

Country

Official certificate to the EU

Part I : Details of consignment	I.1. Consignor Name Address Country ISO Code		I.2. Certificate reference I.3. Central competent authority I.4. Local competent authority		I.2.a. IMSOC reference Specimen not to be used for imports into the EU																	
	I.5. Consignee Name Address Country ISO Code		I.6. Responsible for the consignment in EU Name Address Country ISO Code																			
	I.7. Country of origin	ISO Code	I.8. Region of origin	Code	I.9. Country of destination	ISO 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 Frozen <input type="checkbox"/> Chilled <input type="checkbox"/> Ambient <input type="checkbox"/>		I.17. Accompanying documents Type Number																			
I.19. Container No / Seal No																						
I.20. Certified as																						
Travelling circulations/animal acts <input type="checkbox"/>	Pharmaceutical use <input type="checkbox"/>	Quarantine establishments <input type="checkbox"/>	Technical use <input type="checkbox"/>	Artificial reproduction <input type="checkbox"/>	Products for human consumption <input type="checkbox"/>	Slughter <input type="checkbox"/>	Petfood <input type="checkbox"/>	Other <input type="checkbox"/>	Fattening <input type="checkbox"/>	Furher keeping <input type="checkbox"/>	Registered animal <input type="checkbox"/>	Canning industry <input type="checkbox"/>	Game resting <input type="checkbox"/>	Feed stuff <input type="checkbox"/>	Approved Bodies <input type="checkbox"/>	Manufacture of petfood <input type="checkbox"/>	Breeding/production <input type="checkbox"/>	Pets <input type="checkbox"/>	Production of petfood <input type="checkbox"/>	Trade samples <input type="checkbox"/>	Relaying area/purification centre <input type="checkbox"/>	Furher processing <input type="checkbox"/>
I.21. For transit <input type="checkbox"/> Non-EU ISO Code		I.22. For internal market <input type="checkbox"/>																				
I.24. Total number of packages		I.25. Total quantity		I.23. For re-entry <input type="checkbox"/>																		
I.24. Total number of packages		I.25. Total quantity		I.26. Total net weight		I.26. Total gross weight																
I.27. Description of consignment																						
<p>1. 23 RESIDUES AND WASTE FROM THE FOOD INDUSTRIES; PREPARED ANIMAL FODDER</p> <p>2309 Preparations of a kind used in animal feeding</p> <p>230910 Dog or cat food, put up for retail sale</p>																						
Commodity		Species		Nature of commodity		Manufacturing plant		Net weight														
Batch number										Package count												

II. Health information

Part II : Certification

SPECIMEN

Part II : Certification	II. Health information		
	<p>I, the undersigned official veterinarian, declare that I have read and understood Regulation (EC) No 1069/2009 of the European Parliament and of the Council (1a), and in particular Article 10 of that Regulation, and Commission Regulation (EU) No 142/2011 (1b), and in particular Chapter II of Annex XIII and Chapter II of Annex XIV thereto, and certify that the dogchews described above:</p>		
<p>II.1. have been prepared exclusively with the following animal by-products:</p>			
<p>(2) <input type="checkbox"/> either [- carcasses and parts of animals slaughtered or, in the case of game, bodies or parts of animals killed, and which are fit for human consumption in accordance with Union legislation, but are not intended for human consumption for commercial reasons;]</p>			
<p>(2) <input type="checkbox"/> and/or [- carcasses and the following parts originating either from animals that have been slaughtered in a slaughterhouse and were considered fit for slaughter for human consumption following an ante-mortem inspection or bodies and the following parts of animals from game killed for human consumption in accordance with Union legislation:</p> <p>(i) carcasses or bodies and parts of animals which are rejected as unfit for human consumption in accordance with Union legislation, but which did not show any signs of disease communicable to humans or animals;</p> <p>(ii) heads of poultry;</p> <p>(iii) hides and skins, including trimmings and splitting thereof, horns and feet, including the phalanges and the carpus and metacarpus bones, tarsus and metatarsus bones;</p> <p>(iv) pig bristles;</p> <p>(v) feathers;]</p>			
<p>(2) <input type="checkbox"/> and/or [- blood of animals which did not show any signs of disease communicable through blood to humans or animals, obtained from animals that have been slaughtered in a slaughterhouse after having been considered fit for slaughter for human consumption following an ante-mortem inspection in accordance with Union legislation;]</p>			
<p>(2) <input type="checkbox"/> and/or [- animal by-products arising from the production of products intended for human consumption, including degreased bone, greaves and centrifuge or separator sludge from milk processing;]</p>			
<p>(2) <input type="checkbox"/> and/or [- aquatic animals, and parts of such animals, except sea mammals, which did not show any signs of disease communicable to humans or animals;]</p>			
<p>(2) <input type="checkbox"/> and/or [- animal by-products from aquatic animals originating from plants or establishments manufacturing products for human consumption;]</p>			
<p>(2) <input type="checkbox"/> and/or [- material from animals which have been treated with certain substances which are prohibited by Council Directive 96/22/EC (2a), the import of the material being permitted in accordance with Article 35(a)(ii) of Regulation (EC) No 1069/2009;]</p>			
<p>II.2. have been subjected</p>			
<p>(2) <input type="checkbox"/> either [in the case of dogchews made from hides and skins of ungulates or from fish, to a treatment sufficient to destroy pathogenic organisms (including salmonella); and the dogchews are dry;]</p>			
<p>(2) <input type="checkbox"/> and/or [in the case of dogchews made from animal by-products other than hides and skins of ungulates or from fish, to a heat treatment of at least 90°C throughout their substance;]</p>			
<p>II.3. were examined by random sampling of at least five samples from each processed batch taken during or after storage at the processing plant and complies with the following standards (3):</p> <p>Salmonella: absence in 25g: n = 5, c = 0, m = 0, M = 0,</p> <p>Enterobacteriaceae: n = 5, c = 2, m = 10, M = 300 in 1 gramme;</p>			

