

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			I.9. Country of destination		
	ISO Code			ISO Code		
	I.8. Region of origin			I.10. Region of destination		
	Code					
	I.11. Place of Dispatch			I.12. Place of destination		
	Name			Name		
Address			Address			
Approval Number			Approval Number			
Country			Country			
ISO Code			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			
Ambient <input type="checkbox"/> Chilled <input type="checkbox"/> Frozen <input type="checkbox"/> Controlled temperature <input type="checkbox"/>			Commercial document reference			
			Date of issue			
			Country			
			Place of issue			
I.19. Container No / Seal No						
I.20. Certified as						
Production of petfood <input type="checkbox"/>		Production <input type="checkbox"/>		Technical use <input type="checkbox"/>		
Fattening <input type="checkbox"/>		Breeding <input type="checkbox"/>		Animal Feedingstuff <input type="checkbox"/>		
Human consumption <input type="checkbox"/>		Other <input type="checkbox"/>		Pharmaceutical use <input type="checkbox"/>		
Relaying <input type="checkbox"/>				Breeding and production <input type="checkbox"/>		
				Slaughter <input type="checkbox"/>		
				Artificial reproduction <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.24. Total quantity		I.25. Total gross weight		
I.28. Description of consignment						
1. 16 PREPARATIONS OF MEAT, OF FISH OR OF CRUSTACEANS, MOLLUSCS OR OTHER AQUATIC INVERTEBRATES						
1602 Other prepared or preserved meat, meat offal or blood						
160290 Other, including preparations of blood of any animal						
Commodity	Species	Quantity	Net weight	Package count		
Identification number			Identification system			

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Part II: Certification	II. Health information								
	<p>I, the undersigned state/official veterinarian certify <input type="checkbox"/> [that the certificate is based on the following pre-export certificates (see attached list in case more than two)(1):</p> <table border="0"> <tr> <td>Date:</td> <td>Number:</td> <td>Country of origin:</td> <td>Administrative territory:</td> <td>Approval number of the Establishment:</td> <td>Name and quantity (net weight) of the product:</td> </tr> </table>				Date:	Number:	Country of origin:	Administrative territory:	Approval number of the Establishment:
Date:	Number:	Country of origin:	Administrative territory:	Approval number of the Establishment:	Name and quantity (net weight) of the product:				

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	II. Health information		
Part II: Certification	II.1.	Products, manufactured from meat, sub-products and fats of all animal species, poultry and other meat products, destined for human food were processed at establishments, approved by the Competent Veterinary Service in the EU for supplying their production for export and operating under its constant supervision.	
	II.2.	Raw materials from which the product is manufactured are obtained from clinically healthy animals which have been subjected to veterinary inspection prior to slaughter, their carcasses and internal organs - to post mortem veterinary- sanitary inspection, conducted by the State/official Veterinary Service.	
	II.3.	Beef and mutton from which canned meat, salamis and other ready for consumption meat products are manufactured are derived from animals that originate from herds where there is no case of Bovine spongiform encephalopathy (BSE) or scrapie respectively and do not belong to birth cohorts of BSE positive animals. The meat is obtained from animals which have been tested for BSE, with negative results, when they are over 72 months at slaughterhouse if no classical BSE case in animals younger than 5 years has been detected in the Member State over the last 3 years. In other cases, the meat is obtained from bovine animals which have been tested for BSE, with negative results, when they are over 48 months at slaughterhouse. Specified risk materials (SRM) were removed according to the OIE Code recommendations.	
	II.4.	Meat and meat products were obtained from the slaughtered animals, which were not fed by feed of animal origin, excluding milk proteins.	
	II.5.	Animals, from which meat is derived, were not subjected to the exposure of natural or synthetically estrogenic, hormonal substances, thyreostatics, antibiotics, other drugs and pesticides, used prior to slaughter no later than authorised by instructions on how to use them.	
	II.6.	Meat products are recognised fit for human consumption.	
	II.7.	Products originate from meat processing establishments or coldstores, in the administrative territory of the EU Member State free from the diseases appearing on list A in the OIE Code of 2003 and for which the species from which the product originates is susceptible, including(2): <ul style="list-style-type: none">- <input type="checkbox"/> African swine fever - during the last 3 years in the territory of the EU, excluding <input type="checkbox"/> Sardinia; the administrative territories envisaged by the applicable Commission Implementing Regulation of the EU introducing changes to Commission Implementing Regulation (EU) 2021/605;- <input type="checkbox"/> African swine fever - during the last 3 years in the territory of the EU excluding Sardinia . This product was treated using technologies that guarantees the destruction of the ASF virus according to Annex 1 to Rosselkhoznazor letter of 28.06.2021 No. FS-KS-7/18163; <ul style="list-style-type: none">· rinderpest during the last 12 months and foot-and-mouth diseases during the last 6 months in the territory of the EU Member State;	
	II.8.	Microbiological, chemical-toxicological and radiological characteristics of the product correspond to actual veterinary and sanitary rules and requirements of the Russian Federation.	
	II.9.	Products have identification marks.	
	II.10.	Single-use containers and packaging material correspond to hygienic requirements.	
	II.11.	Means of transport are treated and prepared in accordance with the rules approved in the EU.	
	Notes		
	Part I		
	<ul style="list-style-type: none">· Box I.6.: Pre-export certificates numbers.· Box I.11.: Place of origin: name, number and address of the dispatch establishment.· Box I.16.: Point of crossing the border of the Russian Federation.· Box I.18.: Temperature of storage and transport.· Box I.19.: State the total gross weight and total net weight.· Box I.25.: Identification of goods		
	Customs code and title: Use the appropriate Harmonised System (HS) code.		
	Manufacturing plant, cold store: State the name, address and the approval number of the manufacturing plant or cold store, if appropriate.		

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Part II: Certification	II. Health information		
	Part II		
	<ul style="list-style-type: none">· (1) Delete if not relevant and confirm by signature and stamp· (2) Administrative territories, zones and time periods may be modified with a mutual agreement on the basis of the Memorandum of 4 April 2006 on zoning and regionalisation.		
Signature and stamp must be in a different colour to that in the printed certificate			
Certifying Officer			
Name (in capital letters)		Qualification and title	
Date of signature		Signature	
Stamp			