, set so that the particle size after reduction is no greater than 50 millimetres. The effectiveness of the equipment must be checked daily and its condition recorded. If checks disclose the existence of particles larger than 50 millimetres, the process must be stopped and repairs made before the process is resumed.	
	Time, temperature and pressure		
	2.	The animal by-products with the particle size of no greater than 50 millimetres must be heated to a core temperature of more than 133 °C for at least 20 minutes without interruption at a pressure (absolute) of at least 3 bars. The pressure must be produced by the evacuation of all air in the sterilisation chamber and the replacement of the air by steam (“saturated steam”); the heat treatment may be applied as the sole process or as pre- or post-process sterilisation phase.	
	3.	The processing may be carried out in batch or continuous systems.’	
(3)	Delete as appropriate.		
-	The signature and stamp must be in a different colour to that of the printing.		

EUROPEAN UNION

Part II: Certification	II. Health information			
	- Note for the person responsible for the load in Great Britain, Channel Islands or Isle of Man: this certificate is only for veterinary purposes and has to accompany the consignment until it reaches the factory of destination from the border control post.			
	Certifying Officer			
	Name (in capital letters)	Qualification and title		
	Date of signature	Signature		
	Stamp			

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Část I	I.1. Odesílatel Název Adresa Země <span style="float: right;">Kód ISO</span>		I.2. Referenční číslo IMSOC <b>Specimen not to be used for exports from EU</b> I.2.a. Local Reference																																				
	I.5. Příjemce Název Adresa Země <span style="float: right;">Kód ISO</span>		I.3. Ústřední příslušný orgán I.4. Local competent authority																																				
	I.7. Země původu <span style="float: right;">Kód ISO</span>		I.9. Country of destination <span style="float: right;">Kód ISO</span>																																				
	I.8. Region of origin <span style="float: right;">Kód</span>		I.10. Region určení <span style="float: right;">Kód</span>																																				
	I.11. Place of Dispatch Název Adresa Číslo schválení Země <span style="float: right;">Kód ISO</span>		I.12. Místo určení Název Adresa Číslo schválení Země <span style="float: right;">Kód ISO</span>																																				
	I.13. Místo nakládky Název Adresa Číslo schválení Země <span style="float: right;">Kód ISO</span>		I.14. Date and time of departure																																				
	I.15. Dopravní prostředky <table border="1" style="width: 100%; border-collapse: collapse;"> <thead> <tr> <th style="width: 25%;">Typ</th> <th style="width: 25%;">Doklad</th> <th style="width: 50%;">Identifikace</th> </tr> </thead> <tbody> <tr><td> </td><td> </td><td> </td></tr> <tr><td> </td><td> </td><td> </td></tr> <tr><td> </td><td> </td><td> </td></tr> <tr><td> </td><td> </td><td> </td></tr> </tbody> </table>		Typ	Doklad	Identifikace													I.16 Entry Point																					
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I.19. Č. kontejneru / č. plomby																																							
