CHAPTER 3(C)

Health certificate

For dogchews intended for ~~dispatch to or for~~ transit through[2] the European Union

COUNTRY: UNITED STATES **Veterinary certificate to EU**

Part I : Details of dispatched consignment

I.1. Consignor Name Address Tel.	I.2. Certificate reference No I.2.a. I.3. Central competent authority **APHIS-VS** I.4. Local competent authority
I.5. ~~Consignee~~ Border inspection post through which consignment is intended to leave the EU Name Address ~~Postal code~~ ~~Tel.~~	I.6. Person responsible for the load in EU Name Address Postal code Tel.
I.7. Country of origin: **US** ISO code: **US-0** I.8. Region of origin Code	I.9. Country of destination ISO code I.10. Region of destination Code
I.11. Place of origin Name Approval number Address Name Approval number Address Name Approval number Address	I.12. Place of destination Custom warehouse ☐ Approval number Name Address Postal code
I.13. Place of loading	I.14. Date of departure
I.15. Means of transport Aeroplane ☐ Ship ☐ Railway wagon ☐ Road vehicle ☐ Other ☐ Identification Documentation references	I.16. Entry BIP in EU I.17.
I.18. Description of commodity	I.19. Commodity code (HS code) **42.05.00** I.20. Quantity
I.21. Temperature of product Ambient ☐ Chilled ☐ Frozen ☐	I.22. Number of packages

COUNTRY: UNITED STATES **Dogchews**

	I.2. Certificate reference No / I.2.a.
I.23. Seal/Container No	I.24. Type of packaging
I.25. Commodities certified for: Animal feedingstuff ☒ Technical use ☐	
I.26. For transit through EU to third country ☒ Third country ISO code	I.27. For import or admission into EU ☐

I.28. Identification of the commodities

Species (Scientific name)	Approval number of establishments Manufacturing plant	Net weight	Batch number

II.	**Health information**	II.a. Certificate reference No	II.b.

Part II: Certification

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, and Commission Regulation (EU) No 142/2011[1b], and in particular Annex XIII, Chapter II and Annex XIV, Chapter II thereof, and certify that the dogchews described above:

II.1. have been prepared exclusively with the following animal by-products:

[2]~~either~~ [- ~~carcases 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;]~~

[2]~~and/or~~ [- ~~carcases 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:~~

~~(i) carcases 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;~~

~~(ii) heads of poultry;~~

~~(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, of animals other than ruminants;~~

~~(iv) pig bristles;~~

~~(v) feathers;]~~

[2]~~and/or~~ [- ~~blood of animals which did not show any signs of disease communicable through blood to humans or animals, obtained from animals other than ruminants 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;]~~

[2]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;]

[2]~~and/or~~ [- ~~animal by-products from aquatic animals originating from plants or establishments manufacturing products for human consumption;]~~

[2]~~and/or~~ [- ~~material from animals which have been treated with certain substances which are prohibited pursuant to Directive 96/22/EC, the import of the material being permitted in accordance with Article 35(a)(ii) of Regulation (EC) No 1069/2009;]~~

II.2. have been subjected

[2]~~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;]~~

[2]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;]

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]:

Salmonella:	absence in 25g: n = 5, c = 0, m = 0, M = 0,
Enterobacteriaceae:	n = 5, c = 2, m = 10, M = 300 in 1 gram;

II.4. have undergone all precautions to avoid contamination with pathogenic agents after treatment;

II.5. were packed in new packaging;

II.6.

[2]either [the product does not contain and is not derived from specified risk material as defined in Annex V to Regulation (EC) No 999/2001 of the European Parliament and of the Council[4] or mechanically separated meat obtained from bones of bovine, ovine or caprine animals; and the animals from which this product is derived have not been slaughtered after stunning by means of gas injected into the cranial cavity or killed by the same method or slaughtered by laceration of central nervous tissue by means of an elongated rod-shaped instrument introduced into the cranial cavity;]

[2]~~or~~ ~~[the product does not contain and is not derived from bovine, ovine or 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 by a decision in accordance with Article 5(2) of Regulation (EC) No 999/2001;]~~

II.7. in addition as regards TSE:

II. Health information	II.a. Certificate reference No	II.b.

~~(2)either [in case of animal by-products intended for feeding ruminants and containing milk or milk products of ovine or caprine origin, the ovine and caprine animals from which these products are derived have been kept continuously since birth or for the last three years on a holding where no official movement restriction is imposed due to a suspicion of TSE and which has satisfied the following requirements for the last three years:~~

~~(i) it has been subject to regular official veterinary checks;~~

~~(ii) no classical scrapie case, as defined in point 2(g) of Annex I to Regulation (EC) No 999/2001, has been diagnosed or, following the confirmation of a classical scrapie case:~~

~~— all animals in which classical scrapie was confirmed have been killed and destroyed, and~~

~~— all goats and sheep on the holding have been killed and destroyed, except for breeding rams of the ARR/ARR genotype and breeding ewes carrying at least one ARR allele and no VRQ allele;~~

~~(iii) ovine and caprine animals, with the exception of sheep of the ARR/ARR prion genotype, are introduced into the holding only if they come from a holding which complies with the requirements set out in points (i) and (ii).]~~

~~(2)or [in case of animal by-products intended for feeding ruminants and containing milk or milk products of ovine or caprine origin, and destined to a Member State listed in the Annex to Commission Regulation (EC) No 546/2006(5), the ovine and caprine animals from which these products are derived have been kept continuously since birth or for the last seven years on a holding where no official movement restriction is imposed due to a suspicion of TSE and which has satisfied the following requirements for the last seven years:~~

~~(i) it has been subject to regular official veterinary checks;~~

~~(ii) no classical scrapie case, as defined in point 2(g) of Annex I to Regulation (EC) No 999/2001, has been diagnosed or, following the confirmation of a classical scrapie case:~~

~~— all animals in which classical scrapie was confirmed have been killed and destroyed, and~~

~~— all goats and sheep on the holding have been killed and destroyed, except for breeding rams of the ARR/ARR genotype and breeding ewes carrying at least one ARR allele and no VRQ allele;~~

~~(iii) ovine and caprine animals, with the exception of sheep of the ARR/ARR prion genotype, are introduced into the holding only if they come from a holding which complies with the requirements set out in points (i) and (ii).]~~

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 import commodity.
— Box reference I.12: Place of destination: this box is to be filled in only if it is a certificate for transit commodity. The products in transit can 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); information is to be provided in the event of unloading and reloading.
— Box reference I.19: Alternatively, commodity codes 2309 and 4101 may be chosen.
— Box reference I.23: for bulk containers, the container number and the seal number (if applicable) should be given.
— Box reference I.25: technical use: any use other than for animal consumption.
— Box reference I.26 and I.27: fill in according to whether it is a transit or an import certificate.

Part II:

(1a) OJ L 300, 14.11.2009, p. 1.
(1b) OJ L 54, 26.2.2011, p. 1.
(2) Delete as appropriate.
(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 94, 1.4.2006, p. 28.

— 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 has to accompany the consignment until it reaches the border inspection post.

Official veterinarian/Official inspector

Name (in capital letters): Qualification and title:

Date: Signature:

Stamp: