COMMISSION IMPLEMENTING REGULATION (EU) 2019/628

of 8 April 2019

concerning model official certificates for certain animals and goods and amending Regulation (EC) No 2074/2005 and Implementing Regulation (EU) 2016/759 as regards these model certificates

(Text with EEA relevance)

THE EUROPEAN COMMISSION,

Having regard to the Treaty on the Functioning of the European Union,

Having regard to Regulation (EU) 2017/625 of the European Parliament and of the Council of 15 March 2017 on official controls and other official activities performed to ensure the application of food and feed law, rules on animal health and welfare, plant health and plant protection products, amending Regulations (EC) No 999/2001, (EC) No 396/2005, (EC) No 1069/2009, (EC) No 1107/2009, (EU) No 1151/2012, (EU) No 652/2014, (EU) 2016/429 and (EU) 2016/2031 of the European Parliament and of the Council, Council Regulations (EC) No 1/2005 and (EC) No 1099/2009 and Council Directives 98/58/EC, 1999/74/EC, 2007/43/EC, 2008/119/EC and 2008/120/EC and repealing Regulations (EC) No 854/2004 and (EC) No 882/2004 of the European Parliament and of the Council, Council Directives 89/608/EEC, 89/662/EEC, 90/425/EEC, 91/496/EEC, 96/23/EC, 96/93/EC and 97/78/EC and Council Decision 92/438/EEC (Official Controls Regulation) (1), and in particular points (a), (c) and (e) of the first paragraph of Articles 90 and 126(3) thereof,

Whereas:

- Regulation (EU) 2017/625 lays down rules for official controls and other control activities performed by the (1)competent authorities of the Member States in order to verify compliance with Union legislation in the area of, among others, food safety at all stages of the production, processing and distribution process. In particular, it provides for official certification when considered appropriate to ensure compliance with EU rules on animals and goods.
- (2)Point (a) of the first paragraph of Article 90 of Regulation (EU) 2017/625 empowers the Commission to adopt, by means of implementing acts, rules concerning model official certificates and rules for the issuance of such certificates, where requirements are not laid down in the rules referred to in Article 1(2) of that Regulation.
- (3) Consignments of animals and goods shall be accompanied by an official certificate issued either on paper or in electronic form. Therefore, it is appropriate to establish common requirements as regards issuance of official certificates in both cases in addition to the requirements laid down in Chapter VII of Title II of Regulation (EU) 2017/625.
- (4) Model certificates are included in the electronic system Traces, set up by Commission Decision 2003/623/EC (2), to facilitate and accelerate administrative procedures at Union borders and to enable electronic communication between the competent authorities which helps preventing possible fraudulent or deceptive practices in respect of the official certificates.
- Since 2003, computer technology has evolved considerably and the Traces system has been modified to improve (5) the quality, processing and secure exchange of data. As a result, the format of the model certificates and the notes on their completion laid down in this Regulation should be adapted to the Traces system, for example by reflecting the use of multiple Combined Nomenclature (CN) codes or providing traceability for triangular trade, where the country of dispatch is not the country of origin of the consignment.
- In accordance with Article 133(4) of Regulation (EU) 2017/625, the Traces system is to be integrated into the (6) Information Management System for Official Controls (IMSOC). The model health certificates laid down in this Regulation should therefore be adapted to IMSOC.
- (7) Point (c) of the first paragraph of Article 90 of Regulation (EU) 2017/625 empowers the Commission to lay down by means of implementing acts rules concerning the procedures to be followed for the issuance of replacement certificates.

 ^{(&}lt;sup>1</sup>) OJ L 95, 7.4.2017, p. 1.
 (²) Commission Decision 2003/623/EC of 19 August 2003 concerning the development of an integrated computerised veterinary system known as Traces (OJ L 216, 28.8.2003, p. 58).

- To avoid misuse and abuse, it is important to define the cases where a replacement certificate may be issued and (8)the requirements that need to be met by such certificates. In particular, these cases should be limited to obvious administrative errors, such as transposed numbers in the container number or seal number, spelling errors in addresses or in product descriptions.
- (9) Article 126(2)(c) of Regulation (EU) 2017/625 establishes the requirement that consignments of certain animals and goods are to be accompanied by an official certificate, an official attestation or any other evidence that the consignment complies with the applicable rules referred to in Article 1(2) of that Regulation.
- (10)Commission Delegated Regulation (EU) 2019/625 (3) provides for a list of goods and animals intended for human consumption, in particular products of animal origin, live insects and sprouts and seeds intended for the production of sprouts, that need to be accompanied by an official certificate upon the entry into the Union if intended for placing on the market. To facilitate official controls upon the entry into the Union, model official certificates should be laid down for such goods and animals intended for human consumption in accordance with Point (a) of the first paragraph of Article 90 and Article 126(3) of Regulation (EU) 2017/625.
- (11)Model certificates required for public health reasons are currently laid down in various legal acts. It is appropriate to consolidate these model certificates in one single legal act by making cross-references to them.
- With respect to certification of certain products of animal origin for animal health reasons, common model (12)certificates are used. The requirements for certification for animal health reasons should be revised by 21 April 2021, which is the date of application of Regulation (EU) 2016/429 of the European Parliament and of the Council (*). The common model certificates should be maintained until that revision.
- For reasons of harmonisation and clarity, model certificates currently laid down in Commission Regulation (EC) (13)No 2074/2005 (5), Commission Regulation (EU) No 211/2013 (6) and Commission Implementing Regulation (EU) 2016/759 (7) should be incorporated into this Regulation. As a result, Regulation (EC) No 2074/2005 and Implementing Regulation (EU) 2016/759 should be amended accordingly and Regulation (EU) No 211/2013 should be repealed.
- (14)To facilitate the verification of compliance with EU requirements, it seems appropriate to introduce additional new model health certificates for the entry of rendered animal fats and greaves, insects and reptile meat intended for placing on the market. Such model certificates also make it easier for competent authorities in third countries to understand EU requirements and therefore facilitate the entry of animal fats and greaves, insects and reptile meat into the Union.
- Point (e) of the first paragraph of Article 90 of Regulation (EU) 2017/625 empowers the Commission to adopt, (15)by means of implementing acts, rules concerning the format of documents that are to accompany animals or goods after official controls have been performed. In accordance with Article 5 of Commission Delegated Regulation (EU) 2019/624 (8), such health certificates are to accompany animals to the slaughterhouse after ante-mortem inspection has been carried at the holding of provenance. The format of such certificates should therefore be laid down in this Regulation.

⁽³⁾ Commission Delegated Regulation (EU) 2019/625 of 4 March 2019 supplementing Regulation (EU) 2017/625 of the European Parliament and of the Council with regard to requirements for the entry into the Union of consignments of certain animals and goods intended for human consumption (see page 18 of this Official Journal).

Regulation (EU) 2016/429 of the European Parliament and of the Council of 9 March 2016 on transmissible animal diseases and amending and repealing certain acts in the area of animal health ('Animal Health Law') (OJ L 84, 31.3.2016, p. 1).

Commission Regulation (EC) No 2074/2005 of 5 December 2005 laying down implementing measures for certain products under Regulation (EC) No 853/2004 of the European Parliament and of the Council and for the organisation of official controls under Regulation (EC) No 854/2004 of the European Parliament and of the Council and Regulation (EC) No 882/2004 of the European Parliament and of the Council, derogating from Regulation (EC) No 852/2004 of the European Parliament and of the Council and amending Regulations (EC) No 853/2004 and (EC) No 854/2004 (OJ L 338, 22.12.2005, p. 27). Commission Regulation (EU) No 211/2013 of 11 March 2013 on certification requirements for imports into the Union of sprouts and

seeds intended for the production of sprouts (OJ L 68, 12.3.2013, p. 26).

Commission Implementing Regulation (EU) 2016/759 of 28 April 2016 drawing up lists of third countries, parts of third countries and territories from which Member States are to authorise the introduction into the Union of certain products of animal origin intended for human consumption, laying down certificate requirements, amending Regulation (EC) No 2074/2005 and repealing Decision 2003/812/EC (OJ L 126, 14.5.2016, p. 13). Commission Delegated Regulation (EU) 2019/624 of 8 February 2019 concerning specific rules for the performance of official controls

on the production of meat and for production and relaying areas of live bivalve molluscs in accordance with Regulation (EU) 2017/625 of the European Parliament and of the Council (see page 1 of this Official Journal).

- (16) In the case of emergency slaughter outside the slaughterhouse, it is appropriate for reasons of harmonisation and clarity, to lay down a model certificate in this Regulation for the declaration to be issued by the (official) veterinarian in accordance with point 6 of Chapter VI of Section I of Annex III of Regulation (EC) No 853/2004 of the European Parliament and of the Council (⁹).
- (17) As Regulation (EU) 2017/625 applies with effect from 14 December 2019, this Regulation should also apply from that date.
- (18) It is appropriate to introduce a transitional period to take into account the consignments of animals and goods shipped and certified, if required, before the date of application of this Regulation.
- (19) The measures provided for in this Regulation are in accordance with the opinion of the Standing Committee on Plants, Animals, Food and Feed,

HAS ADOPTED THIS REGULATION:

Article 1

Subject matter and scope

- 1. This Regulation lays down:
- (a) rules for the uniform application of Articles 88 and 89 of Regulation (EU) 2017/625 as regards the signature and issuance of official certificates and the guarantees of reliability for official certificates, in order to comply with the requirements of Article 126(2)(c) of that Regulation;
- (b) requirements for model official certificates which are not submitted in IMSOC;
- (c) requirements for model official certificates which are submitted in IMSOC;
- (d) requirements for replacement certificates.
- 2. This Regulation also sets out:
- (a) model official certificates for the entry into the Union of animals, products of animal origin, composite products, germinal products and animal by-products and notes for its completion;
- (b) specific model official certificates for the entry into the Union of the following animals and goods intended for human consumption and placing on the market:
 - (i) products of animal origin for which such certificate is required in accordance with Article 13 of Delegated Regulation (EU) 2019/625;
 - (ii) live insects;
 - (iii) sprouts and seeds intended for the production of sprouts;
- (c) model official certificates in the case of ante-mortem inspection at the holding of provenance or in the case of emergency slaughter outside the slaughterhouse.

Article 2

Definitions

For the purpose of this Regulation, the following definitions shall apply:

(1) 'placing on the market' means placing on the market as defined in point (8) of Article 3 of Regulation (EC) No 178/2002 of the European Parliament and of the Council (¹⁰);

^(*) Regulation (EC) No 853/2004 of the European Parliament and of the Council of 29 April 2004 laying down specific hygiene rules for food of animal origin (OJ L 139, 30.4.2004, p. 55).

^{(&}lt;sup>10</sup>) Regulation (EC) No 178/2002 of the European Parliament and of the Council of 28 January 2002 laying down the general principles and requirements of food law, establishing the European Food Safety Authority and laying down procedures in matters of food safety (OJ L 31, 1.2.2002, p. 1).

- (2) 'sprouts' means sprouts as defined in point (a) of Article 2 of Commission Implementing Regulation (EU) No 208/2013 (¹¹);
- (3) 'slaughterhouse' means a slaughterhouse as defined in point 1.16 of Annex I to Regulation (EC) No 853/2004;
- (4) 'fresh meat' means fresh meat as defined in point 1.10 of Annex I to Regulation (EC) No 853/2004;
- (5) 'meat' means meat as defined in point 1.1 of Annex I to Regulation (EC) No 853/2004;
- (6) 'poultry' means poultry as defined in point 1.3 of Annex I to Regulation (EC) No 853/2004;
- (7) 'wild game' means wild game as defined in point 1.5 of Annex I to Regulation (EC) No 853/2004;
- (8) 'eggs' means eggs as defined in point 5.1 of Annex I to Regulation (EC) No 853/2004;
- (9) 'egg products' means egg products as defined in point 7.3 of Annex I to Regulation (EC) No 853/2004;
- (10) 'meat preparations' means meat preparations as defined in point 1.15 of Annex I to Regulation (EC) No 853/2004;
- (11) 'meat products' means meat products as defined in point 7.1 of Annex I to Regulation (EC) No 853/2004;
- (12) 'treated stomachs, bladders and intestines' means treated stomachs, bladders and intestines as defined in point 7.9 of Annex I to Regulation (EC) No 853/2004;
- (13) 'bivalve molluscs' means bivalve molluscs as defined in point 2.1 of Annex I to Regulation (EC) No 853/2004;
- (14) 'fishery products' means fishery products as defined in point 3.1 of Annex I to Regulation (EC) No 853/2004;
- (15) 'raw milk' means raw milk as defined in point 4.1 of Annex I to Regulation (EC) No 853/2004;
- (16) 'dairy products' means dairy products as defined in point 7.2. of Annex I to Regulation (EC) No 853/2004;
- (17) 'colostrum' means colostrum as defined in point 1 of Section IX of Annex III of Regulation (EC) No 853/2004;
- (18) 'colostrum-based products' means colostrum-based products as defined in points 2 of Section IX of Annex III of Regulation (EC) No 853/2004;
- (19) 'frogs' legs' means frogs' legs as defined in point 6.1 of Annex I to Regulation (EC) No 853/2004;
- (20) 'snails' means snails as defined in point 6.2 of Annex I to Regulation (EC) No 853/2004;
- (21) 'rendered animal fat' means rendered animal fat defined in point 7.5 of Annex I to Regulation (EC) No 853/2004;
- (22) 'greaves' means greaves as defined in point 7.6 of Annex I to Regulation (EC) No 853/2004;
- (23) 'gelatine' means gelatine as defined in point 7.7 of Annex I to Regulation (EC) No 853/2004;
- (24) 'collagen' means collagen as defined in point 7.8 of Annex I to Regulation (EC) No 853/2004;
- (25) 'honey' means honey as defined in point 1 of Part IX of Annex II to Regulation (EU) No 1308/2013 of the European Parliament and of the Council (¹²);

 ^{(&}lt;sup>11</sup>) Commission Implementing Regulation (EU) No 208/2013 of 11 March 2013 on traceability requirements for sprouts and seeds intended for the production of sprouts (OJ L 68, 12.3.2013, p. 16).
 (¹²) Regulation (EU) No 1308/2013 of the European Parliament and of the Council of 17 December 2013 establishing a common

^{(&}lt;sup>12</sup>) Regulation (EU) No 1308/2013 of the European Parliament and of the Council of 17 December 2013 establishing a common organisation of the markets in agricultural products and repealing Council Regulations (EEC) No 922/72, (EEC) No 234/79, (EC) No 1037/2001 and (EC) No 1234/2007 (OJ L 347, 20.12.2013, p. 671).

- (26) 'apiculture products' means apiculture products as defined in point 2 of Part IX of Annex II to Regulation (EU) No 1308/2013;
- (27) 'reptile meat' means reptile meat as defined in point (16) of Article 2 of Delegated Regulation (EU) 2019/625;
- (28) 'insects' means insects as defined in point (17) of Article 2 of Delegated Regulation (EU) 2019/625;
- (29) 'reefer vessel' means a reefer vessel as defined in point (26) of Article 2 of Delegated Regulation (EU) 2019/625;
- (30) 'freezer vessel' means a freezer vessel as defined in point 3.3 of Annex I to Regulation (EC) No 853/2004;
- (31) 'factory vessel' means a factory vessel as defined in point 3.2 of Annex I to Regulation (EC) No 853/2004;
- (32) 'production area' means a production area as defined in point 2.5 of Annex I to Regulation (EC) No 853/2004;
- (33) 'dispatch centre' means a dispatch centre as defined in point 2.7 of Annex I to Regulation (EC) No 853/2004;
- (34) 'mechanically separated meat' means mechanically separated meat as defined in point 1.14 of Annex I to Regulation (EC) No 853/2004;
- (35) 'game-handling establishment' means a game-handling establishment as defined in point 1.18 of Annex I to Regulation (EC) No 853/2004;
- (36) 'cutting plant' means a cutting plant as defined in point 1.17 of Annex I to Regulation (EC) No 853/2004;
- (37) 'farmed game' means farmed game as defined in point 1.6 of Annex I to Regulation (EC) No 853/2004.

Article 3

Requirements for model official certificates not submitted in IMSOC

The model official certificates for those animals, products of animal origin, composite products, germinal products, animal by-products, sprouts and seeds intended for the production of sprouts originating from third countries or regions thereof which are required by Union legislation for the entry into the Union and are not submitted in IMSOC, shall meet the following requirements:

- (1) In addition to the signature of the certifying officer, the certificate shall bear an official stamp. The colour of signature shall be different to the colour of the printing. This requirement also applies to stamps other than those embossed or watermarked.
- (2) Where the model certificate contains statements, the statements which are not relevant shall be crossed out, initialled and stamped by the certifying officer, or completely removed from the certificate.
- (3) The certificate shall consist of:
 - (a) a single sheet of paper; or
 - (b) several sheets of paper where all sheets are indivisible and constitute an integral whole; or
 - (c) a sequence of pages numbered so as to indicate that it is a particular page in a finite sequence.
- (4) Where the certificate consists of a sequence of pages, each page shall indicate the unique code as referred to in Article 89(1)(a) of Regulation (EU) 2017/625 and bear the signature of the certifying officer and the official stamp.
- (5) The certificate shall be issued before the consignment to which it relates leaves the control of the competent authorities of the third country issuing the certificate.

Requirements for model official certificates submitted in IMSOC

1. The model official certificates for the entry into the Union of animals, products of animal origin, composite products, germinal products and animal by-products originating from third countries or regions thereof, submitted in IMSOC, shall be based on the model official certificate laid down in Annex I.

2. Part II of the model official certificates referred to in paragraph 1 shall include the specific health guarantees and the information as required in Part II of the relevant model official certificates for those animals, products of animal origin, composite products, germinal products and animal by-products originating from third countries or regions thereof which are required by Union legislation for the entry into the Union.

3. The official certificate shall be submitted in IMSOC before the consignment to which it relates leaves the control of the competent authorities of the third country issuing the certificate.

4. The requirements laid down in this Article shall not affect the nature, content and format of the official certificates or attestations referred to in Article 73(2)(b) and (c) and Article 129(2)(a) of Regulation (EU) 2017/625.

Article 5

Replacement certificates

1. Competent authorities may issue a replacement certificate only in the case of administrative errors in the initial certificate or where the initial certificate has been damaged or lost.

2. The replacement certificate shall not modify information in the initial certificate concerning the identification, traceability and health guarantees of consignments.

- 3. In addition, the replacement certificate shall:
- (a) make clear reference to the unique code referred to in Article 89(1)(a) of Regulation (EU) 2017/625 and the date of issue of the initial certificate, and clearly state that it replaces the initial certificate;
- (b) have a new certificate number different to that of the initial certificate;
- (c) carry the date when it was issued, as opposed to the date of issue of the initial certificate; and
- (d) be presented in its original to the competent authorities, except in the case of electronic replacement certificates submitted in IMSOC.

Article 6

Notes on the completion of model official certificates

The model official certificates referred to in Articles 12, 13 and 15 to 27 shall be completed on the basis of the notes set out in Annex II.

Article 7

Model official certificates for the entry into the Union for placing on the market of fresh meat of ungulates

To meet the certification requirements laid down in Articles 88, 89 and Article 126(2)(c) of Regulation (EU) 2017/625, the model official certificates 'BOV', 'OVI', 'POR', 'EQU', 'RUF', 'RUW', 'SUF', 'SUW' and 'EQW' set out in Part 2 of Annex II to Commission Regulation (EU) No 206/2010 (¹³) shall be used for the entry into the Union for placing on the market of fresh meat of ungulates.

^{(&}lt;sup>13</sup>) Commission Regulation (EU) No 206/2010 of 12 March 2010 laying down lists of third countries, territories or parts thereof authorised for the introduction into the European Union of certain animals and fresh meat and the veterinary certification requirements (OJ L 73, 20.3.2010, p. 1).

Model official certificates for the entry into the Union for placing on the market of meat of poultry, ratites and wild game birds, eggs and egg products

To meet the certification requirements laid down in Articles 88, 89 and Article 126(2)(c) of Regulation (EU) 2017/625, the model official certificates 'POU', 'POU-MI/MSM', 'RAT', 'RAT-MI/MSM', 'WGM', 'WGM-MI/MSM', 'E' and 'EP' set out in Part 2 of Annex I to Commission Regulation (EC) No 798/2008 (14) shall be used for the entry into the Union for placing on the market of meat of poultry, ratites and wild game birds, egg and egg products.

Article 9

Model official certificates for the entry into the Union for placing on the market of meat of wild leporidae, of certain wild land mammals and of farmed rabbits

To meet the certification requirements laid down in Articles 88, 89 and Article 126(2)(c) of Regulation (EU) 2017/625, the model official certificates 'WL', 'WM' and 'RM' set out in Annex II to Commission Regulation (EC) No 119/2009 (15) shall be used for the entry into the Union for placing on the market of meat of wild leporidae, of certain wild land mammals and of farmed rabbits.

Article 10

Model official certificate for the entry into the Union for placing on the market of meat preparations

To meet the certification requirements laid down in Articles 88, 89 and Article 126(2)(c) of Regulation (EU) 2017/625, the model official certificate set out in Annex II to Commission Decision 2000/572/EC (16) shall be used for the entry into the Union for placing on the market of meat preparations.

Article 11

Model official certificates for the entry into the Union for placing on the market of certain meat products and treated stomachs, bladders and intestines

To meet the certification requirements laid down in Articles 88, 89 and Article 126(2)(c) of Regulation (EU) 2017/625, the model official certificate set out in Annex III to Commission Decision 2007/777/EC (17) shall be used for the entry into the Union for placing on the market of certain meat products and treated stomachs, bladders and intestines. However, in the case of the entry into the Union for placing on the market of casings, the animal health certificate set out in Annex I A to Commission Decision 2003/779/EC (18) shall be used.

Article 12

Model official certificates for the entry into the Union for placing on the market of live bivalve molluscs, echinoderms, tunicates and marine gastropods

To meet the certification requirements laid down in Articles 88, 89 and Article 126(2)(c) of Regulation (EU) 2017/625, the model official certificate set out in Chapter A of Part I of Annex III to this Regulation shall be used for the entry into the Union for placing on the market of live bivalve molluscs, echinoderms, tunicates and marine gastropods. In the

⁽¹⁴⁾ Commission Regulation (EC) No 798/2008 of 8 August 2008 laying down a list of third countries, territories, zones or compartments from which poultry and poultry products may be imported into and transit through the Community and the veterinary certification requirements (OJ L 226, 23.8.2008, p. 1). ⁽¹⁵⁾ Commission Regulation (EC) No 119/2009 of 9 February 2009 laying down a list of third countries or parts thereof, for imports into,

or transit through, the Community of meat of wild leporidae, of certain wild land mammals and of farmed rabbits and the veterinary certification requirements (OJ L 39, 10.2.2009, p. 12).

⁽¹⁶⁾ Commission Decision 2000/572/EC of 8 September 2000 laying down the animal and public health and veterinary certification (1⁷) Commission Decision 2007/777/EC of 29 November 2007 laying down the animal and public health conditions and model certificates

for imports of certain meat products and treated stomachs, bladders and intestines for human consumption from third countries and repealing Decision 2005/432/EC (OJ L 312, 30.11.2007, p. 49). (¹⁸) Commission Decision 2003/779/EC of 31 October 2003 laying down animal health requirements and veterinary certification for the

import of animal casings from third countries (OJ L 285, 1.11.2003, p. 38).

case of the entry into the Union and placing on the market of processed bivalve molluscs belonging to the species *Acanthocardia tuberculatum*, the model official certification set out in Chapter B of Part I of Annex III to this Regulation shall be added to the certificate referred to in the first sentence.

Article 13

Model official certificates for the entry into the Union for placing on the market of fishery products

1. To meet the certification requirements laid down in Articles 88, 89 and Article 126(2)(c) of Regulation (EU) 2017/625, the model official certificate set out in Chapter A of Part II of Annex III to this Regulation shall be used for the entry into the Union for placing on the market of fishery products.

2. In the case of fishery products caught by vessels flying the flag of a Member State and transferred in third countries with or without storage, the model certificate set out in Chapter B of Part II of Annex III to this Regulation shall be used.

3. To meet the certification requirements laid down in Articles 88, 89 and Article 126(2)(c) of Regulation (EU) 2017/625, the model official certificate to be signed by the captain, set out in Chapter C of Part II to Annex III to this Regulation shall be used when fishery products are imported directly from a reefer, freezer or factory vessel as provided for in Article 11(3) of Delegated Regulation (EU) 2019/625.

Article 14

Model official certificates for the entry into the Union for placing on the market of raw milk, colostrum, dairy products and colostrum-based products

To meet the certification requirements laid down in Articles 88, 89 and Article 126(2)(c) of Regulation (EU) 2017/625, the model official certificates 'Milk-RM', 'Milk-RMP', 'Milk-HTB', 'Milk-HTC' and 'Colostrum-C/CPB' set out in Part 2 of Annex II to Commission Regulation (EU) No 605/2010 (¹⁹) shall be used for the entry into the Union for placing on the market of raw milk, colostrum, dairy products and colostrum-based products.

Article 15

Model official certificate for the entry into the Union for placing on the market of chilled, frozen or prepared frogs' legs intended for human consumption

To meet the certification requirements laid down in Articles 88, 89 and Article 126(2)(c) of Regulation (EU) 2017/625, the model official certificate set out Part III of Annex III to this Regulation shall be used for the entry into the Union for placing on the market of chilled, frozen or prepared frogs' legs intended for human consumption.

Article 16

Model official certificate for the entry into the Union for placing on the market of chilled, frozen, shelled, cooked, prepared or preserved snails intended for human consumption

To meet the certification requirements laid down in Articles 88, 89 and Article 126(2)(c) of Regulation (EU) 2017/625, the model official certificate set out in Part IV of Annex III to this Regulation shall be used for the entry into the Union for placing on the market of chilled, frozen, shelled, cooked, prepared or preserved snails intended for human consumption.

^{(&}lt;sup>19</sup>) Commission Regulation (EU) No 605/2010 of 2 July 2010 laying down animal and public health and veterinary certification conditions for the introduction into the European Union of raw milk, dairy products, colostrum and colostrum-based products intended for human consumption (OJ L 175, 10.7.2010, p. 1).