Part II : Certification	II. Health information		
	II.4.	have undergone all precautions to avoid contamination with pathogenic agents after treatment;	
	II.5.	were packed in new packaging;	
	(2)	[II.6. the dogchews described above	
	(2)	○ either [is derived from other ruminants than bovine, ovine or caprine animals.]]	
	(2)	○ or [is derived from bovine, ovine or caprine animals and does not contain and is not derived from:	
	(2)	○ either [bovine, ovine and caprine materials other than those derived from animals born, continuously reared and slaughtered in a country or region classified as posing a negligible BSE risk in accordance with Decision 2007/453/EC.]]	
	(2)	○ or [(a) 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 (4);	
		(b) mechanically separated meat obtained from bones of bovine, ovine or caprine animals, except from those animals that were born, continuously reared and slaughtered in a country or region classified as posing a negligible BSE risk in accordance with Commission Decision 2007/453/EC (5), in which there has been no indigenous BSE case,	
		(c) animal by-product or derived product obtained from bovine, ovine or caprine animals which have been killed, after stunning, by laceration of the 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, except for those animals that were born, continuously reared and slaughtered in a country or region classified as posing a negligible BSE risk in accordance with Decision 2007/453/EC.]]]	

Part II : Certification	II. Health information			
	Notes			
	Part I:			
	Box reference I.6:	Person responsible for the consignment in the European Union: this box is to be filled in only if it is a certificate for transit commodity; it may be filled in if the certificate is for a commodity to be imported into the European Union.		
	Box reference I.12:	Place of destination: this box is to be filled in only if it is a certificate for a transit commodity. Products in transit may only be stored in free zones, free warehouses and custom warehouses.		
	Box reference I.15:	Registration number (railway wagons or container and lorries), flight number (aircraft) or name (ship); the information is to be provided in the event of unloading and reloading in the European Union.		
	Box reference I.19:	for bulk containers, the container number and the seal number (if applicable) must be given.		
	Box reference I.20:	technical use: any use other than feeding of farmed animals, other than fur animals, and the production or manufacturing of pet food.		
	Box reference I.21 and I.22:	fill in according to whether it is a transit or an import certificate.		
	Box reference I.25:	05.11, 23.09, 41.01 or 42.05.		
Box reference I.25:	Species: select from the following: Aves, Ruminantia, Suidae, Mammalia Other Than Ruminantia or Suidae, Pesca, Mollusca, Crustacea, Invertebrates Other Than Mollusca And Crustacea.			
Part II:				
(1a)	OJ L 300, 14.11.2009, p. 1.			
(1b)	OJ L 54, 26.2.2011, p. 1.			
(2)	Delete as appropriate.			
(2a)	OJ L 125, 23.5.1996, p. 3.			
(3)	Where:			
	n =	number of samples to be tested;		
	m =	threshold value for the number of bacteria; the result is considered satisfactory if the number of bacteria in all samples does not exceed m;		
	M =	maximum value for the number of bacteria; the result is considered unsatisfactory if the number of bacteria in one or more samples is M or more; and		
	c =	number of samples the bacterial count of which may be between m and M, the sample still being considered acceptable if the bacterial count of the other samples is m or less.		
(4)	OJ L 147, 31.5.2001, p. 1.			
(5)	OJ L 172, 30.6.2007, p. 84.			
-	The signature and the stamp must be in a different colour to that of the printing.			
-	Note for the person responsible for the consignment in the European Union: This certificate is only for veterinary purposes and must accompany the consignment until it reaches the border inspection post of entry into the European Union.			
Official veterinarian or Official inspector				
Name (in capital letters)		Qualification and title		
Date of signature		Signature		
Stamp				