I.20. Certified as <table style="width: 100%;"> <tr> <td style="width: 25%;">Transhumance <input type="checkbox"/></td> <td style="width: 25%;">Pollination <input type="checkbox"/></td> <td style="width: 25%;">Krmivo pro zvířata <input type="checkbox"/></td> <td style="width: 25%;">Použití pro farmaceutické účely <input type="checkbox"/></td> </tr> <tr> <td>Game Restocking <input type="checkbox"/></td> <td>Výkrmová <input type="checkbox"/></td> <td>Evidování koňovité <input type="checkbox"/></td> <td>Karanténa <input type="checkbox"/></td> </tr> <tr> <td>Umělé rozmnožování <input type="checkbox"/></td> <td>Training <input type="checkbox"/></td> <td>Další postup <input type="checkbox"/></td> <td>Racing <input type="checkbox"/></td> </tr> <tr> <td>Production of petfood <input type="checkbox"/></td> <td>Breeding and production <input type="checkbox"/></td> <td>Consignments according to Regulation No 999/2001 <input type="checkbox"/></td> <td>Competition <input type="checkbox"/></td> </tr> <tr> <td>Storage <input type="checkbox"/></td> <td>Breeding <input type="checkbox"/></td> <td>Ornamental use/research <input type="checkbox"/></td> <td>Laboratory <input type="checkbox"/></td> </tr> <tr> <td>Rodent food <input type="checkbox"/></td> <td>Zvířata v zájmovém chovu <input type="checkbox"/></td> <td>Jiné <input type="checkbox"/></td> <td>Production <input type="checkbox"/></td> </tr> <tr> <td>Technické použití <input type="checkbox"/></td> <td>Ornamental bird food <input type="checkbox"/></td> <td>Sádkování <input type="checkbox"/></td> <td>Porážka <input type="checkbox"/></td> </tr> <tr> <td>Schválené orgány <input type="checkbox"/></td> <td>Lidská spotřeba <input type="checkbox"/></td> <td>Krmivo pro domácí zvířata <input type="checkbox"/></td> <td>Sales <input type="checkbox"/></td> </tr> <tr> <td>Organic fertilizers <input type="checkbox"/></td> <td>Unregistered equidae <input type="checkbox"/></td> <td>Cirkus/výstava <input type="checkbox"/></td> <td>Category 3 fish oil/fish meal for detoxification according to Regulation 2015/786 <input type="checkbox"/></td> </tr> </table>				Transhumance <input type="checkbox"/>	Pollination <input type="checkbox"/>	Krmivo pro zvířata <input type="checkbox"/>	Použití pro farmaceutické účely <input type="checkbox"/>	Game Restocking <input type="checkbox"/>	Výkrmová <input type="checkbox"/>	Evidování koňovité <input type="checkbox"/>	Karanténa <input type="checkbox"/>	Umělé rozmnožování <input type="checkbox"/>	Training <input type="checkbox"/>	Další postup <input type="checkbox"/>	Racing <input type="checkbox"/>	Production of petfood <input type="checkbox"/>	Breeding and production <input type="checkbox"/>	Consignments according to Regulation No 999/2001 <input type="checkbox"/>	Competition <input type="checkbox"/>	Storage <input type="checkbox"/>	Breeding <input type="checkbox"/>	Ornamental use/research <input type="checkbox"/>	Laboratory <input type="checkbox"/>	Rodent food <input type="checkbox"/>	Zvířata v zájmovém chovu <input type="checkbox"/>	Jiné <input type="checkbox"/>	Production <input type="checkbox"/>	Technické použití <input type="checkbox"/>	Ornamental bird food <input type="checkbox"/>	Sádkování <input type="checkbox"/>	Porážka <input type="checkbox"/>	Schválené orgány <input type="checkbox"/>	Lidská spotřeba <input type="checkbox"/>	Krmivo pro domácí zvířata <input type="checkbox"/>	Sales <input type="checkbox"/>	Organic fertilizers <input type="checkbox"/>	Unregistered equidae <input type="checkbox"/>	Cirkus/výstava <input type="checkbox"/>	Category 3 fish oil/fish meal for detoxification according to Regulation 2015/786 <input type="checkbox"/>
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I.23. Celkový počet balení	I.24. Celkové množství	I.25. Celková čistá hmotnost	I.25. Celková hrubá hmotnost																																				

I.28. Description of consignment

**1. 35 ALBUMINOIDNÍ LÁTKY; MODIFIKOVANÉ ŠKROBY; KLIHY; ENZYMY****3503** Želatina (včetně želatiny v pravouhlých (včetně čtvercových) fóliích, též povrchově upravená nebo barvená) a deriváty želatiny; vyzina; jiné klišy živočišného původu, kromě kaseinových klišů čísla | 3501