Model official certificate for the entry into the Union for placing on the market of rendered animal fats and greaves intended for human consumption

To meet the certification requirements laid down in Articles 88, 89 and Article 126(2)(c) of Regulation (EU) 2017/625, the model official certificate set out in Part V of Annex III to this Regulation shall be used for the entry into the Union for placing on the market of rendered animal fats and greaves intended for human consumption.

Article 18

Model official certificate for the entry into the Union for placing on the market of gelatine intended for human consumption

To meet the certification requirements laid down in Articles 88, 89 and Article 126(2)(c) of Regulation (EU) 2017/625, the model official certificate set out in Part VI of Annex III to this Regulation shall be used for the entry into the Union for placing on the market of gelatine intended for human consumption.

Article 19

Model official certificate for the entry into the Union for placing on the market of collagen intended for human consumption

To meet the certification requirements laid down in Articles 88, 89 and Article 126(2)(c) of Regulation (EU) 2017/625, the model official certificate set out in Part VII of Annex III to this Regulation shall be used for the entry into the Union for placing on the market of collagen intended for human consumption.

Article 20

Model official certificate for the entry into the Union for placing on the market of raw materials for the production of gelatine and collagen intended for human consumption

To meet the certification requirements laid down in Articles 88, 89 and Article 126(2)(c) of Regulation (EU) 2017/625, the model official certificate set out in Part VIII of Annex III to this Regulation shall be used for the entry into the Union for placing on the market of raw materials for the production of gelatine and collagen intended for human consumption.

Article 21

Model official certificate for the entry into the Union for placing on the market of treated raw materials for the production of gelatine and collagen intended for human consumption

To meet the certification requirements laid down in Articles 88, 89 and Article 126(2)(c) of Regulation (EU) 2017/625, the model official certificate set out in Part IX of Annex III to this Regulation shall be used for the entry into the Union for placing on the market of treated raw materials for the production of gelatine and collagen intended for human consumption.

Article 22

Model official certificate for the entry into the Union for placing on the market of honey and other apiculture products intended for human consumption

To meet the certification requirements laid down in Articles 88, 89 and Article 126(2)(c) of Regulation (EU) 2017/625, the model official certificate set out in Part X of Annex III to this Regulation shall be used for the entry into the Union for placing on the market of honey and other apiculture products intended for human consumption.

Model official certificate for the entry into the Union for placing on the market of highly refined chondroitin sulphate, hyaluronic acid, other hydrolysed cartilage products, chitosan, glucosamine, rennet, isinglass and amino acids intended for human consumption

To meet the certification requirements laid down in Articles 88, 89 and Article 126(2)(c) of Regulation (EU) 2017/625, the model official certificate set out in Part XI of Annex III to this Regulation shall be used for the entry into the Union for placing on the market of highly refined chondroitin sulphate, hyaluronic acid, other hydrolysed cartilage products, chitosan, glucosamine, rennet, isinglass and amino acids intended for human consumption.

Article 24

Model official certificate for the entry into the Union for placing on the market of reptile meat intended for human consumption

To meet the certification requirements laid down in Articles 88, 89 and Article 126(2)(c) of Regulation (EU) 2017/625, the model official certificate set out in Part XII of Annex III to this Regulation shall be used for the entry into the Union for placing on the market of reptile meat intended for human consumption.

Article 25

Model official certificate for the entry into the Union for placing on the market of insects intended for human consumption

To meet the certification requirements laid down in Articles 88, 89 and Article 126(2)(c) of Regulation (EU) 2017/625, the model official certificate set out in Part XIII of Annex III to this Regulation shall be used for the entry into the Union for placing on the market of insects intended for human consumption.

Article 26

Model official certificate for the entry into the Union for placing on the market of other products of animal origin intended for human consumption and not covered by Articles 7 to 25

To meet the certification requirements laid down in Articles 88, 89 and Article 126(2)(c) of Regulation (EU) 2017/625, the model official certificate set out in Part XIV of Annex III to this Regulation shall be used for the entry into the Union for placing on the market of other products of animal origin intended for human consumption and not covered by Articles 7 to 25 of this Regulation.

Article 27

Model official certificate for the entry into the Union for placing on the market of sprouts and seeds intended for the production of sprouts

To meet the certification requirements laid down in Articles 88, 89 and Article 126(2)(c) of Regulation (EU) 2017/625, the model official certificate set out in Part XV of Annex III to this Regulation shall be used for the entry into the Union for placing on the market of sprouts and seeds intended for the production of sprouts.

Article 28

Model official certificates in case of ante-mortem inspection at the holding of provenance

To meet the certification requirements laid down in Articles 88, 89 and Article 126(2)(c) of Regulation (EU) 2017/625, the model official certificates set out in Annex IV to this Regulation shall be used in the case of ante-mortem inspection at the holding of provenance in accordance with Articles 5 and 6 of Delegated Regulation (EU) 2019/624.

Model official certificate in case of emergency slaughter outside the slaughterhouse

To meet the certification requirements laid down in Articles 88, 89 and Article 126(2)(c) of Regulation (EU) 2017/625, the model official certificate set out Annex V to this Regulation shall be used in the case of emergency slaughter outside the slaughterhouse in accordance with Article 4 of Delegated Regulation (EU) 2019/624.

Article 30

Amendments to Regulation (EC) No 2074/2005

Regulation (EC) No 2074/2005 is amended as follows:

(1) Article 6 is deleted;

(2) Annex VI is deleted.

Article 31

Amendments to Implementing Regulation (EU) 2016/759

Implementing Regulation (EU) 2016/759 is amended as follows:

(1) Article 2 is deleted;

(2) Annex II is deleted.

Article 32

Repeal

Regulation (EU) No 211/2013 is repealed. References to Regulation (EU) No 211/2013 shall be construed as references to this Regulation and read in accordance with the correlation table set out in Annex VI to this Regulation.

Article 33

Transitional provisions

Consignments of products of animal origin accompanied by the relevant certificates issued accordance with Regulation (EC) No 2074/2005, Regulation (EU) No 211/2013 and Implementing Regulation (EU) 2016/759 may be accepted for the entry into the Union until 13 March 2020 provided that the certificate was signed before 14 December 2019.

Until 13 March 2020, consignments of rendered animal fats and greaves may enter the Union when using the certificate for meat products set out in Annex III to Decision 2007/77/EC and consignments of reptile meat, insects and other products of animal origin referred to in Article 26 may enter the Union without certificate set out in Annex III of this Regulation.

Article 34

Entry into force and application

This Regulation shall enter into force on the twentieth day following that of its publication in the Official Journal of the European Union.

It shall apply from 14 December 2019.

This Regulation shall be binding in its entirety and directly applicable in all Member States.

Done at Brussels, 8 April 2019.

For the Commission The President Jean-Claude JUNCKER

ANNEX I

MODEL OFFICIAL CERTIFICATES FOR THE ENTRY INTO THE UNION OF ANIMALS, PRODUCTS OF ANIMAL ORIGIN, COMPOSITE PRODUCTS, GERMINAL PRODUCTS AND ANIMAL BY-PRODUCTS

cou	OUNTRY							Official certificate to the EU				
	l.1.	Consignor/Exporter						I.2.	Certificate referen	ce No	I.2.a IMSOC reference	ce No
		Name						1.3.	I.3. Central Competent Authority			
		Address						1.4.	I.4. Local Competent Authority			
nent		Tel. No										
nsignr	l.5.	Consignee/Importer						I.6. Operator responsible for the consignment				
ed co		Name							Name			
patch		Address							Address			
of dis		Postal code							Postal code			
ails c	Tel. No											
Part I: Details of dispatched consignment	1.7.	Country of origin	ISO	I.8. Re	gion	of origin	Code	1.9.	Country of destination	ISO	I.10. Region of destination	Code
å	l.11	.11 Place of dispatch						Place of destination	on			
		Name		Approva	l No				Name			
		Address							Address			
	l.13.	Place of loading						I.14. Date and time of departure				
	l.15.	Means of transport						l.16.	Entry BCP			
		Aeroplane	Ves	ssel C		Other		l.17.	Accompanying do	cuments		
		Road vehicle	Rai	Iway [Туре			
	Identification:							No				
	I.18. Transport conditions											
		Ambient	Chi	lled [Frozen						
	l.19.	Container No/Seal No)			1		<u> </u>				

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COUNTRY							Official certificate	to the EU	
I.20. Goods certified as Canning industry Animal feedingstu Human consumpt Breeding/product Game restocking	uff tion ion	Fattening Quarantine Further process Slaughter Artificial reprodu	Ction	Approv Relayir	aceutical use red body		Trade samples Circus/exhibition Pets Other		
I.21. For transit				1.22. F	or internal mark	et			
Third country		ISO		R	efinitive import e-entry emporary admis	ssion			
I.23. Total number of p	I.24. Quantity Total numb	ber	Total n	Total net weight (Kg)		Total gross weight ((Kg)		
I.25. Description of goo	ods			_					
No	Code and CN	title							
Species (scie Ag	-	Bre	eed/Categor Sex	У	Identificatior Quant	-	Identificatio Test		
Species (Scie	entific name)		Nature of commodity					Treatment type	
Zone		Ma	anufactur	ring plant		Cold sto	ore		
Final consumer	Number o package		Net weight		Batch	No	Type of pac	kaging	
	Stamp			Signature					

COUNTRY

EN

Certificate model (**)

	II.	Health information (*)	II.a.	Certificate reference No	II.b.	IMSOC reference No
;						
-						
_						
	Certi	fying officer				
	Nam	e (in capital letters)		Qualification and title		
	Date			Signature		
	Stan	qr				
	<i>(</i>), -	pecify sanitary requirement to be completed				

(**) To be replaced by the specific title of each model of certificate

ANNEX II

NOTES ON THE COMPLETION OF THE MODEL OFFICIAL CERTIFICATES FOR THE ENTRY INTO THE UNION OF ANIMALS, PRODUCTS OF ANIMAL ORIGIN, COMPOSITE PRODUCTS, GERMINAL PRODUCTS, AND ANIMAL BY-PRODUCTS

General

To positively select any option, please tick or mark the relevant box with a cross (X).

Whenever mentioned, 'ISO' means the international standard two-letter code for a country, in accordance with the international standard ISO 3166 alpha-2 (¹).

Only one of the options may be selected in boxes I.15, I.18, I.20 and I.22.

If the consignee, the entry border control post (BCP) or the transport details (that is to say, the means and date) change after the certificate has been issued, the operator responsible for the consignment must advise the competent authority of the Member State of entry. Such a change shall not result in a request for a replacement certificate.

Part I: Details of the dispatched consignment

- Country: The name of the third country issuing the certificate.
- Box I.1. Consignor/Exporter: the name and address (street, city and region, province or state, as appropriate) of the natural or legal person dispatching the consignment that must be located in the third country, except for the re-entry of consignments originating from the European Union.
- Box I.2. Certificate reference No: the unique mandatory code assigned by the competent authority of the third country in accordance with its own classification. This box is compulsory for all certificates not submitted in IMSOC.
- Box I.2.a IMSOC reference No: the unique reference code automatically assigned by IMSOC, if the certificate is registered in IMSOC. This box must not be completed if the certificate is not submitted in IMSOC.
- Box I.3. Central competent authority: name of the central authority in the third country issuing the certificate.
- Box I.4. Local competent authority: if applicable, the name of the local authority in the third country issuing the certificate.
- Box I.5. Consignee/Importer: name and address of the natural or legal person to whom the consignment is intended in the Member State or third country of destination in the case of transit. However, this information is not compulsory for consignments in transit through the European Union.
- Box I.6. Operator responsible for the consignment:

The name and address of the person in the European Union in charge of the consignment when presented to the BCP and who makes the necessary declarations to the competent authorities either as the importer or on behalf of the importer.

For products in transit through the European Union: the name and address are compulsory.

For certain animals: the name and address are compulsory if required by the relevant European Union legislation.

For animals and products for the placing on the market: the name and address are optional.

Box I.7. Country of origin:

For products: the name and ISO code of the country where the goods were produced, manufactured and packaged (labelled with the identification mark).

⁽¹⁾ List of country names and code elements under: http://www.iso.org/iso/country_codes/iso-3166-1_decoding_table.htm

For animals: the country of residence during the required period as set out in the relevant European Union health certificate. For registered horses re-entering the European Union, the country of origin means the country from which they were last consigned.

In the case of trade involving more than one third country (triangular trade), a separate certificate must be completed for each country of origin.

- Box I.8. Region of origin: if applicable, for animals or products affected by the regionalisation measures in accordance with European Union legislation. The code of approved regions, zones or compartments must be stated as defined in the relevant European Union legislation.
- Box I.9. Country of destination: the name and ISO code of the European Union country of destination of the animals or products.

If the products are in transit, the name and ISO code of the third country of destination is required.

- Box I.10. Region of destination: see box I.8.
- Box I.11. Place of dispatch: the name, address and approval number, if required by the European Union legislation, of the holdings or establishments from which the animals or the products come from.

For animals: a holding or any other officially monitored agricultural, industrial or commercial establishment, including zoos, amusement parks, wildlife and hunting reserves, where animals are regularly kept or bred.

For germinal products: semen collection or storage centres, or embryo collection or production teams.

For other products: any unit of a company in the food or animal by-product sector. Only the establishment shipping the products is to be named. In the case of trade involving more than one third country (triangular trade), the place of dispatch is the last third-country establishment of the export chain from which the final consignment is transported to the European Union.

Box I.12. Place of destination:

Except in the case of storage of products in transit, this information is optional.

For the placing on the market: the place where the animals or products are sent for final unloading. Give the name, address and approval number of the holdings or establishments of the place of destination, if applicable.

For storage of products in transit: the name, address and approval number of the warehouse in a free zone, the customs warehouse or the ship supplier.

Box I.13. Place of loading:

For animals: the name of the city or the place where the animals are loaded and if they are assembled beforehand, the details of the official assembly centre.

For products: the name of the city and category (for example, establishment, holding, port or airport) of the final place where the products are to be loaded in the means of transport for the journey to the European Union. In the case of a container, state where it is to be placed aboard the final means of transport to the European Union. In the case of a ferry, indicate the place where the truck embarked.

Box I.14. Date and time of departure:

For animals: the date and time at which the animals are scheduled to leave in their means of transport (aeroplane, vessel, railway or road vehicle).

For products: the date when the means of transport departs (aeroplane, vessel, railway or road vehicle).

Box I.15. Means of transport: means of transport leaving the country of dispatch.

Mode of transport: aeroplane, vessel, railway, road vehicle or other. 'Other' means modes of transport not covered by Council Regulation (EC) No 1/2005 (²).

⁽²⁾ Council Regulation (EC) No 1/2005 of 22 December 2004 on the protection of animals during transport and related operations and amending Directives 64/432/EEC and 93/119/EC and Regulation (EC) No 1255/97 (OJ L 3, 5.1.2005, p. 1).

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Identification of the means of transport: for aeroplanes the flight number, for vessels the ship name(s), for railways the train identity and wagon number, for road transports the registration number plate with trailer number plate if applicable.

In the case of a ferry, state the identification of the road vehicle, the registration number plate with trailer number plate if applicable, and the name of the scheduled ferry.

- Box I.16. Entry BCP: state the name of the BCP and its identification code assigned by IMSOC.
- Box I.17. Accompanying documents:

The type and reference number of document must be stated when a consignment is accompanied by the other documents such as CITES permit, permit for invasive alien species (IAS) or a commercial document (for example, the airway bill number, the bill of lading number or the commercial number of the train or road vehicle)

- Box I.18. Transport conditions: category of required temperature during the transport of products (ambient, chilled, frozen). Only one category may be selected.
- Box I.19. Container No/Seal No: if applicable, the corresponding numbers.

The container number must be provided if the goods are transported in closed containers.

Only the official seal number must be stated. An official seal applies if a seal is affixed to the container, truck or rail wagon under the supervision of the competent authority issuing the certificate.

Box I.20. Goods certified as: state the purpose for the placing on the market of the animals or intended use for products as specified in the relevant European Union health certificate.

Animal feedingstuffs: concerns only animal by-products intended for animal feed as referred to in Regulation (EC) No 1069/2009 of the European Parliament and of the Council (³).

Approved body: movement of animals to an approved body, an institute or a centre in accordance with Council Directive 92/65/EEC (⁴).

Artificial reproduction: concerns only germinal products.

Breeding/production: for breeding and production animals, including aquaculture animals intended for farming.

Canning industry: concerns, for example, tuna intended for the canning industry.

Circus/exhibition: for registered circus and exhibition animals and aquatic animals for aquariums or similar businesses not for further sale.

Fattening: concerns ovine and caprine animals only.

Further process: concerns only products which have to be further processed before being placed on the market.

Game restocking: concerns only game for the purpose of rebuilding stocks.

Human consumption: concerns only products intended for human consumption for which a health or veterinary certificate is required by European Union legislation.

Other: intended for purposes not listed elsewhere in this classification, including aquatic animals intended for put-and-take fisheries.

 ^{(&}lt;sup>3</sup>) Regulation (EC) No 1069/2009 of the European Parliament and of the Council of 21 October 2009 laying down health rules as regards animal by-products and derived products not intended for human consumption and repealing Regulation (EC) No 1774/2002 (Animal by-products Regulation) (OJ L 300, 14.11.2009, p. 1).
 (⁴) Council Directive 92/65/EEC of 13 July 1992 laying down animal health requirements governing trade in and imports into the

^(*) Council Directive 92/65/EEC of 13 July 1992 laying down animal health requirements governing trade in and imports into the Community of animals, semen, ova and embryos not subject to animal health requirements laid down in specific Community rules referred to in Annex A (I) to Directive 90/425/EEC (OJ L 268, 14.9.1992, p. 54).

Pets: commercial movements into the Union of dogs, cats, ferrets and birds. For ornamental aquatic animals intended for pet shops or similar businesses for further sale.

Pharmaceutical use: animal by-products unfit for human or animal consumption, as referred to in Regulation (EC) No 1069/2009.

Quarantine: refers to Commission Implementing Regulation (EU) No 139/2013 (5) for birds other than poultry, to Directive 92/65/EEC for carnivores, primates and bats, and to Council Directive 2006/88/EC (6) for aquaculture animals.

Registered equidae: in accordance with Council Directive 2009/156/EC (7).

Relaying: concerns only aquaculture animals.

Slaughter: for animals destined directly or via an assembly centre to a slaughterhouse.

Technical use: animal by-products unfit for human or animal consumption, as referred to in Regulation (EC) No 1069/2009.

Trade samples: as defined in point 39 of Annex I to Commission Regulation (EU) No 142/2011 (8).

- Box I.21. For transit: only for the transit of animals or products through the European Union from one third country to another third country or from one part of a third country to another part of the same third country. State the name and ISO code of the third country of destination.
- Box I.22. For internal market: for all consignments destined to the European Union market.

Definitive import: this option must only be used for consignments intended to be placed under the customs procedure 'release for free circulation' in the European Union.

For certain animals (for example, registered equidae) only one of the following options must be selected:

Re-entry: this option must only be used for animals authorised for re-entry, such as registered horses for racing, competition and cultural events re-entering the European Union after their temporary export.

Temporary admission: this option must only be used for the entry of animals authorised for temporary entry into the European Union, such as registered horses for a period of less than 90 days.

- Box I.23. Total number of packages: the number of boxes, cages or stalls, in which the animals are being transported, the number of cryogenic containers for germinal products or the number of packages for products. In the case of bulk consignments, this box is optional.
- Box I.24. Quantity:

For animals: the total number of heads or straws expressed as units.

For germinal products: the total number of straws expressed as units.

For products and aquatic animals, except ornamental fish: the total gross and net weight in kilograms.

Total net weight: this is defined as the mass of the goods themselves without immediate containers or any packaging.

Total gross weight: overall weight in kilograms. This is defined as the aggregate mass of the products and of the immediate containers and all their packaging, but excluding transport containers and other transport equipment.

^{(&}lt;sup>5</sup>) Commission Implementing Regulation (EU) No 139/2013 of 7 January 2013 laying down animal health conditions for imports of certain birds into the Union and the quarantine conditions thereof (OJ L 47, 20.2.2013, p. 1).

Council Directive 2006/88/EC of 24 October 2006 on animal health requirements for aquaculture animals and products thereof, and on

the prevention and control of certain diseases in aquatic animal field requirements for aquaticative during and products directly and on the prevention and control of certain diseases in aquatic animals (OJ L 328, 24.11.2006, p. 14). Council Directive 2009/156/EC of 30 November 2009 on animal health conditions governing the movement and importation from third countries of equidae (OJ L 192, 23.7.2010, p. 1). Commission Regulation (EU) No 142/2011 of 25 February 2011 implementing Regulation (EC) No 1069/2009 of the European (7)

Parliament and of the Council laying down health rules as regards animal by-products and derived products not intended for human consumption and implementing Council Directive 97/78/EC as regards certain samples and items exempt from veterinary checks at the border under that Directive (OJ L 54, 26.2.2011, p. 1).

Box I.25. Description of goods: State the relevant Harmonised System code (HS code) and the title defined by the World Customs Organisation as referred to in Council Regulation (EEC) No 2658/87 (⁹). This customs description shall be supplemented, if necessary, by additional information required to classify the animals or the products in veterinary terms. In addition, state any specific requirements relating to the animals or to the nature/processing of the products as defined in the relevant European Union model health or veterinary certificate.

Zone: for animals or products affected by the setting up of approved zones or compartments in accordance with European Union legislation. The zones or production areas (for example, in the case of bivalve molluscs) must be indicated as published in the European Union lists of approved establishments.

For animals: the species, breed or category, identification method, identification number, age, sex, quantity or net weight, and test.

For germinal products: collection or production date, approval number of the centre or team, identification of the straw, and quantity. In addition, as regards donor animals, the species, breed or category, and identification.

For products: the species, types of products, type of treatment, approval number of establishments together with ISO country code (slaughter house, processing plant, cold store), number of packages, type of packaging, batch number, net weight, and final consumer (i.e. products are packed for final consumer).

Species: the scientific name or as defined in accordance with European Union legislation.

Type of packaging: identify the type of packaging according to the definition given in Recommendation No 21 (¹⁰) of UN/CEFACT (United Nations Centre for Trade Facilitation and Electronic Business).

Part II: Certification

This part must be completed by an official veterinarian or an official inspector.

Box II. Health information: please complete this part in accordance with the specific European Union health requirements relating to the animal species or to the nature of the products and as defined in the equivalence agreements with certain third countries or in other European Union legislation, such as that for certification.

Where there are no animal or public health attestations for the consignment, then the whole of this section shall be deleted or invalidated or not be present at all in accordance with the footnotes for Part II of the specific European Union health certificates.

- Box II.a. Certificate reference No: same reference code as in box I.2.
- Box II.b. IMSOC reference No: same reference code as in box I.2.a.
- Certifying officer: Official veterinarian or official inspector as defined by the relevant European Union legislation: the name in capital letters, qualification and title, where applicable, identification number and original stamp of the competent authority and date of signature.

^{(&}lt;sup>9</sup>) Council Regulation (EEC) No 2658/87 of 23 July 1987 on the tariff and statistical nomenclature and on the Common Customs Tariff (OJ L 256, 7.9.1987, p. 1).

