

Part I : Details of consignment	I.1. Consignor		I.2. IMSOC Reference		
	Name		I.2.a. Local Reference		
	Address				
	Country	ISO Code			
	I.5. Consignee		I.3. Central competent authority		
	Name		I.4. Local competent authority		
	Address				
	Country	ISO Code			
	I.7. Country of origin		ISO Code	I.9. Country of destination	
				ISO Code	
I.8. Region of origin		Code	<del>I.10. Region of destination</del>		
I.11. Place of Dispatch		I.12. Place of destination			
Name		Name			
Address		Address			
Approval Number		Approval Number			
Country	ISO Code	Country		ISO Code	
I.13. Place of Loading		I.14. Date and time of departure			
Name					
Address					
Approval Number					
Country	ISO Code				
I.15. Means of Transport		I.16 Entry Point			
Mode	International transport document	Identification			
I.18. Transport conditions		I.17. Accompanying documents			
Chilled <input type="checkbox"/> Controlled temperature <input type="checkbox"/> Frozen <input type="checkbox"/> Ambient <input type="checkbox"/>		Commercial document reference		Date of issue	
		Country		Place of issue	
I.19. Container No / Seal No					
I.20. Certified as Human consumption <input type="checkbox"/>					
I.21. For transit through a third country <input type="checkbox"/>		I.22. For transit through Member State(s) <input type="checkbox"/>			
Country	ISO Code	Country		ISO Code	
EU Exit 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 of consignment					
<b>1. 35 ALBUMINOIDAL SUBSTANCES; MODIFIED STARCHES; GLUES; ENZYMES</b> <b>3502</b> Albumins (including concentrates of two or more whey proteins, containing by weight more than 80   % whey proteins, calculated on the dry matter), albuminates and other albumin derivatives <b>350220</b> Milk albumin, including concentrates of two or more whey proteins Other: <b>35022091</b> Dried (for example, in sheets, scales, flakes, powder)					
Commodity	Species	Product Description	Batch number	Date of manufacture	
Expiration Date	Package count	Manufacturing plant	Net weight		

Part II: Certification	II. Health information		
	<p>II. Health information</p> <p>The health authority, represented by the undersigned official veterinarian, certifies that:</p> <p>II.1. The products have been produced in an establishment that has been subject to health approval by the competent authority and applies the Hazard Analysis and Critical Control Points (HACCP) system, Good Manufacturing Practice (GMP) programmes and Sanitation Standard Operating Procedures (SSOP).</p> <p>II.2. The products comply with the industrialisation process specifications to ensure the health control of food-safety critical points and the inactivation of pathogens for animal health.</p> <p>The milk was subjected to one of the following treatments or equivalent:</p> <p>(either) (1) ○ [II.2.1. High-Temperature Short-Time (HTST) pasteurisation at at least 72° C for at least 15 seconds if the pH is less than 7]</p> <p>(or) (1) ○ [II.2.2. High-Temperature Short-Time (HTST) pasteurisation on two (2) consecutive occasions if the pH is higher than or equal to 7]</p> <p>(or) (1) ○ [II.2.3. Slow pasteurisation at a temperature of at least 63°C for at least 30 minutes]</p> <p>(or) (1) ○ [II.2.4. An ultra-high temperature (UHT) treatment at not less than 135° C in combination with a suitable holding time.]</p> <p>(or) (1) ○ [II.2.5. A HTST treatment combined with another physical treatment by either: lowering the pH below 6 for one hour or additional heating equal to or greater than 72°C combined with desiccation]</p> <p>○ II.2.6. Dairy products derived from raw milk(2):</p> <p>II.2.6.1. The milk used to make the product comes from herds free from Brucellosis and Tuberculosis.</p> <p>II.2.6.2. The product has undergone a maturing process of at least 60 days at a temperature of 2°C or above.</p> <p>II.3. They are fit for human consumption.</p> <p>II.4. Additional animal health attestation: the products comply with the requirements mentioned below:</p> <p>II.4.1. They come from herds and primary production establishments that were not subject to health restrictions at the time of the milk collection.</p> <p>II.4.2. The primary production establishment and the area of at least 10 km surrounding it have not been under quarantine or subject to animal movement restrictions in the sixty (60) days prior to dispatch.</p> <p>II.4.3. The product was subject to an identity check at the place of loading.</p> <p>II.4.4. The necessary precautions have been taken after treatment to avoid contact of the milk or its products with any micro-organism that is potentially pathogenic to animals that cause notifiable infectious or contagious diseases according to the OIE list.</p>		
	<p>Notes</p> <p>Part I</p> <p>(1) Delete what does not apply.</p> <p>(2) Only applicable to countries that are free from foot-and-mouth disease.</p> <p>- The signature and the stamp must be in a different colour to that of the printing.</p> <p>- The certificate must be issued in Spanish and in the language of the EU Member State, on paper with the letterheads, logos and stamps of the issuing health authority.</p>		
	<p>Certifying Officer</p> <p>Name (in capital letters)</p> <p>Date of signature</p> <p>Stamp</p>	<p>Qualification and title</p> <p>Signature</p>	