Komodita	Druh	Množství	Číslo šarže	Výrobní zařízení
Chladírenské zařízení	Bourárna	Datum zmrazení	Datum výroby	Datum porážky
Čistá hmotnost	Product Description	Počet balení	Identifikační značka	

Část I

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II. Informace týkající se zdraví

I, the undersigned official, declare that I have read and understood Regulation (EC) No 1069/2009 of the European Parliament and of the Council and in particular Articles 8 and 10 thereof, and Commission Regulation (EU) No 142/2011, and in particular Annex XIV, Chapter II thereof, and certify that the photographic gelatine described above:

Part II: Certification

- II.1. consists exclusively of photographic gelatine for photographic uses and is not intended for any other purpose;
- II.2. has been prepared and stored in a plant registered and supervised by the competent authority in accordance with Article 23 of Regulation (EC) No 1069/2009, which does not produce gelatine for food, feed or other uses intended for dispatch to Great Britain, Channel Islands or Isle of Man;
- II.3. has been prepared with Category 3 animal by-products and/or bovine vertebral column classified as Category 1 material;
- II.4. has been wrapped, packaged in new containers, stored and transported in sealed, leak-proof labelled containers in a vehicle under satisfactory hygiene conditions;
- II.5. has been produced by a process ensuring that the raw material is:
- (3) either  [treated by pressure sterilisation as referred to in definition No 19 of Article 3 of Regulation (EC) No 1069/2009 (2);]
- (3) or  [subjected to:
- (i) treatment with acid for at least two days, washing with water and treatment with an alkaline solution for at least 20 days; the pH must be adjusted and material purified by means of filtration and sterilised at 138-140°C for 4 seconds; or
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- II.6. has been wrapped and packaged in wrappings and packages carrying the words 'PHOTOGRAPHIC GELATINE FOR THE PHOTOGRAPHIC INDUSTRY ONLY'.



II. Informace týkající se zdraví

## Notes

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

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

## Part I:

- Box reference I.5: The intended destination of the photographic gelatine can only be Great Britain, Channel Islands or Isle of Man.
- Box reference I.9: Country of destination: only applicable to Great Britain, Channel Islands or Isle of Man.
- Box reference I.11 and I.12: Approval number: the registration number of the establishment or plant, which has been issued by the competent authority.
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- Box reference I.23: Identification of container/seal number: only where applicable.
- Box reference I.25: technical use: any use other than for animal consumption.

## Part II:

- (2) Pressure sterilisation (method 1) is also referred to in Chapter III of Annex IV to Regulation (EU) No 142/2011 as follows:
- ‘Reduction
1. If the particle size of the animal by-products to be processed is more than 50 millimetres, the animal by-products must be reduced in size using appropriate equipment, set so that the particle size after reduction is no greater than 50 millimetres. The effectiveness of the equipment must be checked daily and its condition recorded. If checks disclose the existence of particles larger than 50 millimetres, the process must be stopped and repairs made before the process is resumed.
- Time, temperature and pressure
2. The animal by-products with the particle size of no greater than 50 millimetres must be heated to a core temperature of more than 133 °C for at least 20 minutes without interruption at a pressure (absolute) of at least 3 bars. The pressure must be produced by the evacuation of all air in the sterilisation chamber and the replacement of the air by steam (“saturated steam”); the heat treatment may be applied as the sole process or as pre- or post-process sterilisation phase.
  3. The processing may be carried out in batch or continuous systems.’
- (3) Delete as appropriate.
- The signature and stamp must be in a different colour to that of the printing.

II. Informace týkající se zdraví

- Note for the person responsible for the load in Great Britain, Channel Islands or Isle of Man: this certificate is only for veterinary purposes and has to accompany the consignment until it reaches the factory of destination from the border control post.

Certifying Officer

Name (in capital letters)

Qualification and title

Datum podpisu

Podpis

Razítko

Part II: Certification

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