⁽¹⁰⁾ Last version: Revision 9 Annexes V and VI as published on: http://www.unece.org/tradewelcome/un-centre-for-trade-facilitation-and-ebusiness-uncefact/outputs/cefactrecommendationsrec-index/list-of-trade-facilitation-recommendations-n-21-to-24.html

ANNEX III

MODEL OFFICIAL CERTIFICATES FOR THE ENTRY INTO THE UNION FOR PLACING ON THE MARKET OF ANIMALS AND GOODS INTENDED FOR HUMAN CONSUMPTION

PART I

CHAPTER A: MODEL OFFICIAL CERTIFICATE FOR THE ENTRY IN THE UNION FOR PLACING ON THE MARKET OF LIVE BIVALVE MOLLUSCS, ECHINODERMS, TUNICATES AND MARINE GASTROPODS

cou	OUNTRY							Official certificate to the EU			
	l.1.	Consignor/Exporter Name					1.2.	Certificate referen	ce No	I.2.a IMSOC referen	ce No
							I.3. Central Competent Authority				
L.		Address					I.4. Local Competent Authority				
nment		Tel. No									
nsig	I.5.	Consignee/Importer					I.6. Operator responsible for the consignment				
cor		Name					Name				
hed											
atcl		Address						Address			
disp	Postal code							Postal code			
of (Tel. No										
Part I: Details of dispatched consignment	1.7.	Country of origin	ISO	I.8. Region	of origin	Code	1.9.	Country of destination	ISO	I.10. Region of destination	Code
Part I:	l.11	Place of dispatch					l.12.	Place of destination	on		
		Name		Approval No	1			Name			
		Address						Address			
	I.13.	Place of loading					l.14.	Date and time of c	departure	e	
	115	Means of transport					116	Entry BCP			
	1.15.	means of transport					1.10.				
		Aeroplane	Ves	isel 🛛	Other		l.17.	Accompanying do	cuments	3	
		Road vehicle	Rai	lway 🛛							
								Туре			
		Identification:						No			
	119	Transport conditions									
	1.10.	Transport conditions									
		Ambient	Chi	lled 🛛	Frozen						
	1.19.	Container No/Seal No									

COU	NTRY				Official certificate to the EU
1.20.	Goods certified as				
	Human consumption				
I.21.				1.22.	
1.23.	Total number of packages	1.24. (Quantity		
		-	Fotal number	Total net weight (Kg)	Total gross weight (Kg)
1.25.	Description of goods				
	No Code and CN	l title			
	Species (Scientific name)		Nat	ture of commodity	Treatment type
			Cutting pl	ant/manufacturing plant	Cold store
F	inal consumer Number packag		Net weight	Batch No	Type of packaging

Part II: Certification

EN

Live bivalve molluscs, echinoderms, tunicates and marine gastropods II. Health information II.a. Certificate reference number II.b. II.1 (1) Public health attestation for live bivalve molluscs, echinoderms, tunicates and marine gastropods II. (1) Public health attestation for live bivalve molluscs, echinoderms, tunicates and marine gastropods II.1 (1) Public health attestation for live bivalve molluscs, echinoderms, tunicates and marine gastropods I. the undersigned, declare that I am aware of the relevant provisions of Regulation (EC) No 178/2002 of the European Parliament and of the Council of 28 January 2002 laying down the general principles and requirements of food law, establishing the European Food Safety Authority and laying down procedures in matters of food safety (OJ L 31, 1.2.2002, p. 1), Regulation (EC) No 852/2004 of the European Parliament and of the Council of 29 April 2004 on the

hygiene of foodstuffs (OJ L 139, 30.4.2004, p. 1) and Regulation (EC) No 853/2004 of the European Parliament and of the Council of 29 April 2004 laying down specific hygiene rules for food of animal origin (OJ L 139, 30.4.2004, p. 55)) and Regulation (EU) 2017/625 of the European Parliament and of the Council of 15 March 2017 on official controls and other official activities performed to ensure the application of food and feed law, rules on animal health and welfare, plant health and plant protection products, amending Regulations (EC) No 999/2001, (EC) No 396/2005, (EC) No 1069/2009, (EC) No 1107/2009, (EU) No 1151/2012, (EU) No 652/2014, (EU) 2016/429 and (EU) 2016/2031 of the European Parliament and of the Council, Council Regulations (EC) No 1/2005 and (EC) No 1099/2009 and Council Directives 98/58/EC, 1999/74/EC, 2007/43/EC, 2008/119/EC and 2008/120/EC and repealing Regulations (EC) No 854/2004 and (EC) No 882/2004 of the European Parliament and of the Council, Council Directives 89/608/EEC, 89/662/EEC, 90/425/EEC, 91/496/EEC, 96/23/EC, and 97/78/EC and Council Decision 92/438/EEC (Official Controls Regulation) (OJ L 95, 7.4.2017, p. 1) and certify that the (⁴) [live bivalve molluscs] (⁴) [live echinoderms] (⁴) [live tunicates] (⁴) [live marine gastropods] described above were produced in accordance with those requirements, in particular that they:

- come from (an) establishment(s) implementing a programme based on the HACCP principles in accordance with Article 5 of Regulation (EC) No 852/2004;
- have been harvested, where necessary relayed and transported in accordance with Section VII, Chapters I and II of Annex III to Regulation (EC) No 853/2004;
- were handled, where necessary purified, and packaged in compliance with Section VII, Chapters III and IV of Annex III to Regulation (EC) No 853/2004;
- satisfy the health standards laid down in Section VII, Chapter V of Annex III to Regulation (EC) No 853/2004 and the criteria laid down in Commission Regulation (EC) No 2073/2005 of 15 November 2005 on microbiological criteria for foodstuffs (OJ L 338, 22.12.2005, p. 1);
- have been packaged, stored and transported in compliance with Section VII, Chapters VI and VIII of Annex III to Regulation (EC) No 853/2004;
- have been marked and labelled in accordance with Section I of Annex II and Section VII, Chapter VII of Annex III to Regulation (EC) No 853/2004;
- in the case of *Pectinidae*, marine gastropods and *Holothuroidea* that are not filter feeders harvested outside classified production areas, comply with the specific requirements laid down in Section VII, Chapter IX of Annex III to Regulation (EC) No 853/2004;
- have satisfactorily undergone the official controls laid down in Articles 42 to 58 of Commission Implementing Regulation (EU) 2019/627 of 15 March 2019 laying down uniform practical arrangements for the performance of official controls on products of animal origin intended for human consumption in accordance with Regulation (EU) 2017/625 of the European Parliament and of the Council and amending Commission Regulation (EC) No 2074/2005 as regards official controls (OJ L 131, 17.5.2019, p. 51) and Article 7 of Commission Delegated Regulation (EU) 2019/624 of 8 February 2019 concerning specific rules for the performance of official controls on the production of meat and for production and relaying areas of live bivalve molluscs in accordance with Regulation (EU) 2017/625 of the European Parliament and of the Council (OJ L 131, 17.5.2019, p. 1); and
- fulfil the guarantees covering live animals and products thereof, if of aquaculture origin, provided by the residue plans submitted in accordance with Council Directive 96/23/EC of 29 April 1996 on measures to monitor certain substances and residues thereof in live animals and animal products and repealing Directives 85/358/EEC and 86/469/EEC and Decisions 89/187/EEC and 91/664/EEC (OJ L 125, 23.5.1996, p. 10), and in particular Article 29 thereof.
- II.2 (²) (⁴) Animal health attestation for live bivalve molluscs of aquaculture origin

II.2.1 (3) (4) [Requirements for species susceptible to Bonamia exitiosa, Perkinsus marinus and Mikrocytos mackini

I, the undersigned official inspector, hereby certify that the live bivalve molluscs referred to in Part I of this certificate:

Live bivalve molluscs, echinoderms, tunicates and marine

COUN	ITRY	Live bivalve molluscs, ec	gastropod			
II.	Health information	II.a. Certificate reference number	II.b.			
	marinus] (⁴) [<i>Mikrocytos mackini</i>] in accordance animal health requirements for aquaculture an	or compartment declared free from (⁴) [<i>Bonamia exitiosa</i>] (⁴) [<i>Perkinsus</i> ince with Chapter VII of Council Directive 2006/88/EC of 24 October 2006 on animals and products thereof, and on the prevention and control of certain 11.2006, p. 14) or the relevant OIE Standard by the competent authority of				
	 where the relevant diseases are notifiab relevant disease must be immediately inv 		orts of suspicion of infection of the			
	— all introduction of species susceptible to t	he relevant diseases come from an are	ea declared free of the disease.]			
II.2.2	(³) (⁴) [Requirements for species susceptible State, zone or compartment declared diseas relevant disease					
	I, the undersigned official inspector, hereby cer	tify that the live bivalve molluscs referr	red to above:			
(⁶) originate from a country/territory, zone or compartment declared free from (⁴) [<i>Marteilia refringens</i>] (⁴) [<i>Bo ostreae</i>] in accordance with Chapter VII of Directive 2006/88/EC or the relevant OIE Standard by the com authority of my country,						
 where the relevant diseases are notifiable to the competent authority and reports of suspicion of infection of the relevant disease must be immediately investigated by the official services, and 						
	(ii) all introduction of species susceptible to the relevant diseases come from an area declared free of the disease.]					
1.2.3	Transport and labelling requirements					
	I, the undersigned official inspector, hereby cer	tify that:				
11.2.3	 .1 the live bivalve molluscs referred to above are their health status, 	placed under conditions, including w	ith a water quality, that do not alte			
II.2.3	.2 the transport container or well boat prior to load	ling is clean and disinfected or previou	isly unused; and			
II.2.3	.3 the consignment is identified by a legible label the ship's manifest, with the relevant informa following statement:					
	'Live bivalve molluscs intended for human cons	sumption in the Union'.				
certi	s notes in Annex II of Commission Implementing ficates for certain animals and goods and am 2016/759 as regards these model certificates ((ending Regulation (EC) No 2074/20	-			
Part	I:					
_	Box reference I.8: Region of origin: indicate the pr	oduction area.				
Part	11:					
(1)	Part II.1 <u>does not</u> apply to countries with spe Agreements or other Union legislation.	cial public health certification requir	ements laid down in Equivalenc			
(²)	Part II.2 does not apply to:					
	(a) non-viable molluscs, which means molluscs from which they were obtained,	no longer able to survive as living ani	mals if returned to the environmer			

(b) live bivalve molluscs placed on the market for human consumption without further processing, provided that they are packed in retail-sale packages which comply with the provisions for such packages in Regulation (EC) No 853/2004,

Live bivalve molluscs, echinoderms, tunicates and marine

cou	NTRY	Live bivalve molluscs, ec	hinoderms, tunicates and marine gastropods				
II.	Health information	II.a. Certificate reference number	II.b.				
	(c) live bivalve molluscs destined for processing 2006/88/EC, or for dispatch centres, purifica treatment system inactivating the pathogens reducing the risk of transmitting diseases to the	tion centres or similar businesses w in question, or where the effluent is	hich are equipped with an effluent subject to other types of treatment				
	(d) live bivalve molluscs which are intended for further processing before human consumption without temporary storage at the place of processing and packed an labelled for that purpose in accordance with Regulation (EC) No 853/2004.						
(3)	Part II.2.1 and II.2.2 <u>only</u> apply to species susceptible to one or more of the diseases referred to in the title. Susceptible species are listed in Annex IV to Directive 2006/88/EC.						
(4)	Keep as appropriate.						
(5)	For consignments of species susceptible to <i>Bonamia exitiosa, Perkinsus marinus</i> and <i>Mikrocytos mackini</i> this statement must be kept for the consignment to be authorised into any part of the Union.						
(6)	To be authorised into a Member State, zone or compartment (boxes I.9 and I.10 of Part I of the certificate) declared free from <i>Marteilia refringens</i> or <i>Bonamia ostreae</i> or with a surveillance or eradication programme established in accordance with Article 44(1) or (2) of Directive 2006/88/EC, one of these statements must be kept if the consignment contain species susceptible to the disease(s) for which disease freedom or programme(s) apply(ies). Data on the disease status of each farms and mollusc farming areas in the Union are accessible at http://ec.europa.eu/food/animal/liveanimals/aquaculture/index_en.htm.						
_	The colour of the stamp and signature must be diff	erent to that of the other particulars in	the certificate.				
Offi	cial inspector						
	Name (in capital letters):	Qualification and title:					
	Date:	Signature:					
	Stamp:						

CHAPTER B: ADDITIONAL MODEL OFFICIAL CERTIFICATION FOR PROCESSED BIVALVE MOLLUSCS BELONGING TO THE SPECIES ACANTHOCARDIA TUBERCULATUM

The official inspector hereby certifies that the processed bivalve molluscs of the species *Acanthocardia tuberculatum*, certified in the health certificate reference No:

- were harvested in production areas clearly identified, monitored and authorised by the competent authority in accordance with Article 12 of Commission Delegated Regulation (EU) 2019/625 of 4 March 2019 supplementing Regulation (EU) 2017/625 of the European Parliament and of the Council with regard to requirements for the entry into the Union of consignments of certain animals and goods intended for human consumption (OJ L 131, 17.5.2019, p. 18) and where the paralytic shellfish poisoning (PSP) level in the edible parts of these molluscs is lower than 300 µg for 100g;
- 2. were transported in containers or vehicles sealed by the competent authority, directly to the establishment:

(name and official approval number of the establishment, authorised specially by the

(name and official approval number of the establishment, authorised specially by the competent authority to carry out their treatment);

- 3. were accompanied while being transported to this establishment by a document issued by the competent authority which authorises the transport, attesting to the nature and quantity of the product, area of origin and establishment of destination;
- 4. were subjected to the heat treatment outlined in the Annex to Commission Decision 96/77/EC of 18 January 1996 establishing the conditions for the harvesting and processing of certain bivalve molluscs coming from areas where the paralytic shellfish poison level exceeds the limits laid down by Council Directive 91/495/EEC (OJ L 15, 20.1.1996, p. 46); and
- 5. do not contain a PSP level detectable by the bioassay method, as demonstrated by the attached analytical report(s) of the test carried out on each lot included in the consignment covered by this certification.

The official inspector hereby certifies that the competent authority has verified that the 'own health' checks carried out in the establishment referred to in point 2 are specifically applied to the heat treatment referred to in point 4.

The undersigned official inspector hereby declares that he/she is aware of the provisions of Decision 96/77/EC and that the attached analytical report(s) correspond(s) to the test carried out on the products after processing.

Official inspector						
Name (in capitals):	Qualification and title:					
Date:	Signature:					
Stamp:						

PART II CHAPTER A: MODEL OFFICIAL CERTIFICATE FOR THE ENTRY IN THE UNION FOR PLACING ON THE MARKET OF FISHERY PRODUCTS

cou	COUNTRY						Official certificate to the EU				
	l.1.	Consignor/Exporter						I.2.	Certificate referen	ice No	I.2.a IMSOC reference No
		Name						I.3.	I.3. Central Competent Authority		
		Address						I.4.	4. Local Competent Authority		
nent		Tel. No									
signr	1.5.	Consignee/Importer						I.6.	Operator respons	ible for th	ne consignment
d con		Name							Name		
spatche		Address							Address		
of dis		Postal code						Postal code			
ails c		Tel. No									
Part I: Details of dispatched consignment	1.7.	Country of origin	ISO	I.8. R	egion	of origin	Code	I.9. Country of ISO destination		ISO	I.10.
Ра	l.11	Place of dispatch						l.12.	Place of destination	on	
		Name		Approv	al No				Name		
		Address							Address		
	I.13.	Place of loading						l.14.	Date and time of o	departure)
	l.15.	Means of transport						I.16.	Entry BCP		
		Aeroplane	Ves	sel		Other		l.17.	Accompanying do	ocuments	
		Road vehicle	Rai	lway	way 🛛				Туре		
		Identification:						No			
	I.18. Transport conditions										
		Ambient 🛛	Chi	lled		Frozen					
	l.19.	Container No/Seal No	I								

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COUNTRY			Official certificate to the EU	
I.20. Goods certified as Canning industry □				
Human consumption				
l.21.		1.22.		
	1			
I.23. Total number of packages	I.24. Quantity			
	Total number	Total net weight (Kg)	Total gross weight (Kg)	
I.25. Description of goods		·		
No Code and CN	title			
Species (Scientific name)	Nat	ture of commodity	Treatment type	
	Vessel	l/manufacturing plant	Cold store	
Final consumer Number package		Batch No	Type of packaging	

	COUNT	ſRY	Fishery products						
	11.	Health information	II.a. Certificate reference number	II.b.					
	II.1.	(¹) Public health attestation							
Part II: Certification		I, the undersigned, declare that I am aware of the Parliament and of the Council of 28 January 2 establishing the European Food Safety Autho 1.2.2002, p. 1), Regulation (EC) No 852/2004 of hygiene of foodstuffs (OJ L 139, 30.4.2004, p. 1 the Council of 29 April 2004 laying down specir and Regulation (EU) 2017/625 of the European other official activities performed to ensure the a health and plant protection products, amending (EC) No 1107/2009, (EU) No 1151/2012, (EU Parliament and of the Council, Council Regula 98/58/EC, 1999/74/EC, 2007/43/EC, 2008/119/E (EC) No 882/2004 of the European Parliame 90/425/EEC, 91/496/EEC, 96/23/EC, 96/93/EC Regulation) (OJ L 95, 7.4.2017, p. 1) and certify with those requirements, in particular that they:	bles and requirements of food law, matters of food safety (OJ L 31, ne Council of 29 April 2004 on the of the European Parliament and of origin (OJ L 139, 30.4.2004, p. 55)) March 2017 on official controls and on animal health and welfare, plant No 396/2005, (EC) No 1069/2009, (EU) 2016/2031 of the European 1099/2009 and Council Directives Regulations (EC) No 854/2004 and ectives 89/608/EEC, 89/662/EEC, ion 92/438/EEC (Official Controls						
		 come from (an) establishment(s) implement points (HACCP) principles in accordance w 							
		 have been caught and handled on board frozen and thawed hygienically in complia Annex III to Regulation (EC) No 853/2004; 							
	-	 satisfy the health standards laid down in S the criteria laid down in Commission Reg criteria for foodstuffs (OJ L 338, 22.12.2005) 	gulation (EC) No 2073/2005 of 15 N						
		 have been packaged, stored and transport Regulation (EC) No 853/2004; 	ted in compliance with Section VIII,	Chapters VI to VIII of Annex III to					
		 have been marked in accordance with Sec 	tion I of Annex II to Regulation (EC) N	lo 853/2004;					
		 fulfil the guarantees covering live animals plans submitted in accordance with Coun- substances and residues thereof in live a 86/469/EEC and Decisions 89/187/EEC an thereof; and 	cil Directive 96/23/EC of 29 April 19 nimals and animal products and rep	96 on measures to monitor certain bealing Directives 85/358/EEC and					
		 have satisfactorily undergone the official Regulation (EU) 2019/627 of 15 March 20 official controls on products of animal origi 2017/625 of the European Parliament No 2074/2005 as regards official controls (019 laying down uniform practical arr in intended for human consumption in and of the Council and amendii	angements for the performance of n accordance with Regulation (EU)					
	II.2	$(^2)$ $(^4)$ Animal health attestation for fish and cr	ustaceans of aquaculture origin						
	II.2.1	(³) (⁴) [Requirements for species susceptible yellowhead disease	to epizootic haematopoietic necr	osis (EHN), taura syndrome and					
		I, the undersigned official inspector, hereby certition this certificate:	fy that the aquaculture animals or pro	ducts thereof referred to in Part I of					
		(⁵) originate from a country/territory, zone or [yellowhead disease] in accordance with Chapi health requirements for aquaculture animals and in aquatic animals (OJ L 328, 24.11.2006, p. 14)	er VII of Council Directive 2006/88/ I products thereof, and on the preven	EC of 24 October 2006 on animal tion and control of certain diseases					
		(i) where the relevant diseases are notifiable relevant disease must be immediately inve		orts of suspicion of infection of the					
		(ii) all introduction of species susceptible to the	e relevant diseases come from an are	ea declared free of the disease, and					

COUNT			Fishery products					
II.	Health information	II.a. Certificate reference number	II.b.					
	(iii) species susceptible to the relevant of	diseases are not vaccinated against the rele	vant diseases.]					
II.2.2	necrosis (IHN), infectious salmon ana	ptible to viral haemorrhagic septicaemia emia (ISA), koi herpes virus (KHV) and v nt declared disease free or subject to	white spot disease intended for a					
	I, the undersigned official inspector, here this certificate:	by certify that the aquaculture animals or pro	oducts thereof referred to in Part I c					
	(⁶) originate from a country/territory, zone or compartment declared free from (⁴) [VHS] (⁴) [IHN] (⁴) [ISA] (⁴) [KHV] (⁴) [White spot disease] in accordance with Chapter VII of Directive 2006/88/EC or the relevant OIE Standard by the competent authority of my country,							
		otifiable to the competent authority and rep ely investigated by the competent authority,	orts of suspicion of infection of the					
	(ii) all introduction of species suscepti and	ble to the relevant diseases come from an	area declared free of the disease					
	(iii) species susceptible to the relevant	diseases are not vaccinated against the rele	vant diseases.]					
II.2.3	Transport and labelling requirements							
	l, the undersigned official inspector, here	by certify that:						
II.2.3.	1 the aquaculture animals referred to abor health status;	ve are placed under conditions in which th	e water quality does not alter thei					
II.2.3.	3.2. prior to loading the transport container or well boat is clean and disinfected or previously unused; and							
II.2.3.	2.3.3. the consignment is identified by a legible label on the exterior of the container, or when transported by well boat, in the ship's manifest, with the relevant information referred to in boxes I.7 to I.11 of Part I of this certificate, and the following statement:							
	'(⁴) [Fish] (⁴) [Crustaceans] intended fo	r human consumption in the Union'.						
Notes								
certif		enting Regulation (EU) 2019/628 of 8 Apr Id amending Regulation (EC) No 2074/20 ates (OJ L 131, 17.5.2019, p. 101)						
Part I	:							
— E	Box reference I.8: Region of origin: For froze	en or processed bivalve molluscs, indicate th	ne production area.					
		for whole fish initially frozen in brine at -9° nce with the requirements of Section VIII, (consumption' for the other cases.						
	Box reference I.25: Insert the appropriate H 0304, 0305, 0306, 0307, 0308, 0511, 1504,	Harmonised System (HS) code(s) using he 1516, 1518, 1603, 1604, 1605 or 2106.	adings such as: 0301, 0302, 0303					
— I	Box reference I.25: Nature of com	modity: specify whether aquaculture or wild	origin.					
	Treatment typ	e: specify whether live, chilled, frozen or pro	cessed.					
	Manufacturing	<i>plant</i> : includes factory vessel, freezer vesse	el, reefer vessels, cold					
store	and processing plant.							
Part I	:							
	Part II.1 of this certificate <u>does not</u> apply equivalence agreements or other EU legisla	to countries with special public health cert tion	ification requirements laid down in					

staceans that cannot survive as living animals prated before dispatch, pereof, which are placed on the market for l										
rated before dispatch, hereof, which are placed on the market for l										
ereof, which are placed on the market for										
	(c) aquaculture animals and products thereof, which are placed on the market for human consumption without further processing, provided that they are packed in retail-sale packages which comply with the provisions for such packages in Regulation (EC) No 853/2004,									
(d) crustaceans destined for processing establishments authorised in accordance with Article 4(2) of Directive 2006/88/EC, or for dispatch centres, purification centres or similar businesses which are equipped with an effluent treatment system that inactivates the pathogens in question, or where the effluent undergoes other types of treatment reducing the risk of transmitting diseases to the natural waters to an acceptable level, and										
For consignments of species susceptible to EHN, taura syndrome and/or yellowhead disease this statement must be kept for the consignment to be authorised into any part of the EU.										
In order to be authorised into a Member State, zone or compartment (boxes I.9 and I.10 of Part I of the certificate) declared free from VHS, IHN, ISA, KHV or white spot disease or with a surveillance or eradication programme drawn up in accordance with Article 44(1) or (2) of Directive 2006/88/EC, one of these statements must be kept if the consignment contain species susceptible to the disease(s) for which disease freedom or programme(s) apply(ies). Data on the disease status of each farm and mollusc farming area in the Union are accessible at http://ec.europa.eu/food/animal/liveanimals/aquaculture/index_en.htm.										
The colour of the stamp and signature must be different to that of the other particulars in the certificate.										
Official inspector										
Name (in capital letters): Qualification and title:										
Date: Signature:										
Stamp:										
	ng establishments authorised in accordar , purification centres or similar businesses w pathogens in question, or where the effluent ises to the natural waters to an acceptable lew further processing before human consumption labelled for that purpose in accordance with R <u>aby</u> apply to species susceptible to one or mo- pole species are listed in Annex IV to Directive is to EHN, taura syndrome and/or yellowhead of any part of the EU. State, zone or compartment (boxes I.9 and I.10 e spot disease or with a surveillance or en- of Directive 2006/88/EC, one of these to the disease(s) for which disease free farm and mollusc farming area in s/aquaculture/index_en.htm. st be different to that of the other particulars in Qualification and title:									

CHAPTER B: MODEL OF OFFICIAL CERTIFICATE FOR FISHERY PRODUCTS CAUGHT BY VESSELS FLYING THE FLAG OF A MEMBER STATE AND TRANSFERRED IN THIRD COUNTRIES WITH OR WITHOUT STORAGE

cour	OUNTRY						Official certificate to the EU					
	I.1. Consignor/Exporter						I.2.	Certificate referen	ice No	I.2.a IMSOC reference No		
	Name I.							1.3.	I.3. Central Competent Authority			
	Address								I.4. Local Competent Authority			
nent		Tel. No										
signn	I.5. Consignee/Importer							I.6. Operator responsible for the consignment				
d con		Name							Name			
Part I: Details of dispatched consignment		Address Postal code Tel. No							Address Postal code			
ırt I: Detai	1.7.	Country of origin	ISC	D I.8.	Region	of origin	jin Code	1.9.	Country of destination	ISO	I.10.	
Ра	l.11	Place of dispatch				I.12.	I.12. Place of destination	on				
		Name	Арр	Approval No			Name					
		Address						Address				
	I.13.	Place of loading						I.14. Date and time of departure				
	l.15.	Means of transport	:					I.16.	I.16. Entry BCP			
		Aeroplane ['essel	ssel 🔲 Other 🗖			I.17.	Accompanying do	ocuments	ents	
		Road vehicle		ailway					Туре			
		Identification:							No			
	l.18.	18. Transport conditions						-				
		Ambient		chilled		Frozen						
	l.19.	Container No/Seal	No			1						

17.5.2019

COUNTRY			Official certificate to the EU					
I.20. Goods certified as Canning industry								
Human consumption								
I.21.		1.22.						
I.23. Total number of packages	I.24. Quantity Total number	Total net weight (Kg)	Total gross weight (Kg)					
I.25. Description of goods								
No Code and CN	title							
Species (Scientific name)	Nat	ture of commodity	Treatment type					
Zone Vesse		//manufacturing plant	Cold store					
Final consumer Number of package		Batch No	Type of packaging					

Part II: Certification

cou	INTRY		Fishery prod	ucts transferred in third countries							
11.	Hea	alth information	II.a. Certificate reference number	II.b.							
11.1.	Put	Public health attestation									
	I, the undersigned official inspector, declare that I am aware of the relevant provisions of Regulation (EC) No 178/2002 of the European Parliament and of the Council of 28 January 2002 laying down the general principles and requirements of food law, establishing the European Food Safety Authority and laying down procedures in matters of food safety (OJ L 31, 1.2.2002, p. 1), Regulation (EC) No 852/2004 of the European Parliament and of the Council of 29 April 2004 on the hygiene of foodstuffs (OJ L 139, 30.4.2004, p. 1) and Regulation (EC) No 853/2004 of the European Parliament and of the Council of 29 April 2004 laying down specific hygiene rules for food of animal origin (OJ L 139, 30.4.2004, p. 55)) and Regulation (EU) 2017/625 of the European Parliament and of the Council of 15 March 2017 on official controls and other official activities performed to ensure the application of food and feed law, rules on animal health and welfare, plant health and plant protection products, amending Regulations (EC) No 1069/2009, (EC) No 1107/2009, (EU) No 1151/2012, (EU) No 652/2014, (EU) 2016/429 and (EU) 2016/2031 of the European Parliament and of the Council Directives 98/58/EC, 1999/74/EC, 2007/43/EC, 2008/119/EC and 2008/120/EC and repealing Regulations (EC) No 854/2004 and (EC) No 882/2004 of the European Parliament and of the Council Directives 89/608/EEC, 89/662/EEC, 90/425/EEC, 91/496/EEC, 96/23/EC, 96/93/EC and 97/78/EC and Council Decision 92/438/EEC (Official Controls Regulation) (OJ L 95, 7.4.2017, p. 1) and certify that the fishery products described above:										
	_	have been landed and unloaded hygienica (indicate approval/registration number(s) requirements laid down in Chapter II of Se	and name of the flag Member State	(s)) in compliance with the relevant							
	_	if applicable, have been stored in approve compliance with the relevant requireme No 853/2004;									
-	 if applicable, have been loaded hygienically on the approved vessel(s)										
	 if applicable, have been loaded in a container										
	_	are accompanied by the print out(s) (**) of	the fishing logbook(s) or relevant par	ts thereof. (**)							
	(**) Electronic format is also accepted.										
Not	es										
cert	See notes in Annex II of Commission Implementing Regulation (EU) 2019/628 of 8 April 2019 concerning model official certificates for certain animals and goods and amending Regulation (EC) No 2074/2005 and Implementing Regulation (EU) 2016/759 as regards these model certificates (OJ L 131, 17.5.2019, p. 101)										
Part	t I:										
_	 Box reference I.11: Place of dispatch: State the name, address and approval number of the cold store in the third country of dispatch or, if the product was not in cold storage, state the name and approval or registration number of the Member State flagged vessel of origin. 										
_	Box reference I.15: State the means of transport leaving the third country of dispatch. In the case of freezer/reefer vessels, state the name of the vessel, approval number and flag State; in the case of a fishing vessel state the registration number and flag State. If the means of transport are containers, trucks or aeroplanes the same indications provided for in the fourth indent of Part II.1 must be stated.										
-	18 °C	eference I.20:Tick 'Canning industry' for who and intended for canning in accordance wi ation (EC) No 853/2004. Tick 'Human consu	th the requirements of Section VIII, (

COU	NTRY	Fishery products transferred in third countries									
II.	Health information	II.a.	Certificate reference number	II.b.							
_	 Box reference I.25: Insert the appropriate Harmonised System (HS) code(s) using headings such as: 0301, 0302, 030 0304, 0305, 0306, 0307, 0308, 0511, 1504, 1516, 1518, 1603, 1604, 1605 or 2106. 										
—	 Box reference I.25: Treatment type: specify whether chilled, frozen or processed. 										
	(*) includes fishing vessel, factory vessel, freezer a	and re	eefer vessel as applicable.								
Offic	cial inspector										
	Name (in capital letters):		Qualification and title:								
	Date:		Signature:								
	Stamp:										

CHAPTER C: MODEL OF OFFICIAL CERTIFICATE TO BE SIGNED BY THE CAPTAIN ACCOMPANYING FROZEN FISHERY PRODUCTS WHEN ENTERING THE UNION FOR PLACING ON THE MARKET DIRECTLY FROM A FREEZER, REEFER OR FACTORY VESSEL

cou	COUNTRY								Official certificate to the EU
	l.1.	I.1. Consignor/Exporter					Certificate referen	nce No	I.2.a IMSOC reference No
		Name				1.3.			
		Address				1.4.			
nent		Tel. No							
Isignr	1.5.	Consignee/Importer				I.6.	Operator respons	ible for t	ne consignment
d con		Name				Name			
Part I: Details of dispatched consignment		Address Postal code Tel. No		Address Postal code					
art I: Det	1.7.	Country of origin	ISO	I.8. Region of origin	Code	1.9.	Country of destination	ISO	I.10.
Å	l.11	Place of dispatch				I.12.	Place of destination	on	
		Name		Approval No			Name		
		Address					Address		
	I.13.					I.14.	Date and time of o	departur	e
	I.15.					I.16.	Entry BCP		
						l.17.	Accompanying do	ocuments	3
							Type No		
	l.18.								
	I.19.								

17.5.2019

COUNTRY				Official certificate to the EU
I.20. Goods certified as Canning industry □				
Human consumption				
l.21.			1.22.	
I.23. Total number of packages		Quantity Total number	Total net weight (Kg)	Total gross weight (Kg)
I.25. Description of goods	1		·	
No Code and CN	title			
Species (Scientific name)				
Final consumer Number package		Net weight	Batch No	Type of packaging

	COUNT	RY					Fishery products
	l.(bis)	Oth	er information				
	Fishing	area(s):				
	IMO/Lic	oyd's r	number (if issued) or call sign of the ve	essel			
	Fishing	perio	d: Start d	ate: .	// Sto	p da	ate://
	11.	Healt	h attestation		II.a. Certificate reference number	•	ll.b.
	II.1	Publi	ic health attestation				
		l, unc	lersigned, declare that:				
Part II: Certification			I am aware of the relevant provision Council of 28 January 2002 laying of European Food Safety Authority and Regulation (EC) No 852/2004 of the foodstuffs (OJ L 139, 30.4.2004, p. 1 Council of 29 April 2004 laying down and Regulation (EU) 2017/625 of the and other official activities performed welfare, plant health and plant protect (EC) No 1069/2009, (EC) No 1107 2016/2031 of the European Parliar No 1099/2009 and Council Directive repealing Regulations (EC) No 854/2 Council Directives 89/608/EEC, 89/6 Council Decision 92/438/EEC (Offici products described above were pro appears on the list of vessels from whether the second	dowr I layii Euro I) an Euro Euro d to e ction 7/200 ment S 98 2004 662/E al Co duce hich i	the general principles and requir ng down procedures in matters of opean Parliament and of the Counci d Regulation (EC) No 853/2004 of cific hygiene rules for food of anima pean Parliament and of the Counci- ensure the application of food and products, amending Regulations (E 9, (EU) No 1151/2012, (EU) No and of the Council, Council Reg /58/EC, 1999/74/EC, 2007/43/EC, and (EC) No 882/2004 of the Euro EC, 90/425/EEC, 91/496/EEC, 96 ontrols Regulation) (OJ L 95, 7.4.2 d in accordance with those requir mports to the Union are permitted (I	eme food cil of al or il of feec EC) 6522 gulat 200 017, eme bein	ents of food law, establishing the I safety (OJ L 31, 1.2.2002, p. 1), f 29 April 2004 on the hygiene of European Parliament and of the igin (OJ L 139, 30.4.2004, p. 55)) 15 March 2017 on official controls d law, rules on animal health and No 999/2001, (EC) No 396/2005, /2014, (EU) 2016/429 and (EU) tions (EC) No 1/2005 and (EC) 08/119/EC and 2008/120/EC and an Parliament and of the Council, EC, 96/93/EC and 97/78/EC and , p. 1) and certify that the fishery ents, in particular that the vessel g 'EU-listed');
			the vessel has a programme based of hazards in accordance with Article 5 of			ol po	ints (HACCP) principles to control
	-	_	the fishery products have been cau, prepared, processed, frozen and Section VIII, Chapters I to IV of Anr danger to public health have been human consumption;	thaw nex I	ed hygienically in compliance w II to Regulation (EC) No 853/2004	∕ith 4.∨is	the requirements laid down in scera and parts that may pose a
		_	the fishery products satisfy the healt (EC) No 853/2004 and, where approp 15 November 2005 on microbiologica	priate	, the criteria laid down in Commiss	sion	Regulation (EC) No 2073/2005 of
			the fishery products have been pact to VIII of Annex III to Regulation (EC)			liano	ce with Section VIII, Chapters VI
		_	the fishery products have been marke	ed in	accordance with Section I of Annex	c II to	o Regulation (EC) No 853/2004;
		_	the fishery products fulfil the guara provided by the residue plans subr measures to monitor certain substan Directives 85/358/EEC and 86/469/E and in particular Article 29 thereof; an	nitteo ices EC a	I in accordance with Council Dire and residues thereof in live animal	ectivo Is ar	e 96/23/EC of 29 April 1996 on nd animal products and repealing
			frozen fishery products have been ke whole fish initially frozen in brine inter of not more than – 9 °C.				

coul	NTRY		Fishery products				
II.	Health attestation	II.a. Certificate reference number	II.b.				
cert	es notes in Annex II of Commission Implementing ificates for certain animals and goods and amo 2016/759 as regards these model certificates (0	ending Regulation (EC) No 2074/20					
Part	l:						
-	Box reference I.2: A unique document number acc	cording to your own classification.					
_	Box reference I.5: The name and address (street, town and post code) of the physical or legal person to whom the consignment is imported directly to in the Member State of destination.						
—	Box reference I.7: The country whose flag is being flown by the vessel issuing this document.						
_	Box reference I.11: The name of the vessel and Delegated Regulation (EU) 2019/625 of 4 Ma Parliament and of the Council with regard to requ and goods intended for human consumption (Ov imported.	rch 2019 supplementing Regulation irements for the entry into the Union	(EU) 2017/625 of the European of consignments of certain animals				
	Box reference I.20: Tick 'Canning industry' for who 18 °C and intended for canning in accordance w Regulation (EC) No 853/2004. Tick 'Human consu	ith the requirements of Section VIII, (
_	Box reference I.25: Insert the appropriate Harmo 0304, 0305, 0306, 0307, 0308, 0511, 1504, 1516,		adings such as: 0301, 0302, 0303,				
_	Box reference I.25: Treatment type: specify wheth	er chilled, frozen or processed.					
	(*) includes fishing vessel, factory vessel, freezer a	and reefer vessel as applicable.					
Cap	tain of the vessel						
	Name (in capital letters):						
	Date:	Signa	ature:				
	Stamp:						

PART III

MODEL OFFICIAL CERTIFICATE FOR THE ENTRY INTO THE UNION FOR PLACING ON THE MARKET OF CHILLED, FROZEN OR PREPARED FROGS' LEGS INTENDED FOR HUMAN CONSUMPTION

cou	OUNTRY							Official certificate to the EU			
	l.1.	Consignor/Exporter						1.2.	Certificate referen	ce No	I.2.a IMSOC reference No
		Name						I.3. Central Competent Authority			
		Address							.4. Local Competent Authority		
nent		Tel. No									
signr	I.5.	Consignee/Importer						I.6.	Operator responsi	ible for th	ne consignment
d con		Name							Name		
Part I: Details of dispatched consignment		Address Postal code							Address Postal code		
ils of		Tel. No									
urt I: Deta	1.7.	Country of origin	ISO	1.8.				1.9.	Country of destination	ISO	I.10.
Å	l.11	Place of dispatch					I.12.	Place of destination	on		
		Name		Approval	No				Name		
		Address							Address		
	l.13.	Place of loading						I.14. Date and time of departure)
	l.15.	Means of transport						I.16.	Entry BCP		
		Aeroplane	Ves	ssel 🛛		Other		l.17.	Accompanying do	cuments	
		Road vehicle	Rai	lway 🛛					Туре		
		Identification:							No		
	l.18.	Transport conditions									
		Ambient 🛛	Chi	lled 🗆		Frozen					
	l.19.	Container No/Seal No				·		<u> </u>			

COUNTRY			Official certificate to the EU
I.20. Goods certified as			
Human consumption			
I.21.		1.22.	
I.23. Total number of packages	I.24. Quantity		
	Total number	Total net weight (Kg)	Total gross weight (Kg)
I.25. Description of goods			
No Code and CN	title		
Species (Scientific name)	Ма	anufacturing plant	Treatment type Cold store
Final consumer Number package		Batch No	Type of packaging

	COUN	TRY	Chilled, frozen or prepared frogs' legs	Model FRG intended for human consumption
	П.	Health information	II.a. Certificate reference No	II.b.
	II.1 .	Public health attestation		
Part II: Certification			ary 2002 laying down the general princ uthority and laying down procedures 004 of the European Parliament and of p. 1) and Regulation (EC) No 853/200 ecific hygiene rules for food of animal or Parliament and of the Council of 15 Mar application of food and feed law, rules ding Regulations (EC) No 999/2001, (EC (EU) No 652/2014, (EU) 2016/429 ar gulations (EC) No 1/2005 and (EC) N 19/EC and 2008/120/EC and repealing ament and of the Council, Council E 3/EC and 97/78/EC and Council Dec	siples and requirements of food law, in matters of food safety (OJ L 31, the Council of 29 April 2004 on the 4 of the European Parliament and of igin (OJ L 139, 30.4.2004, p. 55) and ch 2017 on official controls and other on animal health and welfare, plant C) No 396/2005, (EC) No 1069/2009, id (EU) 2016/2031 of the European o 1099/2009 and Council Directives (Regulations (EC) No 854/2004 and Directives 89/608/EEC, 89/662/EEC, ision 92/438/EEC (Official Controls hese requirements, in particular that hazard analysis and critical control
		 have been handled and, where approving with the requirements of Annex II to R 	priate, prepared, packaged and stored egulation (EC) No 852/2004; and	in a hygienic manner in accordance
			ed, prepared and, where appropriate, ch cordance with the requirements of Secti	
	certifi	s notes in Annex II of Commission Implemen icates for certain animals and goods and 2016/759 as regards these model certificate	amending Regulation (EC) No 2074/2	
	Part I	:		
	— Е	Box reference I.25: Insert the appropriate CN o	code(s) such as: 0208 90 70, 0210 99 39) or 1602 90 99.
	— Е	Box reference I.25: <i>Treatment type</i> : fresh, trea	ted.	
	Part II	l:		
	ר —	The colour of the stamp and signature must be	different from that of the other particula	rs in the certificate.
	Officia	al inspector		
	1	Name (in capital letters):	Q	ualification and title:
	c	Date:	S	ignature:
	5	Stamp:		

PART IV

MODEL OFFICIAL CERTIFICATE FOR THE ENTRY INTO THE UNION FOR PLACING ON THE MARKET OF CHILLED, FROZEN, SHELLED, COOKED, PREPARED OR PRESERVED SNAILS INTENDED FOR HUMAN CONSUMPTION

cou	COUNTRY						Official certificate to the EU				
	l.1.	Consignor/Exporter						1.2.	Certificate referen	ce No	I.2.a IMSOC reference No
		Name						1.3.	Central Competer	nt Author	ity
		Address						1.4.	Local Competent	Authority	1
nent		Tel. No									
Isignr	1.5.	Consignee/Importer						I.6.	Operator responsi	ible for th	ne consignment
ed co		Name							Name		
spatch		Address							Address		
of dis		Postal code							Postal code		
tails		Tel. No		1							
Part I: Details of dispatched consignment	1.7.	Country of origin	ISO	1.8.				1.9.	Country of destination	ISO	I.10.
à	I.11	Place of dispatch						I.12.	Place of destination	on	
		Name		Approval	No				Name		
		Address							Address		
	l.13.	Place of loading					I.14. Date and time of departure				
	l.15.	Means of transport						l.16.	Entry BCP		
		Aeroplane	Ves	ssel 🗆]	Other		I.17.	Accompanying do	cuments	5
		Road vehicle	Rai	lway 🗖]				Туре		
		Identification:							No		
	l.18.	Transport conditions									
		Ambient 🛛	Chi	lled D]	Frozen					
	I.19.	Container No/Seal No)			1		I			

COUNTR	Y				Official certificate to the EU
I.20. God	ods certified as				
Hur	man consumption				
l.21.				1.22.	
I.23. Tota	al number of packages		Quantity		
		T	otal number	Total net weight (Kg)	Total gross weight (Kg)
1.25. Des	scription of goods				
No	Code and CN	title			
	Species (Scientific name)		Ма	nufacturing plant	Treatment type Cold store
Final	consumer Number package		Net weight	Batch No	Type of packaging

	COUN	NTRY	Ch	illed, frozen, shelled, cooked, prep	Model SNS bared or preserved snails intended for human consumption
	П.	Неа	Ith information	II.a. Certificate reference No	II.b.
	II.1.	Pub	lic health attestation		VC.
Part II: Certification		Parl esta 1.2.: hygi the Reg offic heal (EC Parl 98/5 (EC 90/4	e undersigned, declare that I am aware of t iament and of the Council of 28 January 3 ablishing the European Food Safety Author 2002, p. 1), Regulation (EC) No 852/2004 iene of foodstuffs (OJ L 139, 30.4.2004, p. Council of 29 April 2004 laying down specifi julation (EU) 2017/625 of the European Parl ial activities performed to ensure the appl Ith and plant protection products, amending) No 1107/2009, (EU) No 1151/2012, (EL iament and of the Council, Council Regula 58/EC, 1999/74/EC, 2007/43/EC, 2008/119/) No 882/2004 of the European Parliamo (25/EEC, 91/496/EEC, 96/23/EC, 96/93/E julation) (OJ L 95, 7.4.2017, p. 1), and	2002 laying down the general princ ority and laying down procedures i of the European Parliament and of 1) and Regulation (EC) No 853/2004 c hygiene rules for food of animal or liament and of the Council of 15 Mard ication of food and feed law, rules Regulations (EC) No 999/2001, (EC J) No 652/2014, (EU) 2016/429 an ations (EC) No 1/2005 and (EC) No /EC and 2008/120/EC and repealing ent and of the Council, Council D	iples and requirements of food law, n matters of food safety (OJ L 31, the Council of 29 April 2004 on the 4 of the European Parliament and of gin (OJ L 139, 30.4.2004, p. 55) and ch 2017 on official controls and other on animal health and welfare, plant c) No 396/2005, (EC) No 1069/2009, d (EU) 2016/2031 of the European o 1099/2009 and Council Directives Regulations (EC) No 854/2004 and irectives 89/608/EEC, 89/662/EEC,
		l cei	rtify that the snails described above were pr	oduced in accordance with these req	uirements, in particular that they:
		_	come from (an) establishment(s) implem points (HACCP) principles in accordance		
		—	have been handled and, where appropria with the requirements of Annex II to Regul		in a hygienic manner in accordance
	_	—	have been handled and, where appropriat a hygienic manner in accordance with the		
	Note	S			
	certif	ficates	in Annex II of Commission Implementing for certain animals and goods and ame '59 as regards these model certificates (C	ending Regulation (EC) No 2074/2	
	Part	l:			
	_	Box re	ference I.25: Insert the appropriate HS/CN o	code(s) such as: 0307 60 00 or 1605.	
	-	Box re	ference I.25: <i>Treatment type</i> : fresh, treated.		
	Part	II:			
	-	The co	lour of the stamp and signature must be dif	ferent from that of the other particula	rs in the certificate.
	Offici	ial inspe	ector		
		Name	(in capital letters):	Q	ualification and title:
		Date:		Si	gnature:
		Stamp	:		
	L				

PART V

MODEL OFFICIAL CERTIFICATE FOR THE ENTRY INTO THE UNION FOR PLACING ON THE MARKET OF RENDERED ANIMAL FATS AND GREAVES INTENDED FOR HUMAN CONSUMPTION

cou	NTRY										Official certificate to the EU
	l.1.							1.2.	Certificate referen	ce No	I.2.a IMSOC reference No
		Name							I.3. Central Competent Authority		
		Address							Local Competent Authority		
ment		Tel. No									
signı	1.5.	Consignee/Importer						I.6.	Operator responsi	ble for th	ne consignment
hed con		Name							Name		
patcl		Address							Address		
f dis		Postal code							Postal code		
ails o		Tel. No									
Part I: Details of dispatched consignment	1.7.	Country of origin	ISO	1.8.			1.9.	Country of destination	ISO	I.10.	
Ρ	1.11	Place of dispatch					I.12.	Place of destination	on		
		Name		Approv	al No				Name		
		Address							Address		
	l.13.	Place of loading						l.14.	14. Date and time of departure		
	l.15.	Means of transport						l.16.	Entry BCP		
		Aeroplane	Ves	sel		Other		l.17.	Accompanying do	cuments	
		Road vehicle	Rai	lway					Туре		
		Identification:							No		
	l.18.	Transport conditions									
		Ambient	Chi	lled		Frozen					
	l.19.	Container No/Seal No	I			I		I			

COUNTRY				Official certificate to the EU
I.20. Goods certified as				
Human consumption				
l.21.			1.22.	
I.23. Total number of packages	1.24. 0	Quantity		
	ד	otal number	Total net weight (Kg)	Total gross weight (Kg)
I.25. Description of goods				
No Code and CN	title			
Species (Scientific name)		Ма	nufacturing plant	Cold store
Final consumer Number package		Net weight	Batch No	Type of packaging

COUNTRY Rendered animal fats and greaves intended for human consumption П. **Health information** II.a. Certificate reference No II.b. II.1. Public health attestation I, the undersigned, declare that I am aware of the relevant provisions of Regulation (EC) No 178/2002 of the European Parliament and of the Council of 28 January 2002 laying down the general principles and requirements of food law, establishing the European Food Safety Authority and laying down procedures in matters of food safety (OJ L 31, 1.2.2002, p. 1), Regulation (EC) No 852/2004 of the European Parliament and of the Council of 29 April 2004 on the hygiene of foodstuffs (OJ L 139, 30.4.2004, p. 1) and Regulation (EC) No 853/2004 of the European Parliament and of the Council of 29 April 2004 laying down specific hygiene rules for food of animal origin (OJ L 139, 30.4.2004, p. 55) and Regulation (EU) 2017/625 of the European Parliament and of the Council of 15 March 2017 on official controls and other official activities performed to ensure the application of food and feed law, rules on animal health and welfare, plant Part II: Certification health and plant protection products, amending Regulations (EC) No 999/2001, (EC) No 396/2005, (EC) No 1069/2009, (EC) No 1107/2009, (EU) No 1151/2012, (EU) No 652/2014, (EU) 2016/429 and (EU) 2016/2031 of the European Parliament and of the Council, Council Regulations (EC) No 1/2005 and (EC) No 1099/2009 and Council Directives 98/58/EC, 1999/74/EC, 2007/43/EC, 2008/119/EC and 2008/120/EC and repealing Regulations (EC) No 854/2004 and (EC) No 882/2004 of the European Parliament and of the Council, Council Directives 89/608/EEC, 89/662/EEC, 90/425/EEC, 91/496/EEC, 96/23/EC, 96/93/EC and 97/78/EC and Council Decision 92/438/EEC (Official Controls Regulation) (OJ L 95, 7.4.2017, p. 1), and I certify that the rendered animal fats and greaves described above were produced in accordance with these requirements, in particular: that they come from (an) establishment(s) implementing a programme based on the hazard analysis and critical control points (HACCP) principles in accordance with Article 5 of Regulation (EC) No 852/2004; that they have been handled and, where appropriate, prepared, packaged and stored in a hygienic manner in accordance with the requirements of Annex II to Regulation (EC) No 852/2004; and that they comply with the requirements of Section XII of Annex III to Regulation (EC) No 853/2004. II.2. Animal health attestation I, the undersigned official veterinarian, hereby certify, that the rendered animal fats and greaves described above meet the following requirements and come from II.2.1. either third countries, territories and parts thereof appearing in the list authorised for export to the Union of fresh meat in accordance with Part I, of Annex II to Commission Regulation (EU) No 206/2010 of 12 March 2010 laying down lists of third countries, territories or parts thereof authorised for the introduction into the European Union of certain animals and fresh meat and the veterinary certification requirements (OJ L 73, 20.3.2010, p1); II.2.1. or third countries, territories and parts thereof authorised for export to the Union of fresh meat of poultry in accordance with Part 1, of Annex I to Commission Regulation (EC) No 798/2008 of 8 August 2008 laying down a list of third countries, territories, zones or compartments from which poultry and poultry products may be imported into and transit through the Community and the veterinary certification requirements (OJ L 226, 23.8.2008, p. 1); II.2.1. or third countries, territories and parts thereof authorised for export to the Union of meat products of the species of concern subject to the application of the treatment specified for the animal species of origin of the meat product and set out in the list of third countries and territories in Part 1, of Annex II of Commission Decision 2007/777/EC of 29 November 2007 laying down the animal and public health conditions and model certificates for imports of certain meat products and treated stomachs, bladders and intestines for human consumption from third countries and repealing Decision 2005/432/EC (OJ L 312, 30.11.2007, p. 49). Notes See notes in Annex II of Commission Implementing Regulation (EU) 2019/628 of 8 April 2019 concerning model official certificates for certain animals and goods and amending Regulation (EC) No 2074/2005 and Implementing Regulation (EU) 2016/759 as regards these model certificates (OJ L 131, 17.5.2019, p. 101). Part I: Box reference I.25: Insert the appropriate HS/CN code(s) such as: 1501, 1502, 1503 00, 1504, 1506 00 00, 1516 10, 1517, 1518 00 91, 1518 00 95, 1518 00 99 or 2301.

Health information	II.a. Certificate reference No							
		II.b.						
e colour of the stamp and signature must be diffe	erent from that of the other particulars	s in the certificate.						
Official veterinarian								
me (in capital letters):	Qu	alification and title:						
te:	Sig	inature:						
imp:								
r t	eterinarian ne (in capital letters): e:	ne (in capital letters): Qu e: Sig						

PART VI

MODEL OFFICIAL CERTIFICATE FOR THE ENTRY INTO THE UNION FOR PLACING ON THE MARKET OF GELATINE INTENDED FOR HUMAN CONSUMPTION

cou	NTRY									Official certificate to the EU			
	l.1.	Consignor/Exporter						I.2.	Certificate referen	ce No	I.2.a IMSOC reference No		
		Name						1.3.	Central Competer	nt Author	ity		
		Address						1.4.	Local Competent	Authority	,		
ment		Tel. No											
sign	I.5.	Consignee/Importer						1.6.	Operator respons	ible for th	ne consignment		
ned con		Name							Name				
patch		Address							Address				
f dis		Postal code							Postal code				
uils o		Tel. No	No										
Part I: Details of dispatched consignment	1.7.	Country of origin	ISO	1.8.				1.9.	Country of destination	ISO	I.10.		
Ра	l.11	Place of dispatch						I.12.	Place of destination	on			
		Name		Approv	val No				Name				
		Address						Address					
	l.13.	Place of loading						l.14.	I.14. Date and time of departure		•		
	I.15. Means of transport Aeroplane Vess Road vehicle Raily						l.16.	Entry BCP					
			Ves	sel		Other 🗖		I.17.	Accompanying do	documents			
			Rai	Iway 🗖					Туре				
		Identification:							No				
	l.18.	Transport conditions											
		Ambient	Chi	lled		Frozen							
	l.19.	Container No/Seal No						1					

COUNTRY				Official certificate to the EU
I.20. Goods certified as				
Human consumption				
I.21.			1.22.	
I.23. Total number of packages	1.24. 0	Quantity		
	Т	otal number	Total net weight (Kg)	Total gross weight (Kg)
I.25. Description of goods				
No Code and CN	title			
Species (Scientific name)		Ма	nufacturing plant	Cold store
Final consumer Number package		Net weight	Batch No	Type of packaging

Part II: Certification

EN

COUNT	RY	Gelatine i	Model GEL ntended for human consumption
II.	Health information	II.a. Certificate reference No	II.b.

II.1. Public health attestation

I, the undersigned, declare that I am aware of the relevant provisions of Regulation (EC) No 178/2002 of the European Parliament and of the Council of 28 January 2002 laying down the general principles and requirements of food law, establishing the European Food Safety Authority and laying down procedures in matters of food safety (OJ L 31, 1.2.2002, p. 1), Regulation (EC) No 852/2004 of the European Parliament and of the Council of 29 April 2004 on the hygiene of foodstuffs (OJ L 139, 30.4.2004, p. 1) and Regulation (EC) No 853/2004 of the European Parliament and of the Council of 29 April 2004 laying down specific hygiene rules for food of animal origin (OJ L 139, 30.4.2004, p. 55) and Regulation (EU) 2017/625 of the European Parliament and of the Council of 15 March 2017 on official controls and other official activities performed to ensure the application of food and feed law, rules on animal health and welfare, plant health and plant protection products, amending Regulations (EC) No 999/2001, (EC) No 396/2005, (EC) No 1069/2009, (EC) No 1107/2009, (EU) No 1151/2012, (EU) No 652/2014, (EU) 2016/429 and (EU) 2016/2031 of the European Parliament and of the Council, Council Regulations (EC) No 1/2005 and (EC) No 1999/2009 and Council Directives 98/58/EC, 1999/74/EC, 2007/43/EC, 2008/119/EC and 2008/120/EC and repealing Regulations (EC) No 854/2004 and (EC) No 854/2004 of the European Parliament and of the Council, Council Directives 89/608/EEC, 89/662/EEC, 90/425/EEC, 91/496/EEC, 96/23/EC, and 97/78/EC and Council Decision 92/438/EEC (Official Controls Regulation) (OJ L 95, 7.4.2017, p. 1), and

I certify that the gelatine described above was produced in accordance with these requirements, in particular that:

- it comes from (an) establishment(s) implementing a programme based on the hazard analysis and critical control points (HACCP) principles in accordance with Article 5 of Regulation (EC) No 852/2004;
- it has been produced from raw materials that met the requirements of Chapters I and II of Section XIV of Annex III to Regulation (EC) No 853/2004;
- it has been manufactured in compliance with the conditions set out in Chapter III of Section XIV of Annex III to Regulation (EC) No 853/2004;
- it satisfies the criteria of Chapter IV of Section XIV of Annex III to Regulation (EC) No 853/2004 and of Commission Regulation (EC) No 2073/2005 of 15 November 2005 on microbiological criteria for foodstuffs (OJ L 338, 22.12.2005, p. 1);

(¹) and, if of bovine, ovine and caprine animal origin,

it has been derived from animals which have passed ante-mortem and post-mortem inspections,

(1) and, except for gelatine derived from hides and skins,

(¹) either

- [it comes from a country or a region classified in accordance with Commission Decision 2007/453/EC of 29 June 2007 establishing the BSE status of Member States or third countries or regions thereof according to their BSE risk (OJ L 172, 30.6.2007, p. 84) as a country or region posing a negligible BSE risk;
- the gelatine does not contain and is not derived from 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 of 22 May 2001 laying down rules for the prevention, control and eradication of certain transmissible spongiform encephalopathies (OJ L 147, 31.5.2001, p. 1) (²);
- the gelatine does not contain and is not derived from mechanically separated meat obtained from the bones of bovine, ovine or caprine animals, except for gelatine derived from animals that were born, continuously reared and slaughtered in a country or region classified in accordance with Decision 2007/453/EC as a country or region posing a negligible BSE risk in which there has been no indigenous BSE cases;
- the animals, from which the gelatine is derived were not slaughtered after stunning by means of gas injected into the cranial cavity or killed by the same method or slaughtered, after stunning, by laceration of central nervous tissue by means of an elongated rod-shaped instrument introduced into the cranial cavity, except if the animals were born, continuously reared and slaughtered in a country or region classified in accordance with Decision 2007/453/EC as a country or region posing a negligible BSE risk;
- (1) [the animals, from which the gelatine is derived, originate from a country or region classified in accordance with Decision 2007/453/EC as a country or region posing an undetermined BSE risk, and the animals were not fed with meat-and-bone meal or greaves, as defined in the Terrestrial Animal Health Code of the World Organisation for Animal Health];

 Decision 2007/453/EC as a country or region posing an undetermined BSE risk, and the gelatine was p and handled in a manner which ensures that it did not contain and was not contaminated with nerve lymphatic tissues exposed during the deboning process.]] (1) Or [1] (comes from a country or a region classified in accordance with Commission Decision 2007/453/EC of 2007 establishing the BSE status of Member States or third countries or regions thereof according to their I (CJJ L 172, 30.6.2007, p. 84) as a country or region posing a controlled BSE risk; the animals, from which the gelatine is derived, were not killed, after stunning, by laceration of central tissue by means of an elongated rod-shaped instrument introduced into the cranial cavity, or by means injected into the cranial cavity. the gelatine does not contain and is not derived from specified risk material as defined in point 1 of Ant Regulation (EC) No 999/2001, or mechanically separated meat obtained from the bones of bovine, a caprine animals.] (1) Or [1] (1) Or [1] to comes from a country or a region classified in accordance with Decision 2007/453/EC of 29 Jur establishing the BSE status of Member States or third countries or regions thereof according to their E (OJ L 172, 30.6.2007, p. 84) as a country or region with an undetermined BSE risk; the animals, from which the gelatine is derived, were not fed meat-and-bone meal or greaves deriv ruminants, as defined in the Terrestrial Animal health Code of the World Organisation for Animal Health; the animals, from which the gelatine is derived, were not killed, after stunning, by laceration of central tissue by means of an elongated rod-shaped instrument introduced into the cranial cavity, or by means injected into the cranial cavity; the animals, from which the gelatine is derived, were not killed, after stunning, by laceration of central tissue by means of an elongated rod-shaped instrument introduced i	 (1) [the animals, from which the gelatine is derived, originate from a country or region classified in accordance Decision 2007/453/EC as a country or region posing an undetermined BSE risk, and the gelatine was prod and handled in a manner which ensures that it dd not contain and was not contaminated with nervous lymphatic tissues exposed during the deboning process.]] (1) Or (1) Or (1) Cr (2) Cr (2) Cr (2) Cr (3) Cr (4) Cr (4) Cr (5) Cr (4) Cr (5) Cr (6) L 172, 30.6.2007, p. 84) as a country or region posing a controlled BSE risk. (6) L 172, 30.6.2007, p. 84) as a country or region posing a controlled BSE risk. (7) Cr (8) the animals, from which the gelatine is derived, were not killed, after stunning, by laceration of central ner tissue by means of an elongated rod-shaped instrument introduced into the cranil cavity, or by means of injected into the cranil cavity, in the chain and is not derived from specified risk material as defined in point 1 of Annex Regulation (EC) No 999/2001, or mechanically separated meat obtained from the bones of bovine, ovir caprine animals.] (1) Or (1) Cr (1) Cr (1) Cr (2) Cr (3) Cr (4) Cranis active of a country or a region classified in accordance with Decision 2007/453/EC of 29 June : establishing the BSE status of Member States or third countries or regions thereof according to their BSE (CU 1172, 30, 6207, p. 64) as a country or region owith a undetermined BSE risk. (5) The animals	П.	JNTRY			Gelatine intended for human consumption					
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 ruminants, as defined in the Terrestrial Animal health Code of the World Organisation for Animal Health; the animals, from which the gelatine is derived, were not killed, after stunning, by laceration of central tissue by means of an elongated rod-shaped instrument introduced into the cranial cavity, or by means injected into the cranial cavity; the gelatine is not derived from: specified risk material as defined in point 1 of Annex V to Regulation (EC) No 999/2001; nervous and lymphatic tissues exposed during the deboning process; ini) mechanically separated meat obtained from the bones of bovine, ovine or caprine animals. Notes See notes in Annex II of Commission Implementing Regulation (EU) 2019/628 of 8 April 2019 concerning model certificates for certain animals and goods and amending Regulation (EC) No 2074/2005 and Implementing Regulation (EU) 2016/759 as regards these model certificates (OJ L 131, 17.5.2019, p. 101). Part I: Box reference I.25: Insert the appropriate Harmonised System (HS) code(s) using headings such as 3503. 	 ruminants, as defined in the Terrestrial Animal health Code of the World Organisation for Animal Health; the animals, from which the gelatine is derived, were not killed, after stunning, by laceration of central ner tissue by means of an elongated rod-shaped instrument introduced into the cranial cavity, or by means of injected into the cranial cavity; the gelatine is not derived from: (i) specified risk material as defined in point 1 of Annex V to Regulation (EC) No 999/2001; (ii) nervous and lymphatic tissues exposed during the deboning process; (iii) mechanically separated meat obtained from the bones of bovine, ovine or caprine animals. Notes See notes in Annex II of Commission Implementing Regulation (EU) 2019/628 of 8 April 2019 concerning model off cartificates for certain animals and goods and amending Regulation (EC) No 2074/2005 and Implementing Regulation (EU) 2019/6759 as regards these model certificates (OJ L 131, 17.5.2019, p. 101). Part I: Box reference 1.25: Insert the appropriate Harmonised System (HS) code(s) using headings such as 3503. Part II: The removal of specified risk material is not required if the gelatine is derived from animals born, continuously reared slaughtered in a third country or region of a third country classified in accordance with Decision 2007/453/EC as posin negligible BSE risk. The colour of the stamp and signature must be different from that of the other particulars in the certificate. Official veterinarian Name (in capital letters): Qualification and title: 		 [it comes from a country or a region classified in accordance with Decision 2007/453/EC of 29 June 2007 establishing the BSE status of Member States or third countries or regions thereof according to their BSE risk (OJ L 172, 30.6.2007, p. 84) as a country or region with an undetermined BSE risk; 								
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 (i) specified risk material as defined in point 1 of Annex V to Regulation (EC) No 999/2001; (ii) nervous and lymphatic tissues exposed during the deboning process; (iii) mechanically separated meat obtained from the bones of bovine, ovine or caprine animals. Notes See notes in Annex II of Commission Implementing Regulation (EU) 2019/628 of 8 April 2019 concerning model certificates for certain animals and goods and amending Regulation (EC) No 2074/2005 and Implementing Regulation (EU) 2016/759 as regards these model certificates (OJ L 131, 17.5.2019, p. 101). Part I: — Box reference I.25: Insert the appropriate Harmonised System (HS) code(s) using headings such as 3503. 	 (i) specified risk material as defined in point 1 of Annex V to Regulation (EC) No 999/2001; (ii) nervous and lymphatic tissues exposed during the deboning process; (iii) mechanically separated meat obtained from the bones of bovine, ovine or caprine animals. Notes See notes in Annex II of Commission Implementing Regulation (EU) 2019/628 of 8 April 2019 concerning model off perificates for certain animals and goods and amending Regulation (EC) No 2074/2005 and Implementing Regulation (EU) 2016/759 as regards these model certificates (OJ L 131, 17.5.2019, p. 101). Part I: Box reference 1.25: Insert the appropriate Harmonised System (HS) code(s) using headings such as 3503. Part II: 1) Delete as appropriate. 2) The removal of specified risk material is not required if the gelatine is derived from animals born, continuously reared slaughtered in a third country or region of a third country classified in accordance with Decision 2007/453/EC as posinegligible BSE risk. — The colour of the stamp and signature must be different from that of the other particulars in the certificate. Official veterinarian Name (in capital letters): Qualification and title:		tissue by means of an elongated rod-shaped instrument introduced into the cranial cavity, or by means of gas								
 (ii) nervous and lymphatic tissues exposed during the deboning process; (iii) mechanically separated meat obtained from the bones of bovine, ovine or caprine animals. Notes See notes in Annex II of Commission Implementing Regulation (EU) 2019/628 of 8 April 2019 concerning model certificates for certain animals and goods and amending Regulation (EC) No 2074/2005 and Implementing Regulation (EU) 2016/759 as regards these model certificates (OJ L 131, 17.5.2019, p. 101). Part I: — Box reference I.25: Insert the appropriate Harmonised System (HS) code(s) using headings such as 3503.	 (ii) nervous and lymphatic tissues exposed during the deboning process; (iii) mechanically separated meat obtained from the bones of bovine, ovine or caprine animals. Notes See notes in Annex II of Commission Implementing Regulation (EU) 2019/628 of 8 April 2019 concerning model off perifficates for certain animals and goods and amending Regulation (EC) No 2074/2005 and Implementing Regulation (EU) 2016/759 as regards these model certificates (OJ L 131, 17.5.2019, p. 101). Part I: Box reference 1.25: Insert the appropriate Harmonised System (HS) code(s) using headings such as 3503. Part II: 1) Delete as appropriate. 2) The removal of specified risk material is not required if the gelatine is derived from animals born, continuously reared slaughtered in a third country or region of a third country classified in accordance with Decision 2007/453/EC as posineligible BSE risk. — The colour of the stamp and signature must be different from that of the other particulars in the certificate. Official veterinarian Name (in capital letters): Qualification and title:			the	gelatine is not derived from:						
 (iii) mechanically separated meat obtained from the bones of bovine, ovine or caprine animals. Notes See notes in Annex II of Commission Implementing Regulation (EU) 2019/628 of 8 April 2019 concerning model certificates for certain animals and goods and amending Regulation (EC) No 2074/2005 and Implementing Regulation (EU) 2016/759 as regards these model certificates (OJ L 131, 17.5.2019, p. 101). Part I: Box reference I.25: Insert the appropriate Harmonised System (HS) code(s) using headings such as 3503. 	 (iii) mechanically separated meat obtained from the bones of bovine, ovine or caprine animals. Notes See notes in Annex II of Commission Implementing Regulation (EU) 2019/628 of 8 April 2019 concerning model officertificates for certain animals and goods and amending Regulation (EC) No 2074/2005 and Implementing RegulateU) 2016/759 as regards these model certificates (OJ L 131, 17.5.2019, p. 101). Part I: Box reference 1.25: Insert the appropriate Harmonised System (HS) code(s) using headings such as 3503. Part II: Delete as appropriate. The removal of specified risk material is not required if the gelatine is derived from animals born, continuously reared slaughtered in a third country or region of a third country classified in accordance with Decision 2007/453/EC as posineligible BSE risk. The colour of the stamp and signature must be different from that of the other particulars in the certificate. Official veterinarian Name (in capital letters): Qualification and title: 			(i)	specified risk material as defined in p	point 1 of Annex V to Regulation (EC)	No 999/2001;				
Notes See notes in Annex II of Commission Implementing Regulation (EU) 2019/628 of 8 April 2019 concerning model certificates for certain animals and goods and amending Regulation (EC) No 2074/2005 and Implementing Reg EU) 2016/759 as regards these model certificates (OJ L 131, 17.5.2019, p. 101). Part I: — Box reference I.25: Insert the appropriate Harmonised System (HS) code(s) using headings such as 3503. Part II:	Notes See notes in Annex II of Commission Implementing Regulation (EU) 2019/628 of 8 April 2019 concerning model off certificates for certain animals and goods and amending Regulation (EC) No 2074/2005 and Implementing Regulated EU) 2016/759 as regards these model certificates (OJ L 131, 17.5.2019, p. 101). Part I: — Box reference I.25: Insert the appropriate Harmonised System (HS) code(s) using headings such as 3503. Part II: 1) Delete as appropriate. 2) The removal of specified risk material is not required if the gelatine is derived from animals born, continuously reared slaughtered in a third country or region of a third country classified in accordance with Decision 2007/453/EC as posine negligible BSE risk. — The colour of the stamp and signature must be different from that of the other particulars in the certificate. Difficial veterinarian Name (in capital letters): Qualification and title:			(ii)	nervous and lymphatic tissues expos	sed during the deboning process;					
See notes in Annex II of Commission Implementing Regulation (EU) 2019/628 of 8 April 2019 concerning model certificates for certain animals and goods and amending Regulation (EC) No 2074/2005 and Implementing Reg (EU) 2016/759 as regards these model certificates (OJ L 131, 17.5.2019, p. 101). Part I: — Box reference I.25: Insert the appropriate Harmonised System (HS) code(s) using headings such as 3503. Part II:	 See notes in Annex II of Commission Implementing Regulation (EU) 2019/628 of 8 April 2019 concerning model of certificates for certain animals and goods and amending Regulation (EC) No 2074/2005 and Implementing Regulate) 2016/759 as regards these model certificates (OJ L 131, 17.5.2019, p. 101). Part I: Box reference I.25: Insert the appropriate Harmonised System (HS) code(s) using headings such as 3503. Part II: Delete as appropriate. *) Delete as appropriate. *) The removal of specified risk material is not required if the gelatine is derived from animals born, continuously reared slaughtered in a third country or region of a third country classified in accordance with Decision 2007/453/EC as posinelyible BSE risk. The colour of the stamp and signature must be different from that of the other particulars in the certificate. Official veterinarian Name (in capital letters): Qualification and title:			(iii)	mechanically separated meat obtain	ed from the bones of bovine, ovine or	caprine animals.				
See notes in Annex II of Commission Implementing Regulation (EU) 2019/628 of 8 April 2019 concerning model certificates for certain animals and goods and amending Regulation (EC) No 2074/2005 and Implementing Reg (EU) 2016/759 as regards these model certificates (OJ L 131, 17.5.2019, p. 101). Part I: — Box reference I.25: Insert the appropriate Harmonised System (HS) code(s) using headings such as 3503. Part II:	 See notes in Annex II of Commission Implementing Regulation (EU) 2019/628 of 8 April 2019 concerning model of certificates for certain animals and goods and amending Regulation (EC) No 2074/2005 and Implementing Regulated (EU) 2016/759 as regards these model certificates (OJ L 131, 17.5.2019, p. 101). Part I: Box reference I.25: Insert the appropriate Harmonised System (HS) code(s) using headings such as 3503. Part II: Delete as appropriate. (1) Delete as appropriate. (2) The removal of specified risk material is not required if the gelatine is derived from animals born, continuously reared slaughtered in a third country or region of a third country classified in accordance with Decision 2007/453/EC as posinegligible BSE risk. The colour of the stamp and signature must be different from that of the other particulars in the certificate. Official veterinarian Name (in capital letters): Qualification and title: 										
 Box reference I.25: Insert the appropriate Harmonised System (HS) code(s) using headings such as 3503. Part II: 	 Box reference I.25: Insert the appropriate Harmonised System (HS) code(s) using headings such as 3503. Part II: Delete as appropriate. The removal of specified risk material is not required if the gelatine is derived from animals born, continuously reared slaughtered in a third country or region of a third country classified in accordance with Decision 2007/453/EC as posin negligible BSE risk. The colour of the stamp and signature must be different from that of the other particulars in the certificate. Official veterinarian Name (in capital letters): Qualification and title: 	See cer	e notes i tificates	for o	ertain animals and goods and am	ending Regulation (EC) No 2074/20	-				
Part II:	 Part II: ¹) Delete as appropriate. ²) The removal of specified risk material is not required if the gelatine is derived from animals born, continuously reared slaughtered in a third country or region of a third country classified in accordance with Decision 2007/453/EC as posinegligible BSE risk. The colour of the stamp and signature must be different from that of the other particulars in the certificate. Difficial veterinarian Name (in capital letters): 		tl:								
	 (1) Delete as appropriate. (2) The removal of specified risk material is not required if the gelatine is derived from animals born, continuously reared slaughtered in a third country or region of a third country classified in accordance with Decision 2007/453/EC as posinegligible BSE risk. The colour of the stamp and signature must be different from that of the other particulars in the certificate. Official veterinarian Name (in capital letters): 	Par									
	 (1) Delete as appropriate. (2) The removal of specified risk material is not required if the gelatine is derived from animals born, continuously reared slaughtered in a third country or region of a third country classified in accordance with Decision 2007/453/EC as posinegligible BSE risk. The colour of the stamp and signature must be different from that of the other particulars in the certificate. Official veterinarian Name (in capital letters): 	Par 	Box ret	ferend	ce I.25: Insert the appropriate Harmon	ised System (HS) code(s) using headi	ngs such as 3503.				
) Delete as appropriate.	 (2) The removal of specified risk material is not required if the gelatine is derived from animals born, continuously reared slaughtered in a third country or region of a third country classified in accordance with Decision 2007/453/EC as posinegligible BSE risk. The colour of the stamp and signature must be different from that of the other particulars in the certificate. Official veterinarian Name (in capital letters): 			ferend	e I.25: Insert the appropriate Harmon	ised System (HS) code(s) using headi	ngs such as 3503.				
(2) The removal of specified risk material is not required if the gelatine is derived from animals born, continuously reasonable slaughtered in a third country or region of a third country classified in accordance with Decision 2007/453/EC as provide the statement of the sta	 The colour of the stamp and signature must be different from that of the other particulars in the certificate. Official veterinarian Name (in capital letters): Qualification and title: 	 Par	t II:			ised System (HS) code(s) using headi	ngs such as 3503.				
	Name (in capital letters): Qualification and title:	— Par (¹)	t II: Delete The re slaugh	as ap mova tered	propriate. I of specified risk material is not requi in a third country or region of a third	red if the gelatine is derived from anir	nals born, continuously reared ar				
	Name (in capital letters): Qualification and title:	- Par	t II: Delete The rei slaught negligil	as ar mova tered ble B	ppropriate. I of specified risk material is not requi in a third country or region of a third SE risk.	red if the gelatine is derived from anir country classified in accordance with	nals born, continuously reared ar Decision 2007/453/EC as posing				
-		Par	t II: Delete The re slaught negligil The co	as ar mova tered ble B lour c	ppropriate. I of specified risk material is not requi in a third country or region of a third SE risk. If the stamp and signature must be diff	red if the gelatine is derived from anir country classified in accordance with	nals born, continuously reared an Decision 2007/453/EC as posing				
	Date: Signature:	Par	t II: Delete The rei slaught negligil The co	as ap mova tered ble B lour c rinaria	ppropriate. I of specified risk material is not requi in a third country or region of a third SE risk. If the stamp and signature must be diff	red if the gelatine is derived from anir country classified in accordance with ferent from that of the other particulars	nals born, continuously reared an Decision 2007/453/EC as posing in the certificate.				
Liate: Pianoturo		Par (¹) (²)	t II: Delete The rei slaught negligil The co cial veter Name	as ap mova tered ble B lour c rinaria	ppropriate. I of specified risk material is not requi in a third country or region of a third SE risk. If the stamp and signature must be diff	red if the gelatine is derived from anir country classified in accordance with ferent from that of the other particulars	mals born, continuously reared ar Decision 2007/453/EC as posing in the certificate.				
Date: Signature: Signature:	Stamp.	Par (¹) (²)	t II: Delete The rei slaught negligil The co cial veter Name Date:	as ar mova tered ble B lour c rinaria (in ca	ppropriate. I of specified risk material is not requi in a third country or region of a third SE risk. If the stamp and signature must be diff	red if the gelatine is derived from anir country classified in accordance with ferent from that of the other particulars	mals born, continuously reared a Decision 2007/453/EC as posing in the certificate.				

PART VII

MODEL OFFICIAL CERTIFICATE FOR THE ENTRY INTO THE UNION FOR PLACING ON THE MARKET OF COLLAGEN INTENDED FOR HUMAN CONSUMPTION

cou	NTRY							Official certificate to the EU			
	l.1.	Consignor/Exporter						1.2.	Certificate referen	ice No	I.2.a IMSOC reference No
		Name						1.3.	Central Competer	nt Author	ity
		Address						1.4.	Local Competent	Authority	,
ment		Tel. No									
sign	1.5.	Consignee/Importer						I.6.	Operator respons	ible for th	ne consignment
ned con		Name							Name		
patcł		Address							Address		
f dis		Postal code							Postal code		
uils o		Tel. No									
Part I: Details of dispatched consignment	I.7. Country of origin ISO		ISO	1.8.				1.9.	Country of destination	ISO	I.10.
Ра	l.11	Place of dispatch						I.12.	Place of destination	on	
		Name		Approv	al No				Name		
	Address							Address			
	I.13. Place of loading							l.14.	Date and time of o	departure	•
	l.15.	Means of transport						I.16.	Entry BCP		
			Ves	sel		Other 🗖		l.17.	I.17. Accompanying documents		;
			Rai	ilway 🗖					Туре		
		Identification:							No		
	l.18.	Transport conditions									
		Ambient	Chi	lled		Frozen					
	l.19.	Container No/Seal No						1			

COUNTRY				Official certificate to the EU
I.20. Goods certified as				
Human consumption				
I.21.			1.22.	
I.23. Total number of packages	1.24. 0	Quantity		
	т	otal number	Total net weight (Kg)	Total gross weight (Kg)
I.25. Description of goods				
No Code and CN	title			
Species (Scientific name)		Ма	nufacturing plant	Cold store
Final consumer Number package		Net weight	Batch No	Type of packaging

Part II: Certification

EN

COUNT	RY		Collagen i	Model COL ntended for human consumption
П.	Health information	II.a.	Certificate reference No	II.b.
II.1.	Public health attestation			

I, the undersigned, declare that I am aware of the relevant provisions of Regulation (EC) No 178/2002 of the European Parliament and of the Council of 28 January 2002 laying down the general principles and requirements of food law, establishing the European Food Safety Authority and laying down procedures in matters of food safety (OJ L 31, 1.2.2002, p. 1), Regulation (EC) No 852/2004 of the European Parliament and of the Council of 29 April 2004 on the hygiene of foodstuffs (OJ L 139, 30.4.2004, p. 1) and Regulation (EC) No 853/2004 of the European Parliament and of the Council of 29 April 2004 laying down specific hygiene rules for food of animal origin (OJ L 139, 30.4.2004, p. 55) and Regulation (EU) 2017/625 of the European Parliament and of the Council of 15 March 2017 on official controls and other official activities performed to ensure the application of food and feed law, rules on animal health and welfare, plant health and plant protection products, amending Regulations (EC) No 999/2001, (EC) No 396/2005, (EC) No 1069/2009, (EC) No 1107/2009, (EU) No 1151/2012, (EU) No 652/2014, (EU) 2016/29 and (EU) 2016/2031 of the European Parliament and of the Council, Council Council Council Directives 98/58/EC, 1999/74/EC, 2007/43/EC, 2008/119/EC and 2008/120/EC and repealing Regulations (EC) No 854/2004 and (EC) No 854/2004 of the European Parliament and of the Council, Council Directives 89/608/EEC, 89/602/EEC, 90/425/EEC, 91/496/EEC, 96/93/EC and 97/78/EC and Council Decision 92/438/EEC (Official Controls Regulation) (OJ L 95, 7.4.2017, p. 1), and

I certify that the collagen described above was produced in accordance with these requirements, in particular that:

- it comes from (an) establishment(s) implementing a programme based on the hazard analysis and critical control points (HACCP) principles in accordance with Article 5 of Regulation (EC) No 852/2004;
- it has been produced from raw materials that met the requirements of Chapters I and II of Section XV of Annex III to Regulation (EC) No 853/2004;
- it has been manufactured in compliance with the conditions set out in Chapter III of Section XV of Annex III to Regulation (EC) No 853/2004;
- it satisfies the criteria of Chapter IV of Section XV of Annex III to Regulation (EC) No 853/2004 and of Commission Regulation (EC) No 2073/2005 of 15 November 2005 on microbiological criteria for foodstuffs (OJ L 338, 22.12.2005, p. 1);
- (¹) and, if of bovine, ovine and caprine animal origin,

it has been derived from animals which have passed ante-mortem and post-mortem inspections,

(1) and, except for collagen derived from hides and skins,

(¹) either

- [it comes from a country or a region classified in accordance with Commission Decision 2007/453/EC of 29 June 2007 establishing the BSE status of Member States or third countries or regions thereof according to their BSE risk (OJ L 172, 30.6.2007, p. 84) as a country or region posing a negligible BSE risk;
- the collagen does not contain and is not derived from 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 of 22 May 2001 laying down rules for the prevention, control and eradication of certain transmissible spongiform encephalopathies (OJ L 147, 31.5.2001, p. 1) (²);
- the collagen does not contain and is not derived from mechanically separated meat obtained from bones of bovine, ovine or caprine animals, except for collagen derived from animals that were born, continuously reared and slaughtered in a country or region classified in accordance with Decision 2007/453/EC as a country or region posing a negligible BSE risk in which there has been no indigenous BSE cases;
- the animals from which the collagen was derived have not been slaughtered after stunning by means of gas injected into the cranial cavity or killed by the same method or slaughtered, after stunning, by laceration of central nervous tissue by means of an elongated rod-shaped instrument introduced into the cranial cavity, except if the animals were born, continuously reared and slaughtered in a country or region classified in accordance with Decision 2007/453/EC as a country or region posing a negligible BSE risk;
- (1) [the animals, from which the collagen is derived, originate from a country or region classified in accordance with Decision 2007/453/EC as a country or region posing an undetermined BSE risk, and the animals were not fed with meat-and-bone meal or greaves, as defined in the OIE Terrestrial Animal Health Code];

odel COL sumption	Model ntended for human consum	Collagen		NTRY	cou	
	II.b.	II.a. Certificate reference No	formation	Health	П.	
produced	k, and the collagen was prod	gion posing an undetermined BSE r s that it did not contain and was r	he animals, from which the collagen is ision 2007/453/EC as a country or re handled in a manner which ensure phatic tissues exposed during the debo	D		
				(1) or		
	thereof according to their BSE	er States or third countries or region	omes from a country or a region class 7 establishing the BSE status of Memb L 172, 30.6.2007, p. 84) as a country	20		
			animals, from which the collagen is on the by means of an elongated rod-sha ated into the cranial cavity;	tis		
			collagen does not contain and is not ulation (EC) No 999/2001, or mech ine animals.]	R		
				(1) or		
	nereof according to their BSE	States or third countries or regions	omes from a country or a region c blishing the BSE status of Member 3 L 172, 30.6.2007, p. 84) as a country	es		
ived from			animals, from which the collagen is nants, as defined in the Terrestrial An			
			animals, from which the collagen is one of the second of an elongated rod-shated into the cranial cavity;	tis		
 the collagen is not derived from: 						
(i) specified risk material as defined in point 1 of Annex V to Regulation (EC) No 999/2001;						
		ed during the deboning process;	nervous and lymphatic tissues expos	(ii		
	aprine animals.]	d from the bones of bovine, ovine or	mechanically separated meat obtained	(ii		
				25	Note	
		nding Regulation (EC) No 2074/2	nex II of Commission Implementing ertain animals and goods and ame regards these model certificates (C	ificates for	certi	
					Part	
		ed for importing collagen casings.	e I.25: This certificate may also be us	Box refere	—	
7.	igs such as 3504 or 3917.	sed System (HS) code(s) using head	e I.25: Insert the appropriate Harmoni	Box refere	_	
				II:	Part	
			propriate.	Delete as	(¹)	
) The removal of specified risk material is not required if the collagen is derived from animals born, continuously reared and slaughtered in a third country or region of a third country classified in accordance with Decision 2007/453/EC as posing a negligible BSE risk.						
	in the certificate.	erent from that of the other particular	f the stamp and signature must be diff	The colou	_	
			n	ial veterina	Offic	
	ualification and title:	(pital letters):	Name (in o		
	gnature:	5	·	Date:		
	-			Stamp:		

PART VIII

MODEL OFFICIAL CERTIFICATE FOR THE ENTRY INTO THE UNION FOR PLACING ON THE MARKET OF RAW MATERIALS FOR THE PRODUCTION OF GELATINE AND COLLAGEN INTENDED FOR HUMAN CONSUMPTION

cou	NTRY							Official certificate to the EL			
	l.1.	Consignor/Exporter						I.2.	Certificate referen	ice No	I.2.a IMSOC reference No
		Name						1.3.	Central Competer	nt Author	ity
		Address						1.4.	Local Competent	Authority	/
nent		Tel. No									
signr	1.5.	Consignee/Importer						I.6.	Operator responsi	ible for th	ne consignment
ed cor		Name							Name		
patch		Address							Address		
of dis		Postal code							Postal code		
ails		Tel. No				_					
Part I: Details of dispatched consignment	1.7.	Country of origin	ISO	1.8. R	Region	of origin	Code	1.9.	Country of destination	ISO	I.10.
ä	l.11	Place of dispatch					I.12.	Place of destination	on		
		Name		Approv	val No				Name		
	Address								Address		
	1.13.	13. Place of loading							I.14. Date and time of departure		
	I.15.	Means of transport					I.16. Entry BCP				
		Aeroplane	Ves	sel		Other		l.17.	Accompanying do	cuments	3
		Road vehicle	Rai	lway							
									Туре		
		Identification:							No		
	I.18.	3. Transport conditions									
		Ambient 🔲	Chi	lled		Frozen					
	I.19. Container No/Seal No							1			

cou	INTRY				Official certificate to the EU	
1.20.	Goods certified as					
	Human consumption					
I.21.				1.22.		
1.23.	Total number of packages		Quantity			
		T	otal number	Total net weight (Kg)	Total gross weight (Kg)	
1.25.	Description of goods					
	No Code and CN	title				
	Species (Scientific name)		Nat	Nature of commodity		
			Ма			
	Number o package		Net weight	Batch No	Type of packaging	

COUNTRY

П.

II.1.

EN

Official Journal of the European Union Model RCG Raw materials for the production of collagen and gelatine intended for human consumption **Health information** II.a. Certificate reference No II.b. **Public health attestation** I, the undersigned, declare that I am aware of the relevant provisions of Regulation (EC) No 178/2002 of the European Parliament and of the Council of 28 January 2002 laying down the general principles and requirements of food law, establishing the European Food Safety Authority and laying down procedures in matters of food safety (OJ L 31, 1.2.2002, p. 1), Regulation (EC) No 852/2004 of the European Parliament and of the Council of 29 April 2004 on the hygiene of foodstuffs (OJ L 139, 30.4.2004, p. 1), Regulation (EC) No 853/2004 of the European Parliament and of the Council of 29 April 2004 laying down specific hygiene rules for food of animal origin (OJ L 139, 30.4.2004, p. 55) and Regulation (EU) 2017/625 of the European Parliament and of the Council of 15 March 2017 on official controls and other official activities performed to ensure the application of food and feed law, rules on animal health and welfare, plant health and plant protection products, amending Regulations (EC) No 999/2001, (EC) No 396/2005, (EC) No 1069/2009, (EC) No 1107/2009, (EU) No 1151/2012, (EU) No 652/2014, (EU) 2016/429 and (EU) 2016/2031 of the European Parliament and of the Council, Council Regulations (EC) No 1/2005 and (EC) No 1099/2009 and Council Directives 98/58/EC, 1999/74/EC, 2007/43/EC, 2008/119/EC and 2008/120/EC and repealing Regulations (EC) No 854/2004 and (EC) No 882/2004 of the European Parliament and of the Council, Council Directives 89/608/EEC, 89/662/EEC, 90/425/EEC, 91/496/EEC, 96/23/EC, 96/93/EC and 97/78/EC and Council Decision 92/438/EEC (Official Controls Regulation) (OJ L 95, 7.4.2017, p. 1), and I certify that the raw materials described above comply with these requirements, in particular that: (1) [bones, hides and skins of domestic and farmed ruminant animals, pigs and poultry, as well as tendons and sinews described above are derived from animals which were slaughtered in a slaughterhouse and the carcasses of which were found to be fit for human consumption following ante- and post-mortem inspection;] (1) [wild game hides, skins and bones described above are derived from killed animals whose carcasses have been found to be fit for human consumption following post-mortem inspection.] (1) Ifish skins and bones described above are derived from plants that manufacture fishery products for human consumption which are authorised for export;] (¹) and, if of bovine, ovine and caprine animal origin, they have been derived from animals which passed ante-mortem and post-mortem inspections, ⁽¹⁾ and, except for hides and skins of ruminants, (¹) either [they come from a country or a region classified in accordance with Commission Decision 2007/453/EC of 29 June 2007 establishing the BSE status of Member States or third countries or regions thereof according to their BSE risk (OJ L 172, 30.6.2007, p. 84) as a country or region posing a negligible BSE risk; they do not contain and are not derived from 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 of 22 May 2001 laying down rules for the prevention, control and eradication of certain transmissible spongiform encephalopathies (OJ L 147, 31.5.2001, p. 1) (⁶); they do not contain and are not derived from mechanically separated meat obtained from bones of bovine, ovine or caprine animals, except for raw materials derived from animals that were born, continuously reared and slaughtered in a country or region classified in accordance with Decision 2007/453/EC as a country or region

the animals, from which the raw materials are derived, were not slaughtered after stunning by means of gas injected into the cranial cavity or killed by the same method or slaughtered, after stunning, by laceration of central nervous tissue by means of an elongated rod-shaped instrument introduced into the cranial cavity, except if the animals were born, continuously reared and slaughtered in a country or region classified in accordance with Decision 2007/453/EC as a country or region posing a negligible BSE risk;

posing a negligible BSE risk in which there has been no indigenous BSE cases;

Part II: Certification

and/or

and/or

Model RCG Raw materials for the production of collagen and gelatine intended for human consumption

COUN	TRY			w materials for the production of co	human consumption
II.	Hea	Ith information		II.a. Certificate reference No	II.b.
	—	accordance with D	ecision 2007/453/EC as meat-and-bone meal or	aterials are derived, originate from a country or region posing an undeto greaves, as defined in the Terrestria	ermined BSE risk, and the animal
	_	accordance with I materials were pr	Decision 2007/453/EC a oduced and handled ir	aterials are derived, originate from s a country or region posing an uno n a manner which ensures that the issues exposed during the deboning p	determined BSE risk, and the ray ey did not contain and were no
	(¹) 0	r			
	_	2007 establishing	he BSE status of Membe	sified in accordance with Commissior er States or third countries or regions r region posing a controlled BSE risk;	thereof according to their BSE ris
	_	derived, were not	killed, after stunning, b	ls of bovine, ovine and caprine anir y laceration of central nervous tissu ial cavity, or by means of gas injected	e by means of an elongated roc
	_	material as define		rine animal origin do not contain and V to Regulation (EC) No 999/2001, r caprine animals;]	
	(¹) 0	r			
	_	2007 establishing	he BSE status of Member	sified in accordance with Commissior er States or third countries or regions r region with an undetermined BSE ri	thereof according to their BSE ris
	—			are derived, were not fed meat-and-k mal Health Code of the World Organis	
	—	after stunning, by I	aceration of central nerv	of bovine, ovine and caprine animal rous tissue by means of an elongated njected into the cranial cavity;	
	_	the raw materials a	are not derived from:		
		(i) specified risk	material as defined in po	pint 1 of Annex V to Regulation (EC) I	No 999/2001;
		(ii) nervous and	ymphatic tissues expose	ed during the deboning process;	
		(iii) mechanically	separated meat obtaine	d from the bones of bovine, ovine or o	caprine animals.]
II.2.		Animal Health Att	estation (¹)		
				ify that the raw materials described at	Dove:
II.2.1.		consist of animal p	roducts that satisfy the a	nimal health requirements below;	
II.2.2.		have been obtaine	-	egion(s) thereof of (¹) either [:] (¹) c
(¹) eith	ner	[II.2.2.1 anima		ngs and have remained in that territor	ry since birth or for at least the las
		1 iii c ii c s	2 March 2010 laying do htroduction into the Eur ertification requirements mport requirements laid onsumption on a date	pecies referred to in Commission wn lists of third countries, territories ropean Union of certain animals ar s (OJ L 73, 20.3.2010, p. 1), satisfy d down in that Regulation, and th for which import into the Union of from the country or territory thereof Regulation:]	or parts thereof authorised for th nd fresh meat and the veterinar ing all the relevant animal healt nat were slaughtered for huma fresh meat from animals of thos

Model RCG Raw materials for the production of collagen and gelatine intended for

slaughter poultry from (a) third country(ies) listed for that commodity in Part 1 of Annex Commission Regulation (EC) No 798/2008 of 8 August 2008 laying down a list of third counterritories, zones or compartments from which poultry and poultry products may be imported int transit through the Community and the veterinary certification requirements (OJ L 226, 23.8, p. 1), under conditions at least equivalent to those in that Regulation satisfying all the relevant a health import requirements laid down in that Regulation and were slaughtered for human consur-	COUNTI	RY	R	aw materials for the production of	collagen and gelatine intended for human consumptior					
 down a list of third countries or parts thereof, for imports into, or transit through, the Common of meet of will leporidae, of certain wild land memals and of farmed rabbits and the vete certification requirements laid down in that Regulation.]] (1) or (1) or (1) I.2.2.1 poultry that have remained in that territory since hatching or have been imported as day-old bit slaupter poultry from (a) third country(se). Itseed for that commodity in Part. 1 of Annex Commission Regulation (EC) No 788/2008 of 8 August 2008 laying down a list of third country territories, zones or compartments from which poultry and poultry products may be imported in transit through the Community and the veterinary certification requirements (J.U. L226, 23.8, p. 1), under conditions at least equivalent to those in that Regulation requirements (J.U. L226, 23.8, p. 1), under conditions at least equivalent to those in that Regulation statistying all the relevant is health import requirements laid down in that Regulation requirements (J.U. L226, 23.8, p. 1), under conditions at least equivalent to those in maints at of thomas consum on a date for which import in the Union of meat from animals of thomas statistying all the relevant is the country or territory thereof in accordance with Column 6 B of Part 1 to Annex 1 to that Regulation (I) or [II.2.2.1 animals that have been killed in the wild in that territory (⁵) and captured and killed in an area: (i) in which within 25 km there has been no case/outbreak of any of the following diseases for the animals are susceptible: foot and mouth disease, inderpest, Newcaste disease or pathogenic avian influenza during the prior 30 days, nor of classical or African swine fever the prior 40 days, and (ii) in which after killing were transported within 12 hours for chilling either to a collection centr immediately afterwards to a game-handling establishment, or directly to a game-ha establishment.]] II.2.1. have been obtained in an establishm	II.	Health informat	tion	II.a. Certificate reference No	II.b.					
 [II.2.2.1 poultry that have remained in that territory since hatching or have been imported as day-old chi slaughter poultry from (a) third country(ies) listed for that commodity in Part 1 of Annee Commission Regulation (EC) No 798/2008 of 8 August 208 laying down a list of third counterritories, zones or compartments from which poultry and poultry products may be imported in transit through the Community and the veterinary certification requirements (G) L226, 23.8, p. 1), under conditions at least equivalent to those in that Regulation satisfying all the relevant a health linport requirements liad down in that Regulation and were slaughter do furman consur on a date for which import into the Union of meat from animals of those species was authorises the country or territory thereof in accordance with Column 6 B of Part 1 to Annex I to that Regula (!) or [U.2.2.1 animals that have been killed in the wild in that territory (^b) and captured and killed in an area: (i) in which within 25 km there has been no case/outbreak of any of the following diseases for the animals are susceptible foot and mouth disease; rinderpest, Newcastle disease or pathogenic avia influenza during the prior 30 days, nor of classical or African swine fever the prior 40 days, and (ii) in which after killing were transported within 12 hours for chilling either to a collection centr immediately afterwards to a game-handling establishment, or directly to a game-ha establishment] II.2.3. have been obtained in an establishment around which, within a radius of 10 km, there has been no case/out of the following diseases the relevant in fluenza and classical or African swine fever during the prior 30 days or, event of a case of one of those diseases, the preparation of raw materials for export to the Union has authorised on and classical or African swine fever during there than once and an influenza and classical or African swine fever		(¹) or	down a list of third cour of meat of wild leporida certification requiremen	ntries or parts thereof, for imports into e, of certain wild land mammals and its (OJ L 39, 10.2.2009, p. 12), satis	o, or transit through, the Community of farmed rabbits and the veterinary					
 slaughter poultry from (a) third country(ies) listed for that commodity in Part 1 of Anney Commission Regulation (EC) No 798/2008 of 8 August 208 laying down a list of third counterritories, zones or compartments from which poultry and poultry products may be imported in transit through the Community and the veterinary certification requirements (i) L226, 23.8, p. 1), under conditions at least equivalent to those in that Regulation satisfying all the relevant health Import requirements liad down in that Regulation and were sizuphered for human consur on a date for which import into the Union of meat from animals of those species was authorised the country or territory thereof in accordance with Column 6 B of Part 1 to Annex I to that Regulation and were sizuphere and killed in an area: (i) in which within 25 km there has been no case/outbreak of any of the following diseases for the animals are susceptible: foot and mouth disease, rinderpest, Newastie disease or pathogenic avian influenza during the prior 30 days, nor of classical or African swine fever of the prior 40 days, and (ii) that is situated at a distance that exceeds 20 km from the borders separating another term in a country or part thereof, which is not authorised on these dates to export these raw may into the Union, and (iii) in which after killing were transported within 12 hours for chilling either to a collection cent mimediately afterwards to a game-han establishment.] II.2.3. have been obtained in an establishment around which, within a radius of 10 km, there has been no case/out of the following diseases that the animals are susceptible to: foot and mouth disease, inderpest. New disease or highly pathogenic avian influenza, the preparation during the prior 30 days or, event of a case of one of those diseases. In the preparation and the total cleaning and disinfection of the establishment and the total cleaning and disinfection of the establishment turn of the following diseases	(1) or									
 [II.2.2.1 animals that have been killed in the wild in that territory (⁵) and captured and killed in an area: (i) in which within 25 km there has been no case/outbreak of any of the following diseases for the animals are susceptible: foot and mouth disease, inderpest, Newcastle disease or pathogenic avian influenza during the prior 30 days, nor of classical or African swine fever of the prior 40 days, and (ii) that is situated at a distance that exceeds 20 km from the borders separating another territ a country or part thereof, which is not authorised on these dates to export these raw ma into the Union, and (iii) in which after killing were transported within 12 hours for chilling either to a collection centr immediately afterwards to a game-handling establishment, or directly to a game-ha establishment.] II.2.3. have been obtained in an establishment around which, within a radius of 10 km, there has been no case/out of the following diseases that the animals are susceptible to: foot and mouth disease, iniderpest, New disease of night pathogenic avian influenza, and classical or African swine fever during the prior 30 days or, event of a case of one of those diseases, the preparation of raw materials for export to the Union has authorised only after the removal of all meat and the total cleaning and disinfection of the establishment und control of an Oficial veterinarian; II.2.4. have been obtained and prepared without contact with other materials that do not comply with the control required above, and have been handled so as to avoid contamination with pathogenic agents; and II.2.5. have been transported in clean and sealed containers or lorries. Notes See notes in Annex II of Commission Implementing Regulation (EC) No 2074/2005 and Implementing Regu (EU) 2019/528 of 8 April 2019 concerning model o certificates for certain animals and go		[II.2.2.1	slaughter poultry from (a) third country(ies) listed for that commodity in Part 1 of Annex I to Commission Regulation (EC) No 798/2008 of 8 August 2008 laying down a list of third countries, territories, zones or compartments from which poultry and poultry products may be imported into and transit through the Community and the veterinary certification requirements (OJ L 226, 23.8.2008, p. 1), under conditions at least equivalent to those in that Regulation satisfying all the relevant animal health import requirements laid down in that Regulation and were slaughtered for human consumption on a date for which import into the Union of meat from animals of those species was authorised from							
 (i) in which within 25 km there has been no case/outbreak of any of the following diseases for the animals are susceptible: foot and mouth disease, ininderpest, Newcastle disease or pathogenic avian influenza during the prior 30 days, nor of classical or African swine fever of the prior 40 days, and (ii) that is situated at a distance that exceeds 20 km from the borders separating another territi a country or part thereof, which is not authorised on these dates to export these raw main to the Union, and (iii) in which after killing were transported within 12 hours for chilling either to a collection centr immediately afterwards to a game-handling establishment, or directly to a game-ha establishment.] II.2.3. have been obtained in an establishment around which, within a radius of 10 km, there has been no case/out of the following diseases that the animals are susceptible to: foot and mouth disease, ininderpest, New disease or highly pathogenic avian influenza, and classical or African swine fever during the prior 30 days or, event of a case of one of those diseases, the preparation of raw materials for export to the Union has authorised only after the removal of all meat and the total cleaning and disinfection of the establishment und control of an official veterinarian; II.2.4. have been transported in clean and sealed containers or lorries. Notes See notes in Annex II of Commission Implementing Regulation (EU) 2019/628 of 8 April 2019 concerning model o certificates for certain animals and goods and amending Regulation (EC) No 2074/2005 and Implementing Regu (EU) 2016/759 as regards these model certificates (OJ L 131, 17.5.2019, p. 101) Part I: Box reference 1.8: provide the code of territory as appearing in Part 1 of Annex I to Regulation (EC) No 798/2008 are Part 1 of Annex I to Regulation (EC) No 119/2009 and Part 1 of Annex I to Regulation (EC) No 199/2008 and Part 1 of Anne	(1) or									
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Manufacturing plant: includes slaughterhouse, factory vessel, cutting plant, game-ha					206, 0207, 0208, 0302, 0303, 0305					
	— Во	ox reference I.25:	Nature of commodity:	hides, skins, bones, tendons and sine	ews;					
			u ,		essel, cutting plant, game-handling					

сои	NTRY	w materials for the production of c	Model RCG ollagen and gelatine intended for human consumption
۱۱.	Health information	II.a. Certificate reference No	II.b.
Part	11:		
(1)	Delete as appropriate. In the case of products deriv	ved from fishery products, the whole s	section II.2 should be deleted.
(²)	The name and ISO code number of the exporting of	country or territory or zone as laid dow	<i>i</i> n in:
	 the Annex II of Commission Delegated Regu 2017/625 of the European Parliament and of consignments of certain animals and goods int 	the Council with regard to requirement	ents for the entry into the Union of
	— Annex I to Regulation (EC) No 798/2008;		
	— Part 1 of Annex I to Regulation (EC) No 119/20	009;	
	- Part 1 of Annex II to Regulation (EC) No 206/2	010.	
(3)	If parts of the materials were derived from anin Regulation (EU) No 206/2010 for import of that cor of the third country slaughtering the animals shall supplementary guarantees A or F as indicated in co	nmodity into the EU, then the code(s) be stated (the material cannot come) of country(ies) or territory(ies) and
(4)	If the meat comes from slaughter poultry origin Regulation (EC) No 798/2008 for imports of that of and of the third country slaughtering the poultry sha	commodity into the EU, then the coc	
(5)	Only for countries from where game meat intende importation into the Union.	d for human consumption of the san	ne animal species is authorised for
(⁶)	The removal of specified risk material is not requi reared and slaughtered in a third country or region posing a negligible BSE risk.		
_	The signature and the stamp must be in a different	colour to that of the printing.	
NB	Note for the person responsible for the consignment accompany the consignment until it reaches the b manufacturing plant of destination.		
Offic	ial veterinarian		
	Name (in capital letters):	G	qualification and title:
	Date:	S	ignature:
	Stamp:		

PART IX

MODEL OFFICIAL CERTIFICATE FOR THE ENTRY INTO THE UNION FOR PLACING ON THE MARKET OF TREATED RAW MATERIALS FOR THE PRODUCTION OF GELATINE AND COLLAGEN INTENDED FOR HUMAN CONSUMPTION

cou	COUNTRY										Official certificate to the EU
	l.1.	Consignor/Exporter						I.2.	Certificate referen	nce No	I.2.a IMSOC reference No
		Name						1.3.	Central Competer	nt Author	ity
		Address						1.4.	Local Competent	Authority	1
ment		Tel. No									
sign	I.5.	Consignee/Importer						1.6.	Operator respons	ible for th	ne consignment
ned con		Name							Name		
patch		Address							Address		
f dis		Postal code							Postal code		
ils o		Tel. No									
Part I: Details of dispatched consignment	1.7.	Country of origin	ISO	I.8. Re	gion	of origin	Code	1.9.	Country of destination	ISO	I.10.
Ра	l.11	Place of dispatch						I.12.	Place of destination	on	
		Name		Approva	ıl No				Name		
		Address							Address		
	l.13.	Place of loading		1				I.14.	.14. Date and time of departure		
	l.15.	Means of transport						I.16.	6. Entry BCP		
		Aeroplane	Ves	ssel [Other		I.17.	Accompanying do	ocuments	
		Road vehicle	Rai	lway [Туре		
		Identification:							No		
	l.18.	Transport conditions									
		Ambient 🗌	Chi	lled [Frozen					
	l.19.	Container No/Seal N	0			1		I			

COUNTRY			Official certificate to the EU
I.20. Goods certified as			
Human consumption			
I.21.		1.22.	
I.23. Total number of packages	I.24. Quantity		
	Total number	Total net weight (Kg)	Total gross weight (Kg)
I.25. Description of goods			
No Code and CN	title		
Species (Scientific name)	Na	ture of commodity	Cold store
	Ma	anufacturing plant	
Number package	5	Batch No	Type of packaging

C	OUNTF	۲Y			Model TCC oduction of gelatine and collage intended for human consumption
11.	•	Health	n information	II.a. Certificate reference No	II.b.
11.	.1.	Public	c health attestation		
		l, the ı	undersigned, certify that the treated raw m	aterials described above comply with	the following requirements:
		_	they have been derived from establishme	ents under the control of and listed by	the competent authority,
		and			
		_	(¹) [bones, hides and skins of domestic derived from animals which were slaugh fit for human consumption following ante-	tered in a slaughterhouse and the ca	
		(¹) and	d/or		
		_	[wild game hides, skins and bones descr fit for human consumption following post-		whose carcasses were found to be
		(¹) and	d/or		
		_	[fish skins and bones described above consumption which are authorised for ex	•	facture fishery products for huma
		and			
		(¹) eith	ner		
_		—	[they are dried bones of species from bo wild animals, poultry including ratites an are derived from healthy animals slaught	d feathered game for the production	of gelatine and collagen, and the
		(¹) eith	ner		
		_	[crushed to pieces of approximately 15 n at least 30 minutes, a minimum of 80 °C then separated and subsequently washe minimum temperature of 350 °C, or for 700 °C,]	C for at least 15 minutes, or a minimu ed and dried for at least 20 minutes	um of 90 °C for at least 10 minutes in a stream of hot air with an initia
		(¹) or	[sun-dried for a minimum of 42 days at a	n average temperature of at least 20	°C,]
		(1) or	[have undergone an acid treatment such before drying,]	that the pH is maintained at less tha	n 6 to the core for at least one hou
		(1) or	[if they are hides and skins of farmed ru they are derived from healthy animals an		kins or wild game hides and skins
		(¹) eith	ner		
		—	[have undergone an alkali treatment wh days,]	ich ensures a PH> 12 to the core fo	llowed by salting for at least seve
		(1) or	[were dried for at least 42 days at a temp	perature of at least 20 °C,]	
		(1) or	[have undergone an acid treatment that hour,]	provides at least a pH of less than	5 to the core for a minimum of one
		(1) or	[have undergone an alkali treatment whic	ch ensures a pH > 12 to the core for a	at least 8 hours,]]
		(¹) or	[if they are bones, hides or skins of farm hides and skins from third countries, p Commission Implementing Regulation (regions thereof authorised for the entry human consumption, amending Implem 17.5.2019, p. 31), that they have under from establishments registered or appro- with Regulation (EC) No 853/2004,	parts of third countries or regions t (EU) 2019/626 of 5 March 2019 co y into the European Union of certai nenting Regulation (EU) 2016/759 gone any other treatment than those	thereof referred to in Article 15 to oncerning lists of third countries of in animals and goods intended fo as regards these lists (OJ L 131 e listed above, and that they como

coui	NTRY			Model TCG oduction of gelatine and collagen intended for human consumption
П.	Healt	h information	II.a. Certificate reference No	II.b.
	(¹) an	d, if of bovine, ovine and caprine animal or	igin,	
	_	they are derived from animals which pas	sed ante-mortem and post-mortem in	spections,
	(¹) an	d, except for hides and skins of ruminants,		
	(¹) eitl	her		
	_	[they come from a country or a regior 29 June 2007 establishing the BSE stat their BSE risk (OJ L 172, 30.6.2007, p. 8	us of Member States or third count	ries or regions thereof according to
	_	they do not contain and are not derived f (EC) No 999/2001 of the European Pa prevention, control and eradication of ce p. 1) (⁴),	rliament and of the Council of 22 M	lay 2001 laying down rules for the
	_	they do not contain and are not derived ovine or caprine animals, except for tre reared and slaughtered in a country or re region posing a negligible BSE risk in wh	eated raw materials derived from ar gion classified in accordance with De	nimals that were born, continuously ecision 2007/453/EC as a country or
	_	the animals, from which the treated raw gas injected into the cranial cavity or kil central nervous tissue by means of an el if the animals were born, continuously re Decision 2007/453/EC as a country or re	led by the same method or slaughte ongated rod-shaped instrument intro ared and slaughtered in a country or	ered, after stunning, by laceration of duced into the cranial cavity, except
	_	(¹) [the animals, from which the treated r accordance with Decision 2007/453/EC not been fed with meat-and-bone meal c Organisation for Animal Health];	as a country or region posing an unc	determined BSE risk, and they have
	_	(¹) the animals, from which the treated r accordance with Decision 2007/453/EC products were produced and handled contaminated with nervous and lymphatic	c as a country or region posing ar in a manner which ensures that t	n undetermined BSE risk, and the hey did not contain and were not
	(¹) or			
	_	[they come from a country or a region 29 June 2007 establishing the BSE stat their BSE risk (OJ L 172, 30.6.2007, p. 8	us of Member States or third count	ries or regions thereof according to
	_	the animals, from which the treated raw are derived, were not killed, after stunnir shaped instrument introduced into the cra	ig, by laceration of central nervous ti	ssue by means of an elongated rod-
	—	the treated raw materials of bovine, ovi specified risk material as defined in p separated meat obtained from bones of b	oint 1 of Annex V to Regulation (I	
	(¹) or			
	_	[they come from a country or a region 29 June 2007 establishing the BSE stat their BSE risk (OJ L 172, 30.6.2007, p. 8	us of Member States or third count	ries or regions thereof according to
	_	the animals from which the treated ray greaves derived from ruminants, as defi Animal Health,		

Model TCG Treated raw materials for the production of gelatine and collagen

II. Health information II.a. Certificate reference — the animals, from which the treated raw materials of bovine, ovin killed, after stunning, by laceration of central nervous tissue box	
	ne and caprine animal origin are derived were not
introduced into the cranial cavity, or by means of gas injected in	by means of an elongated rod-shaped instrument
 the treated raw materials are not derived from: 	
(i) specified risk material as defined in point 1 of Annex V of F	Regulation (EC) No 999/2001;
(ii) nervous and lymphatic tissues exposed during the debonir	ng process,
(iii) mechanically separated meat obtained from bones of bovi	ne, ovine or caprine animals.]]
II.2. Animal Health Attestation (1)	
I, the undersigned official veterinarian, certify that the treated raw mate	rials described above:
II.2.1. consist of animal products that satisfy the animal health requirements b	pelow,
II.2.2. have been obtained in the country(ies) or region(s) thereof of $(^1)$ [:] $(^2)(^3)$,] (¹) or
II.2.3. have been obtained and prepared without contact with other material above, and have been handled so as to avoid contamination with patho	
II.2.4. have been transported in clean and sealed containers or lorries.	
See notes in Annex II of Commission Implementing Regulation (EU) 2019 certificates for certain animals and goods and amending Regulation (EC (EU) 2016/759 as regards these model certificates (OJ L 131, 17.5.2019, p.	C) No 2074/2005 and Implementing Regulation
Part I:	
 Box reference I.8: Provide the code of the territory as it appears in: 	
 in Part 1 of Annex I to Commission Regulation (EC) No 798/2008 of 8 territories, zones or compartments from which poultry and poultry p the Community and the veterinary certification requirements (OJ L 22 	roducts may be imported into and transit through
 in Part 1 of Annex I to Commission Regulation (EC) No 119/2009 la for imports into, or transit through, the Community of meat of wild farmed rabbits and the veterinary certification requirements (OJ L 39, 	leporidae, of certain wild land mammals and of
 in Part 1 of Annex II to Commission Regulation (EU) No 206/20 countries, territories or parts thereof authorised for the introduction fresh meat and the veterinary certification requirements (OJ L 73, 20. 	into the European Union of certain animals and
 Box reference I.25: Insert the appropriate Harmonised System (HS) cod 0511.99, 1602, 1604, 4101, 4102 or 4103. 	de(s) such as: 0210, 0305, 0505, 0506, 0511 91,
— Box reference I.25: Nature of commodity: hides, skins, bones, to	endons and sinews;
······································	
	rhouse, factory vessel, cutting plant, game nt.

col	JNTRY	Treated raw materials for the	Model TCG production of gelatine and collagen intended for human consumption			
	Health information	II.a. Certificate reference No	II.b.			
Par	t II:					
(¹)	Delete as appropriate. In the case of products deri	ved from fishery products, the whole	e section II.2 should be deleted.			
(²)	The name and ISO code number of the exporting of	country or territory or zone as laid do	own in:			
	— Part 1 of Annex II to Regulation (EC) No 206/2	2010;				
	— Annex I to Regulation (EC) No 798/2008;					
	— Part 1 of Annex I to Regulation (EC) No 119/2	009.				
(3)) If parts of the materials were derived from animals originating from an(other) third country(ies) or regions thereof listed Article 15 or 16 (only when treated as laid down in Part II.1) to Implementing Regulation (EU) 2019/626, the code(s) or country(ies) or region(s) shall be stated.					
(4)) The removal of specified risk material is not required if the treated raw materials are derived from animals born continuously reared and slaughtered in a third country or region of a third country classified in accordance with Decisior 2007/453/EC as posing a negligible BSE risk.					
_	The signature and the stamp must be in a different	t colour to that of the printing.				
NB	B Note for the person responsible for the consignment in EU: this certificate is only for veterinary purposes and has to accompany the consignment until it reaches the border control post. The consignment must be transported directly to the manufacturing plant of destination.					
-	The time of transportation may be included in the o	duration of treatment.				
Offi	cial veterinarian					
	Name (in capital letters):		Qualification and title:			
	Date:		Signature:			
	Stamp:					

PART X

MODEL OFFICIAL CERTIFICATE FOR THE ENTRY INTO THE UNION FOR PLACING ON THE MARKET OF HONEY AND OTHER APICULTURE PRODUCTS INTENDED FOR HUMAN CONSUMPTION

cou	COUNTRY							Official certificate to the EU			
	l.1.							I.2.	Certificate referen	ce No	I.2.a IMSOC reference No
		Name						1.3.	Central Competer	nt Author	ity
		Address						1.4.	Local Competent	Authority	,
ment		Tel. No									
sign	1.5.	Consignee/Importer						1.6.	Operator responsi	ible for th	ne consignment
ed con		Name							Name		
patche		Address							Address		
of dis		Postal code							Postal code		
ails c		Tel. No									
Part I: Details of dispatched consignment	1.7.	Country of origin	ISO	1.8.				1.9.	Country of destination	ISO	I.10.
P	l.11	Place of dispatch						I.12.	Place of destination	on	
		Name		Approv	val No				Name		
		Address							Address		
	I.13.	Place of loading						l.14.	Date and time of o	departure	3
	l.15.	Means of transport						l.16.	Entry BCP		
		Aeroplane	Ves	ssel		Other		l.17.	Accompanying do	cuments	;
		Road vehicle	Rai	lway					-		
									Type No		
		Identification:									
	l.18.	Transport conditions									
		Ambient 🛛	Chi	lled		Frozen					
	l.19.	Container No/Seal No)			1		1			

COUNTRY			Official certificate to the EU
I.20. Goods certified as			
Human consumption			
I.21.		1.22.	
I.23. Total number of packages	I.24. Quantity		
	Total number	Total net weight (Kg)	Total gross weight (Kg)
I.25. Description of goods			
No Code and CN	title		
Species (Scientific name)	Ma	nufacturing plant	Treatment type Cold store
Final consumer Number of package	5	Batch No	Type of packaging

	ſŔŶ	Honey and other apiculture products	Model HON s intended for human consumption
II.	Health information	II.a. Certificate reference No	II.b.
II.1.	Public health attestation		
	I, the undersigned, declare that I am award Parliament and of the Council of 28 Janu establishing the European Food Safety A 1.2.2002, p. 1), Regulation (EC) No 852/2 hygiene of foodstuffs (OJ L 139, 30.4.2004 the Council of 29 April 2004 laying down sp Regulation (EU) 2017/625 of the European official activities performed to ensure the health and plant protection products, amer (EC) No 1107/2009, (EU) No 1151/2012, Parliament and of the Council, Council R 98/58/EC, 1999/74/EC, 2007/43/EC, 2008/ (EC) No 882/2004 of the European Par 90/425/EEC, 91/496/EEC, 96/23/EC, 96/2 Regulation) (OJ L 95, 7.4.2017, p. 1), and	ary 2002 laying down the general princ Authority and laying down procedures 004 of the European Parliament and of 4, p. 1) and Regulation (EC) No 853/200 becific hygiene rules for food of animal or Parliament and of the Council of 15 Mar application of food and feed law, rules iding Regulations (EC) No 999/2001, (EC (EU) No 652/2014, (EU) 2016/429 ar egulations (EC) No 1/2005 and (EC) N (119/EC and 2008/120/EC and repealing liament and of the Council, Council	siples and requirements of food law in matters of food safety (OJ L 31 the Council of 29 April 2004 on the 4 of the European Parliament and o rigin (OJ L 139, 30.4.2004, p. 55) and ch 2017 on official controls and othe on animal health and welfare, plan C) No 396/2005, (EC) No 1069/2009 ad (EU) 2016/2031 of the Europear o 1099/2009 and Council Directives g Regulations (EC) No 854/2004 and Directives 89/608/EEC, 89/662/EEC
	I certify that honey and other apicultur requirements, in particular that they:	re products described above were p	roduced in accordance with these
		plementing a programme based on the nce with Article 5 of Regulation (EC) No	
	 have been handled and, where appr with the requirements of Annex II to F 	opriate, prepared, packaged and stored Regulation (EC) No 852/2004; and	in a hygienic manner in accordance
	accordance with Council Directive 9 residues thereof in live animals and	animals and products thereof provided 6/23/EC of 29 April 1996 on measures animal products and repealing Directive EC (OJ L 125, 23.5.1996, p. 10), and in p	s to monitor certain substances and es 85/358/EEC and 86/469/EEC and
Notes			
certifi	otes in Annex II of Commission Implemer cates for certain animals and goods and 016/759 as regards these model certificat	amending Regulation (EC) No 2074/2	
certifi	cates for certain animals and goods and 016/759 as regards these model certificat	amending Regulation (EC) No 2074/2	
certifi (EU) 2 Part I:	cates for certain animals and goods and 016/759 as regards these model certificat	amending Regulation (EC) No 2074/2 es (OJ L 131, 17.5.2019, p. 101).	
certifi (EU) 2 Part I: — E — E	cates for certain animals and goods and 016/759 as regards these model certificat	amending Regulation (EC) No 2074/2 es (OJ L 131, 17.5.2019, p. 101). al number means registration number.	2005 and Implementing Regulatior
Certifi (EU) 2 Part I: E 1 E	cates for certain animals and goods and 016/759 as regards these model certificat Box reference I.11: place of dispatch: Approva	amending Regulation (EC) No 2074/2 es (OJ L 131, 17.5.2019, p. 101). al number means registration number. armonised System (HS) code(s) using h	2005 and Implementing Regulation eadings such as: 0409, 0410, 0510
Certifi (EU) 2 Part I: E 1 E	Cates for certain animals and goods and 016/759 as regards these model certificat Box reference I.11: place of dispatch: Approva Box reference I.25: Insert the appropriate Ha 521, 1702 or 2106. Box reference I.25: <i>Treatment type</i> : state 'u reatment'.	amending Regulation (EC) No 2074/2 es (OJ L 131, 17.5.2019, p. 101). al number means registration number. armonised System (HS) code(s) using h	2005 and Implementing Regulation eadings such as: 0409, 0410, 0510
Certifi (EU) 2 Part I: E 1 E tu Part II	Cates for certain animals and goods and 016/759 as regards these model certificat Box reference I.11: place of dispatch: Approva Box reference I.25: Insert the appropriate Ha 521, 1702 or 2106. Box reference I.25: <i>Treatment type</i> : state 'u reatment'.	amending Regulation (EC) No 2074/2 es (OJ L 131, 17.5.2019, p. 101). al number means registration number. armonised System (HS) code(s) using h ultrasonication', 'homogenisation', ultrafi	2005 and Implementing Regulation eadings such as: 0409, 0410, 0510 Itration', 'pasteurisation', 'no therma
Certifi (EU) 2 Part I: E 1 E tu Part II T	cates for certain animals and goods and 016/759 as regards these model certificat Box reference I.11: place of dispatch: Approva Box reference I.25: Insert the appropriate Ha 521, 1702 or 2106. Box reference I.25: <i>Treatment type</i> : state 'u reatment'.	amending Regulation (EC) No 2074/2 es (OJ L 131, 17.5.2019, p. 101). al number means registration number. armonised System (HS) code(s) using h ultrasonication', 'homogenisation', ultrafi	2005 and Implementing Regulation eadings such as: 0409, 0410, 0510 Itration', 'pasteurisation', 'no therma
Certifi (EU) 2 Part I: E 1 E tu Part II T Officia	Cates for certain animals and goods and 016/759 as regards these model certificat Box reference I.11: place of dispatch: Approva Box reference I.25: Insert the appropriate Ha 521, 1702 or 2106. Box reference I.25: <i>Treatment type</i> : state 'u reatment'.	amending Regulation (EC) No 2074/2 es (OJ L 131, 17.5.2019, p. 101). al number means registration number. armonised System (HS) code(s) using h ultrasonication', 'homogenisation', ultrafi	2005 and Implementing Regulation eadings such as: 0409, 0410, 0510 Itration', 'pasteurisation', 'no therma
Certifi (EU) 2 Part I: E 1 E tu Part II T Officia	Cates for certain animals and goods and 016/759 as regards these model certificat Box reference I.11: place of dispatch: Approva Box reference I.25: Insert the appropriate Ha 521, 1702 or 2106. Box reference I.25: <i>Treatment type</i> : state 'u reatment'.	amending Regulation (EC) No 2074/2 es (OJ L 131, 17.5.2019, p. 101). al number means registration number. armonised System (HS) code(s) using h ultrasonication', 'homogenisation', ultrafi	2005 and Implementing Regulation eadings such as: 0409, 0410, 0510 Itration', 'pasteurisation', 'no therma

PART XI

MODEL OFFICIAL CERTIFICATE FOR THE ENTRY INTO THE UNION FOR PLACING ON THE MARKET OF HIGHLY REFINED CHONDROITIN SULPHATE, HYALURONIC ACID, OTHER HYDOLYSED CARTILAGE PRODUCTS, CHITOSAN, GLUCOSAMINE, RENNET, ISINGLASS AND AMINO ACIDS INTENDED FOR HUMAN CONSUMPTION

cou	NTRY						Official certificate to the EU			
	l.1.	Consignor/Exporter					1.2.	Certificate referen	ce No	I.2.a IMSOC reference No
		Name					1.3.	Central Competer	nt Author	ity
		Address					I.4.	Local Competent	Authority	1
nent	Tel. No									
signn	1.5.	Consignee/Importer					I.6.	Operator responsi	ible for tl	ne consignment
d con		Name						Name		
Part I: Details of dispatched consignment		Address Postal code Tel. No						Address Postal code		
art I: Deta	1.7.	Country of origin	ISO	1.8.			1.9.	Country of destination	ISO	I.10.
Ра	l.11	Place of dispatch					I.12.	Place of destination	on	
		Name		Approval No				Name		
		Address						Address		
	I.13.	Place of loading					I.14. Date and time of departure			
	I.15.	Means of transport						I.16. Entry BCP		
		Aeroplane	Ves	ssel 🛛	Other		I.17.	Accompanying do	cuments	3
		Road vehicle	Rai	ilway 🛛			Туре			
		Identification:						No		
	I.18.	Transport conditions					-			
		Ambient	Chi	illed	Frozen					
	l.19.	Container No/Seal No)							

COUNTRY				Official certificate to the EU
I.20. Goods certified as				
Human consumption				
I.21.			1.22.	
I.23. Total number of packages	1.24. 0	Quantity		
	т	otal number	Total net weight (Kg)	Total gross weight (Kg)
I.25. Description of goods				
No Code and CN	title			
Species (Scientific name)		Ма	nufacturing plant	Cold store
Final consumer Number of package		Net weight	Batch No	Type of packaging

Model HRP

Highly refined chondroitin sulphate, hyaluronic acid, other hydrolysed cartilage products, chitosan, glucosamine, rennet, isinglass and amino acids intended for human consumption

	COUN	TRY	acids intended for human consumption								
	П.	Health information	II.a. Certificate reference No	II.b.							
	II.1.	Public health attestation									
Part II: Certification		Parliament and of the Council of 28 Janu establishing the European Food Safety / 1.2.2002, p. 1), Regulation (EC) No 852/2 hygiene of foodstuffs (OJ L 139, 30.4.2004 the Council of 29 April 2004 laying down sp Regulation (EU) 2017/625 of the European official activities performed to ensure the health and plant protection products, amer (EC) No 1107/2009, (EU) No 1151/2012, Parliament and of the Council, Council R 98/58/EC, 1999/74/EC, 2007/43/EC, 2008/ (EC) No 882/2004 of the European Par	are of the relevant provisions of Regulation (EC) No 178/2002 of the European nuary 2002 laying down the general principles and requirements of food law, / Authority and laying down procedures in matters of food safety (OJ L 31, //2004 of the European Parliament and of the Council of 29 April 2004 on the //04, p. 1) and Regulation (EC) No 853/2004 of the European Parliament and of specific hygiene rules for food of animal origin (OJ L 139, 30.4.2004, p. 55) and an Parliament and of the Council of 15 March 2017 on official controls and other le application of food and feed law, rules on animal health and welfare, plant ending Regulations (EC) No 999/2001, (EC) No 396/2005, (EC) No 1069/2009, I2, (EU) No 652/2014, (EU) 2016/429 and (EU) 2016/2031 of the European Regulations (EC) No 1/2005 and (EC) No 1099/2009 and Council Directives 20/119/EC and 2008/120/EC and repealing Regulations (EC) No 854/2004 and 'arliament and of the Council, Council Directives 89/608/EEC, 89/662/EEC, 6/93/EC and 97/78/EC and Council Decision 92/438/EEC (Official Controls d								
I certify that the highly refined products described above were produced in accordance with these requirem particular:											
		, , , , , , , , , , , , , , , , , , ,	ent(s) implementing a programme based accordance with Article 5 of Regulation (EC	2							
			vhere appropriate, prepared, packaged a Annex II to Regulation (EC) No 852/2004;	nd stored in a hygienic manner in							
	— that they comply with the requirements of Section XVI of Annex III to Regulation (EC) No 853/2004; and										
	— (¹) if amino acids, that										
	(i) human hair was not used as a source for their manufacture; and										
			on (EC) No 1333/2008 of the European ives ((OJ L 354, 31.12.2008, p. 16).	Parliament and of the Council of							
	Notes See notes in Annex II of Commission Implementing Regulation (EU) 2019/628 of 8 April 2019 concerning model official certificates for certain animals and goods and amending Regulation (EC) No 2074/2005 and Implementing Regulation (EU) 2016/759 as regards these model certificates (OJ L 131, 17.5.2019, p. 101).										
	Part I	:									
		Box reference I.25: Insert the appropriate Hai ex 2932, 3507 or 3503.	rmonised System (HS) code(s) using head	lings such as 2833, ex 3913, 2930,							
	Part I	1:									
	(¹) [Delete as appropriate.									
		 The colour of the stamp and signature must be different from that of the other particulars in the certificate. 									
	Officia	al veterinarian									
	1	Name (in capital letters):		Qualification and title:							
	[Date:		Signature:							
	9	Stamp:									

PART XII

MODEL OFFICIAL CERTIFICATE FOR THE ENTRY INTO THE UNION FOR PLACING ON THE MARKET OF REPTILE MEAT INTENDED FOR HUMAN CONSUMPTION

cou	NTRY							Official certificate to the EU			
	l.1.	Consignor/Exporter						1.2.	Certificate referen	ice No	I.2.a IMSOC reference No
		Name						1.3.	Central Competer	nt Author	ity
		Address						1.4.	Local Competent	Authority	,
ment	Tel. No I.5. Consignee/Importer										
sign								I.6.	1.6. Operator responsible for the consignment		
ned con		Name							Name		
patch		Address							Address		
f dis		Postal code							Postal code		
ils of		Tel. No									
Part I: Details of dispatched consignment	1.7.	Country of origin	ISO	1.8.			1.9.	Country of destination	ISO	I.10.	
Ра	l.11	Place of dispatch					I.12.	Place of destination	on		
		Name		Approval N	lo				Name		
		Address						Address			
	l.13.	Place of loading						l.14.	Date and time of o	departure	3
	l.15.	Means of transport						l.16.	Entry BCP		
		Aeroplane	Ves	sel 🛛		Other		l.17.	7. Accompanying documents		
		Road vehicle	Rail	lway 🛛					Туре		
	Identification:								No		
		Ambient	Chil	Chilled		Frozen					
	l.19.	Container No/Seal No						1			

COUNTRY				Official certificate to the EU
I.20. Goods certified as				
Human consumption				
I.21.			1.22.	
I.23. Total number of packages	1.24. (Quantity		
	r	otal number	Total net weight (Kg)	Total gross weight (Kg)
I.25. Description of goods				
No Code and	CN title			
Species (Scientific name)	Ма	nufacturing plant	Cold store
Final consumer Numb		Net weight	Batch No	Type of packaging

	СОЛИТ	RY		Reptile Meat i	ntended for human consumption					
	п.	Hea	Ith information	II.a. Certificate reference No	II.b.					
	II.1.	Pub	lic health attestation							
Part II: Certification		Parli esta 1.2.2 hygi the (Reg offic heal (EC) 98/5 (EC) 90/4	iament and of the Council of 28 January 2 blishing the European Food Safety Author 2002, p. 1), Regulation (EC) No 852/2004 ene of foodstuffs (OJ L 139, 30.4.2004, p. Council of 29 April 2004 laying down specifi- ulation (EU) 2017/625 of the European Parl ial activities performed to ensure the appli th and plant protection products, amending) No 1107/2009, (EU) No 1151/2012, (EU iament and of the Council, Council Regula 8/EC, 1999/74/EC, 2007/43/EC, 2008/119/) No 882/2004 of the European Parliame	the relevant provisions of Regulation (EC) No 178/2002 of the European 2002 laying down the general principles and requirements of food law, ority and laying down procedures in matters of food safety (OJ L 31, of the European Parliament and of the Council of 29 April 2004 on the 1) and Regulation (EC) No 853/2004 of the European Parliament and of fic hygiene rules for food of animal origin (OJ L 139, 30.4.2004, p. 55) and diament and of the Council of 15 March 2017 on official controls and other lication of food and feed law, rules on animal health and welfare, plant g Regulations (EC) No 999/2001, (EC) No 396/2005, (EC) No 1069/2009, U) No 652/2014, (EU) 2016/429 and (EU) 2016/2031 of the European lations (EC) No 1/2005 and (EC) No 1099/2009 and Council Directives V/EC and 2008/120/EC and repealing Regulations (EC) No 854/2004 and nent and of the Council, Council Directives 89/608/EEC, 89/662/EEC, EC and 97/78/EC and Council Decision 92/438/EEC (Official Controls						
		l cer	tify that the reptile meat described above wa	as produced in accordance with these	requirements, in particular:					
		—	that the reptile meat comes from (an) est and critical control points (HACCP) princip							
		—	that the reptile meat has been handled manner in accordance with the requirement							
	-	_	that <i>Salmonella</i> has been controlled in th equivalent guarantees as the requirement 15 November 2005 on microbiological crite	nts once laid down in Commission	Regulation (EC) No 2073/2005 of					
			that the reptile meat is obtained from an inspection laid down in Article 73 of Comm down uniform practical arrangements for t for human consumption in accordance w Council and amending Commission Rep 17.5.2019, p. 51);	nission Implementing Regulation (EU) he performance of official controls on vith Regulation (EU) 2017/625 of the	2019/627 of 15 March 2019 laying products of animal origin intended European Parliament and of the					
		_	(¹) if crocodile or alligator meat, that the c presence of <i>Trichinella</i> spp. in accorda 10 August 2015 laying down specific rule and	ance with Commission Implementin	g Regulation (EU) 2015/1375 of					
		_	that, when applicable, the food has been a (EU) 2015/2283 of the European Parliam Regulation (EU) No 1169/2011 of the Eu No 258/97 of the European Parliament (OJ L 327, 11.12.2015, p. 1) and listed in t	ent and of the Council of 25 Novemb uropean Parliament and of the Cour and of the Council and Commissio	er 2015 on novel foods, amending noil and repealing Regulation (EC)					
	Notes									
	certifie	cates	n Annex II of Commission Implementing for certain animals and goods and ame 59 as regards these model certificates (C	nding Regulation (EC) No 2074/20						
	Part I:									
	— в	lox ref	erence I.25: Insert the appropriate HS/CN c	ode(s) such as 0208 50 00, 0210 93 (00, 1506, 1601, 1602 or 1603.					

cou	NTRY	Reptile Meat intended for human consumption										
Н.	Health information	II.a. Certificate reference No	II.b.									
Part	Part II:											
(1)	Delete as appropriate.											
_	The colour of the stamp and signature must be different from that of the other particulars in the certificate.											
Offic	cial veterinarian											
	Name (in capital letters):		Qualification and title:									
	Date:		Signature:									
	Stamp:											

PART XIII

MODEL OFFICIAL CERTIFICATE FOR THE ENTRY INTO THE UNION FOR PLACING ON THE MARKET OF INSECTS INTENDED FOR HUMAN CONSUMPTION

cou	NTRY							Official certificate to the EU			
	l.1.	Consignor/Exporter						1.2.	Certificate referen	ce No	I.2.a IMSOC reference No
		Name						1.3.	Central Competer	nt Author	ity
		Address						1.4.	Local Competent	Authority	,
nent		Tel. No									
signr	I.5. Consignee/Importer							I.6.	Operator responsi	ible for th	ne consignment
ed con	Name								Name		
spatch		Address							Address		
of dis	Postal code							Postal code			
ails c		Tel. No									
Part I: Details of dispatched consignment	1.7.	Country of origin	ISO	1.8.			1.9.	Country of destination	ISO	I.10.	
P	l.11	Place of dispatch					I.12.	Place of destination	on		
		Name		Approval No				Name			
		Address						Address			
	l.13.	Place of loading						l.14.	I.14. Date and time of departure		
	l.15.	Means of transport						I.16.	Entry BCP		
		Aeroplane	Ves	sel		Other 🗖		I.17.	Accompanying documents		
		Road vehicle	Rai	lway				Туре			
	Identification:								No		
	Ambient Chilled Frozen										
	l.19.	Container No/Seal No						1			

COUNTRY				Official certificate to the EU
I.20. Goods certified as				
Human consumption				
l.21.			1.22.	
I.23. Total number of packages	1.24. G	Quantity		
	Т	otal number	Total net weight (Kg)	Total gross weight (Kg)
I.25. Description of goods				
No Code and CN	title			
Species (Scientific name)		Cutting pl	ant/manufacturing plant	Cold store
Final consumer Number package		Net weight	Batch No	Type of packaging

	COUN	COUNTRY		Model Insects i	intended for human consun	nption					
	11.	Hea	Ith information	II.a. Certificate reference No	II.b.						
	II.1.	Pub	lic health attestation								
Part II: Certification		Parli esta 1.2.2 hygi the (Reg offic heal (EC) 98/5 (EC) 90/4	e undersigned, declare that I am aware of the iament and of the Council of 28 January 2 blishing the European Food Safety Autho 2002, p. 1), Regulation (EC) No 852/2004 of ene of foodstuffs (OJ L 139, 30.4.2004, p. 1 Council of 29 April 2004 laying down specific ulation (EU) 2017/625 of the European Parli ial activities performed to ensure the applic th and plant protection products, amending) No 1107/2009, (EU) No 1151/2012, (EU iament and of the Council, Council Regula 8/EC, 1999/74/EC, 2007/43/EC, 2008/119/f) No 882/2004 of the European Parliame 25/EEC, 91/496/EEC, 96/23/EC, 96/93/EC ulation (OJ L 95, 7.4.2017, p. 1)), and	2002 laying down the general princip rity and laying down procedures in of the European Parliament and of th 1) and Regulation (EC) No 853/2004 c hygiene rules for food of animal orig ament and of the Council of 15 Marcl cation of food and feed law, rules o Regulations (EC) No 999/2001, (EC)) No 652/2014, (EU) 2016/429 and titions (EC) No 1/2005 and (EC) No EC and 2008/120/EC and repealing f ent and of the Council, Council Dir	ples and requirements of foo matters of food safety (OJ the Council of 29 April 2004 of the European Parliament gin (OJ L 139, 30.4.2004, p. 5 th 2017 on official controls and on animal health and welfare) No 396/2005, (EC) No 1069 d (EU) 2016/2031 of the Eur 1099/2009 and Council Dire Regulations (EC) No 854/200 rectives 89/608/EEC, 89/662	d law, L 31, on the and of 5) and d other , plant /2009, opean ectives 04 and 2/EEC,					
		l cer	tify that the insects described above were pr	roduced in accordance with these req	quirements, in particular:						
		—	that they come from (an) establishment(s) control points (HACCP) principles in accord			critical					
		_	that they have been handled and, where accordance with the requirements of Ann No 852/2004;								
	-	—	that they comply with the requirements No 853/2004, including as regards the use		of Annex III to Regulation	(EC)					
		when applicable, the food has been authorised on the Union market in accordance with Article 6 of Regulatic (EU) 2015/2283 of the European Parliament and of the Council of 25 November 2015 on novel foods, amendin Regulation (EU) No 1169/2011 of the European Parliament and of the Council and repealing Regulation (EC) No 258/97 of the European Parliament and of the Council and Commission Regulation (EC) No 1852/200 (OJ L 327, 11.12.2015, p. 1) and listed in Commission Implementing Regulation (EU) 2017/2470 of 20 December 2017 establishing the Union list of novel foods in accordance with Regulation (EU) 2015/2283 of the European Parliament and of UJ L 351, 30.12.2017, p. 72).									
	Notes See notes in Annex II of Commission Implementing Regulation (EU) 2019/628 of 8 April 2019 concerning model officia certificates for certain animals and goods and amending Regulation (EC) No 2074/2005 and Implementing Regulatio (EU) 2016/759 as regards these model certificates (OJ L 131, 17.5.2019, p. 101).										
	Part I	l:									
	-	Box ref	erence I.25: Insert the appropriate HS/CN c	ode(s) such as 0106 49 00, 0410 or 2	2106.						
	Part I	11:									
	(1)) Delete as appropriate									
		 Box II.1 a programme based on the HACCP principles is not required if the products come directly from a primary producer. 									
	The colour of the stamp and signature must be different from that of the other particulars in the certificate.										
	Officia	al veter	inarian								
		Name	(in capital letters):		Qualification and title:						
		Date:			Signature:						
	· ·	Stamp:									

PART XIV

MODEL OFFICIAL CERTIFICATE FOR THE ENTRY INTO THE UNION FOR PLACING ON THE MARKET OF OTHER PRODUCTS OF ANIMAL ORIGIN INTENDED FOR HUMAN CONSUMPTION NOT COVERED BY ARTICLES 7 TO 25 OF COMMISSION IMPLEMENTING REGULATION (EU) 2019/628

cou	NTRY						Official certificate to the EU			
	l.1.						1.2.	Certificate referen	ice No	I.2.a IMSOC reference No
		Name					1.3.	I.3. Central Competent Authority		
		Address					I.4.	Local Competent	Authority	,
nent		Tel. No								
signr	1.5.	Consignee/Importer					I.6.	Operator respons	ible for th	ne consignment
hed cons		Name						Name		
patcl		Address						Address		
f dis		Postal code						Postal code		
ils o		Tel. No								
Part I: Details of dispatched consignment	1.7.	Country of origin	ISO	1.8.		1.9.	Country of destination	ISO	I.10.	
Ра	l.11	Place of dispatch					I.12.	Place of destination	on	
		Name		Approval No				Name		
		Address						Address		
	l.13.	Place of loading					l.14.	Date and time of o	departure	3
	l.15.	Means of transport					l.16.	.16. Entry BCP		
		Aeroplane	Ves	ssel 🛛	Other		I.17.	Accompanying do	ocuments	
		Road vehicle	Rai	lway 🛛						
								Туре		
		Identification:						No		
	l.18.	Transport conditions								
		Ambient	Chi	lled 🛛	Frozen					
	l.19.	Container No/Seal No				1				

COUNTRY				Official certificate to the EU	
I.20. Goods certified as					
Human consumption					
I.21.			1.22.		
I.23. Total number of packages	1.24. Q	uantity			
	Total number		Total net weight (Kg)	Total gross weight (Kg)	
I.25. Description of goods					
No Code and CN	title				
Species (Scientific name)		Ма	Manufacturing plant		
Final consumer Number of package		Net weight	Batch No	Type of packaging	

 control points (HACCP) principles in that they have been handled and, accordance with the requirements of 	Authority and laying down the general prin Authority and laying down procedures (2004 of the European Parliament and co (04, p. 1) and Regulation (EC) No 853/20 (specific hygiene rules for food of animal of a Parliament and of the Council of 15 Ma (e application of food and feed law, rules (ending Regulations (EC) No 999/2001, (E (2, (EU) No 652/2014, (EU) 2016/429 a (Regulations (EC) No 1/2005 and (EC) (EC) No 1/2005 and (EC) (EC) No 1/2005 and (EC) (EC) and 2008/120/EC and repealir (E) (C) (C) (C) (C) (C) (C) (C) (C) (C) (C	nciples and requirements of food law, is in matters of food safety (OJ L 31 of the Council of 29 April 2004 on the 004 of the European Parliament and or origin (OJ L 139, 30.4.2004, p. 55) and arch 2017 on official controls and other is on animal health and welfare, plan EC) No 396/2005, (EC) No 1069/2009 and (EU) 2016/2031 of the Europear No 1099/2009 and Council Directives ing Regulations (EC) No 854/2004 and Directives 89/608/EEC, 89/662/EEC ecision 92/438/EEC (Official Controls e requirements, in particular: wed on the hazard analysis and critica (EC) No 852/2004; d and stored in a hygienic manner in
I, the undersigned, declare that I am awar Parliament and of the Council of 28 Jam establishing the European Food Safety 1.2.2002, p. 1), Regulation (EC) No 852// hygiene of foodstuffs (OJ L 139, 30.4.200 the Council of 29 April 2004 laying down s Regulation (EU) 2017/625 of the European official activities performed to ensure the health and plant protection products, ame (EC) No 1107/2009, (EU) No 1151/2012 Parliament and of the Council, Council F 98/58/EC, 1999/74/EC, 2007/43/EC, 2008 (EC) No 882/2004 of the European Pa 90/425/EEC, 91/496/EEC, 96/23/EC, 96 Regulation) (OJ L 95, 7.4.2017, p. 1), and I certify that the products described above — that they come from (an) establishm control points (HACCP) principles in — that they have been handled and, accordance with the requirements of	auary 2002 laying down the general prin Authority and laying down procedures (2004 of the European Parliament and co (24, p. 1) and Regulation (EC) No 853/20 (specific hygiene rules for food of animal of n Parliament and of the Council of 15 Ma (e) application of food and feed law, rules (e) No 652/2014, (EU) 2016/429 a (EQ) No 652/2014, (EU) 2016/429 a (E) No 652/2014, (EU) 2016/429 a (E) No 1/2005 and (EC) (E) No 1/2005 and (EC) (E) No 1/2005 and (EC) (E) No 97/78/EC and repealine (E) and 97/78/EC and Council De (E) avere produced in accordance with these (E) implementing a programme bas (E) accordance with Article 5 of Regulation (E) where appropriate, prepared, packaged	nciples and requirements of food law, is in matters of food safety (OJ L 31 of the Council of 29 April 2004 on the 004 of the European Parliament and or origin (OJ L 139, 30.4.2004, p. 55) and arch 2017 on official controls and other is on animal health and welfare, plan EC) No 396/2005, (EC) No 1069/2009 and (EU) 2016/2031 of the Europear No 1099/2009 and Council Directives ing Regulations (EC) No 854/2004 and Directives 89/608/EEC, 89/662/EEC ecision 92/438/EEC (Official Controls e requirements, in particular: wed on the hazard analysis and critica (EC) No 852/2004; d and stored in a hygienic manner in
_		
ficates for certain animals and goods and 2016/759 as regards these model certifica I: Box reference I.25: Insert the appropriate Ha II:	d amending Regulation (EC) No 2074 Ites (OJ L 131, 17.5.2019, p. 101). armonised System (HS) code(s) of the W	/2005 and Implementing Regulation
		Signature:
	 icates for certain animals and goods an 2016/759 as regards these model certifica l: Box reference I.25: Insert the appropriate Ha II: The colour of the stamp and signature must al veterinarian Name (in capital letters): Date: 	Box reference I.25: Insert the appropriate Harmonised System (HS) code(s) of the W II: The colour of the stamp and signature must be different from that of the other particu al veterinarian Name (in capital letters):

PART XV

MODEL OFFICIAL CERTIFICATE FOR THE ENTRY INTO THE UNION FOR PLACING ON THE MARKET OF SPROUTS AND SEEDS INTENDED FOR THE PRODUCTION OF SPROUTS

cou	NTRY										Official certificate to the EU
	l.1.	Consignor/Exporter						I.2.	Certificate referen	ice No	I.2.a IMSOC reference No
		Name				I.3. Central Competent Authority					
		Address						1.4.	Local Competent	Authority	,
ment		Tel. No									
sign	I.5.	Consignee/Importer						1.6.	Operator respons	ible for th	ne consignment
ned con		Name							Name		
patch		Address							Address		
f dis		Postal code							Postal code		
uils o		Tel. No									
Part I: Details of dispatched consignment	1.7.	Country of origin	ISO	1.8.				1.9.	Country of destination	ISO	I.10.
Ра	l.11	Place of dispatch						I.12.	Place of destination	on	
		Name		Approv	val No				Name		
		Address							Address		
	l.13.	Place of loading						l.14.	Date and time of o	departure	•
	l.15.	Means of transport						l.16.	Entry BCP		
		Aeroplane	Ves	sel		Other		I.17.	Accompanying do	ocuments	5
		Road vehicle	Rai	lway					Туре		
		Identification:							No		
	l.18.	Transport conditions									
		Ambient	Chi	lled		Frozen					
	l.19.	Container No/Seal No	<u> </u>					1			

COUNTRY				Official certificate to the EU
I.20. Goods certified as				
Human consumption				
I.21.			1.22.	
I.23. Total number of packages	1.24. 0	Quantity		
	Т	otal number	Total net weight (Kg)	Total gross weight (Kg)
I.25. Description of goods				
No Code and CN	title			
Species (Scientific name)		Ма	nufacturing plant	Cold store
Final consumer Number of package		Net weight	Batch No	Type of packaging

	Certificate for the entry into the Union for placing on the market of sprou COUNTRY seeds intended for the production of sp				
	II .	Health information	II.a. Certificate reference No	II.b.	
		I, the undersigned official inspector, hereby No 852/2004 and certify that:	declare that I am aware of the relev	ant provisions of Regulation (EC)	
	II.1.1. (¹) the seeds described above were produced under conditions which comply with Regulation (EC) No 852/2004 and particular with the general hygiene provisions for primary production and associated operations set out in Part A Annex I thereto;				
tion	II.1.2. (¹)	the sprouts were produced in establishments Commission Regulation (EU) No 210/2013 o pursuant to Regulation (EC) No 852/2004 of th	f 11 March 2013 on the approval of	establishments producing sprouts	
Part II: Certification	II.1.3. (¹) the sprouts were produced under conditions which comply with the traceability requirements laid down in Commission Implementing Regulation (EU) No 208/2013 of 11 March 2013 on traceability requirements for sprouts and seeds intended for the production of sprouts (OJ L 68, 12.3.2013, p. 16).and respect the microbiological criteria laid down in Annex I to Commission Regulation (EC) No 2073/2005 of 15 November 2005 on microbiological criteria for foodstuffs (OJ L 338, 22.12.2005, p. 1).				
	Notes See notes in Annex II of Commission Implementing Regulation (EU) 2019/628 of 8 April 2019 concerning model official certificates for certain animals and goods and amending Regulation (EC) No 2074/2005 and Implementing Regulation (EU) 2016/759 as regards these model certificates (OJ L 131, 17.5.2019, p. 101).				
	Part I:				
	 Box reference I.25: Insert the appropriate HS code(s) such as: 0704 90, 0706 90, 0708 10, 0708 20, 0708 90, 0713 10 0713 33, 0712 34, 0712 35, 0713 39, 0713 40, 0712 50, 0712 60, 0713 90, 0910 99, 1201 10, 1201 90, 1207 50, 1207 99 1209 10, 1209 21, 1209 91 or 1214 90. 				
	— Box	reference I.25: Manufacturing plant: insert the	name of the establishments which pro	oduced the sprouts or seeds.	
	Part II:				
	(¹) Dele	ete as appropriate (e.g. if sprouts or seeds).			
		colour of the signature shall be different to the embossed or are a watermark.	at of the printing. The same rule appli	es to stamps other than those that	
	Official in	spector			
	Nan	ne (in capital letters):		Qualification and title:	
	Date	9:		Signature:	
	Star	np:			

ANNEX IV

MODEL OFFICIAL CERTIFICATES IN THE CASE OF ANTE-MORTEM INSPECTION AT THE HOLDING OF PROVENANCE

Part I: MODEL OFFICIAL CERTIFICATE FOR LIVE ANIMALS

OFFICIAL CERTIFICATE

for live animals transported to the slaughterhouse in the case of ante-mortem inspection at the holding of provenance in accordance with Article 5(2)(f) of Commission Delegated Regulation (EU) 2019/624 (¹).

Na	ame of the official veterinarian:						
No);						
1.	Identification of the animals						
	Species:						
	Number of animals:						
	Identification marking:						
2.	Provenance of the animals						
	Address of holding of provenance:						
	Identification of house (*):						
3.	Destination of the animals						
	The animals will be transported to the following slaughterhouse:						
	by the following means of transport:						
4.	Other relevant information						
5.							
	I, the undersigned, declare that:						
	 the animals described above were examined before slaughter at the above-mentioned holding at						
	— the following observations on the health and welfare of animals were made:						
	 the records and documentation concerning these animals satisfied the legal requirements and do not prohibit the slaughter of the animals, 						
	— I verified the food chain information						
Do	ne at:,						
	(Place)						
on	:						
	(Date)						
Sta	amp						
	(Signature of official veterinarian)						
(*)	optional						

^{(&}lt;sup>1</sup>) Commission Delegated Regulation (EU) 2019/624 of 8 February 2019 concerning specific rules for the performance of official controls on the production of meat and for production and relaying areas of live bivalve molluscs in accordance with Regulation (EU) 2017/625 of the European Parliament and of the Council (OJ L 131, 17.5.2019, p. 1).

Part II: MODEL OFFICIAL CERTIFICATE FOR POULTRY INTENDED FOR THE PRODUCTION OF FOIE GRAS AND DELAYED EVISCERATED POULTRY

OFFICIAL CERTIFICATE

for poultry intended for the production of foie gras and delayed eviscerated poultry slaughtered at the holding of provenance in accordance with Article 6(2) of Commission Delegated Regulation (EU) 2019/624 (¹).

Na	me of the official veterinarian:
No	:
1.	Identification of uneviscerated carcasses
	Species:
	Number:
2.	Provenance of uneviscerated carcasses
	Address of holding:
3.	Destination of uneviscerated carcasses
	The uneviscerated carcasses will be transported to the following cutting plant:
4.	Declaration
	I, the undersigned, declare that:
	 the uneviscerated carcasses described above are of birds which were examined before slaughter on the above- mentioned holding at (time) on
	— the following observations on the health and welfare of animals were made:
	 the records and documentation concerning these animals satisfied the legal requirements and did not prohibit the slaughter of the birds
Do	ne at:,
	(Place)
on	
	(Date)
Sta	amp
	(Signature of official veterinarian)

⁽Signature of official veterinarian)

Commission Delegated Regulation (EU) 2019/624 of 8 February 2019 concerning specific rules for the performance of official controls on the production of meat and for production and relaying areas of live bivalve molluscs in accordance with Regulation (EU) 2017/625 of the European Parliament and of the Council (OJ L 131, 17.5.2019, p. 1). (1)

Part III: MODEL OFFICIAL CERTIFICATE FOR FARMED GAME SLAUGHTERED AT THE HOLDING OF PROVENANCE	
OFFICIAL CERTIFICATE	
for farmed game slaughtered at the holding in accordance with Article 6(3) of Commission Delegated Regulation(EU) 2019/624 (¹).	
Name of the official veterinarian:	
No:	
1. Identification of the animals	
Species:	
Number of animals:	
Identification marking:	
2. Provenance of the animals	
Address of holding of provenance:	
Identification of house (*):	
3. Destination of the animals	
The animals will be transported to the following slaughterhouse:	
by the following means of transport:	
4. Other relevant information	
5. Declaration	
I, the undersigned, declare that:	
(1) the animals described above were examined before slaughter at the above-mentioned holdi at (time) on	
(2) they were slaughtered at the holding at (time) on (date) and the slaughter a bleeding were carried out correctly,	۱d
(3) the following observations on the health and welfare of animals were made:	
(4) the records and documentation concerning these animals satisfied the legal requirements and did not prohibit t slaughter of the animals.	٦e
Done at:	,
(Place)	
on:	
(Date)	
Stamp	
(Signature of official veterinarian)	
(*) optional	

⁽¹⁾ Commission Delegated Regulation (EU) 2019/624 of 8 February 2019 concerning specific rules for the performance of official controls on the production of meat and for production and relaying areas of live bivalve molluscs in accordance with Regulation (EU) 2017/625 of the European Parliament and of the Council (OJ L 131, 17.5.2019, p. 1).

Part IV: MODEL OFFICIAL CERTIFICATE FOR FARMED GAME SLAUGHTERED AT THE HOLDING in accordance with point 3a of Section III of Annex III to Regulation (EC) No 853/2004

OFFICIAL CERTIFICATE

For farmed game slaughtered on the holding in accordance with point 3a of Section III of Annex III to Regulation (EC) No 853/2004 and Article 6(4) of Commission Delegated Regulation (EU) 2019/624 (¹).

Na	Name of the official veterinarian:			
Nc	ť			
1.	Identification of the animals			
	Species:			
	Number of animals:			
	Identification marking:			
2.	Provenance of the animals			
	Address of holding of provenance:			
	Identification of house (*):			
3.	Destination of the animals			
	The animals will be transported to the following slaughterhouse:			
	by the following means of transport:			
4.	Other relevant information			
_				
5.	Declaration			
	I, the undersigned, declare that:			
	(1) the animals described above were examined before slaughter at the above-mentioned holding at (time) on			
	(2) the following observations on the health and welfare of animals were made:			
	(3) the records and documentation concerning these animals satisfied the legal requirements and did not prohibit the slaughter of the animals.			
Do	ne at:,			
	(Place)			
on				
	(Date)			
Sta	amp			
	(Signature of official veterinarian)			
(*)	optional			

^{(&}lt;sup>1</sup>) Commission Delegated Regulation (EU) 2019/624 of 8 February 2019 concerning specific rules for the performance of official controls on the production of meat and for production and relaying areas of live bivalve molluscs in accordance with Regulation (EU) 2017/625 of the European Parliament and of the Council (OJ L 131, 17.5.2019, p. 1).

ANNEX V

MODEL OFFICIAL CERTIFICATE IN THE CASE OF EMERGENCY SLAUGHTER OUTSIDE THE SLAUGHTERHOUSE IN ACCORDANCE WITH ARTICLE 4 OF COMMISSION DELEGATED REGULATION (EU) 2019/624 (1)

MODEL OFFICIAL CERTIFICATE IN THE CASE OF EMERGENCY SLAUGHTER OUTSIDE THE SLAUGH-TERHOUSE

OFFICIAL CERTIFICATE

	In the case of emergency slaughter outside the slaughterhouse
Nar	ne of the official veterinarian:
No:	
1.	Identification of the animals
	Species:
	Number of animals:
	Identification marking:
2.	Place of emergency slaughter
	Address:
	Identification of house (*):
3.	Destination of the animals
	The animals will be transported to the following slaughterhouse:
	by the following means of transport:
4.	Other relevant information
5.	Declaration
	I, the undersigned, declare that:
	(1) the animals described above were examined before slaughter at the above-mentioned holding at (time) on
	(2) they were slaughtered at (time) on (date) and the slaughter and bleeding were carried out correctly,
	(3) the following was the reason for the emergency slaughter:
	(4) the following observations on the health and welfare of animals were made:
	(5) The following treatments were administered to the animal(s):
	(6) the records and documentation concerning these animals satisfied the legal requirements and did not prohibit the slaughter of the animals.
Dor	ne at:,
	(Place)
on:	
Cto.	(Date)
Sta	
	(Signature of official veterinarian)

(*) optional

⁽¹⁾ Commission Delegated Regulation (EU) 2019/624 of 8 February 2019 concerning specific rules for the performance of official controls on the production of meat and for production and relaying areas of live bivalve molluscs in accordance with Regulation (EU) 2017/625 of the European Parliament and of the Council (OJ L 131, 17.5.2019, p. 1).

ANNEX VI

CORRELATION TABLE REFERRED TO IN ARTICLE 32

Regulation (EU) No 211/2013	This Regulation
Article 1	Article 1(2)(b)(ii)
Article 2	Article 2(2)
Article 3	Article 27
Article 4	-
Article 5	_
Annex	Part XV of Annex III