Model health certificate for fat derivatives not intended for human consumption to be used outside the feed chain (FD)

GBHC512 v1.0 Aug-23

Part I. Details of dispatched consignment

I.1 Consignor Name: Address: Tel:	**I.2 Certificate reference no.**	**I.3 Central competent authority**
	I.2.a Not in use	**I.4 Local competent authority**

I.5 Consignee Name: Address: Tel:	**I.6 Person responsible for the load in Great Britain** Name: Address: Tel:

I.7 Country of origin	**ISO code**	**I.8 Region of origin**	**Code**	**I.9 Country of destination**	**ISO code**	**I.10 Region of destination**	**Code**

I.11 Place of origin Name: Approval number: Address: Name: Approval number: Address: Name: Approval number: Address:	**I.12 Place of destination** ☐ Custom warehouse Name: Approval number: Address:
I.13 Place of loading	**I.14 Date of departure**
I.15 Means of transport ☐ Aeroplane ☐ Ship ☐ Railway wagon ☐ Road vehicle ☐ Other Identification: Documentation references:	**I.16 Entry BCP** **I.17** Not in use

II.a. Certificate reference no.	**II.b.**

I.18 Description of commodity		
I.19 Commodity code (HS code)	**I.21 Temperature of products** ☐ Ambient ☐ Chilled ☐ Frozen	**I.23 Seal / Container No.**
I.20 Quantity	**I.22 Number of packages**	**I.24 Type of packaging**

I.25 Commodity certified for

☐ Technical use

I.26 ☐ **For transit through Great Britain to third country** **Third country** **ISO Code**	**I.27** ☐ **For import or admission into Great Britain**

I.28 Identification of the commodities

Species (Scientific name)	Manufacturing plant	Number of packages	Net weight	Batch number

Part II. Certification

Animal Health

I, the undersigned official veterinarian, declare that I am aware of the relevant requirements of the GB regulations, and certify that the fat derivatives described in Part I of this certificate satisfy the health requirements below:

AH/E105 Establishment requirements

Have been prepared and stored in a plant approved, validated and supervised by the competent authority in accordance with GB requirements;

AH/P107 Product requirements

have been prepared from rendered fats exclusively produced from the following animal by-products as set out in the notes for completion:

(a) in case the fat derivatives are intended for uses outside the feed chain, other than in organic fertilisers, soil improvers, cosmetics, pharmaceuticals and medical devices, the following Category 1 materials: (*)[A] (*)[B] (*)[C];

(b) in case the fat derivatives are intended for use in organic fertilisers or soil improvers or other uses outside the feed chain, other than in cosmetics, pharmaceuticals and medical devices, the following Category 2 materials: (*)[D] (*)[E] (*)[F];

(c) the following Category 3 materials: (*)[G] (*)[H] (*)[I] (*)[J] (*)[K] (*)[L] (*)[M] (*)[N] (*)[O] (*)[P];

II.a. Certificate reference no.	**II.b.**

AH/P156 Product requirements

consist of fat derivatives intended for purposes outside the feed chain, other than in cosmetics, pharmaceuticals and medical devices;

AH/P706 Product requirements

in case of fat derivatives produced from animal by-products referred to in AH/P107 (a) and AH/P074 (b):

(a) have been produced using the following methods:

(*)***EITHER*** [transesterification or hydrolysis at least 200°C, under corresponding appropriate pressure, for 20 minutes (glycerol, fatty acids and esters);]

(*)***OR*** [saponification with NaOH 12M (glycerol and soap):

(*)***EITHER*** [in a batch process at 95°C for three hours;]

(*)***OR*** [in a continuous process at 140°C, 2 bars (2000 hPa) for eight minutes;]

(*)***OR*** [hydrogenation at 160°C at 12 bars (12000 hPa) pressure for 20 minutes;]]

(b) are packaged in new containers or in containers that have been cleaned, and all precautions are taken to prevent its contamination which bear labels indicating "NOT FOR HUMAN OR ANIMAL CONSUMPTION";

AH/P707 Product requirements

in case of fat derivatives produced from animal by-products referred to in AH/P107 (c), the fat derivatives have been produced in accordance with one of the processing methods (*)[1]-(*)[2]-(*)[3]-(*)[4]-(*)[5]-(*)[6]-(*)[7] in accordance with GB requirements;

(*) Keep as appropriate.

Official Veterinarian / Official Inspector

By signing this certificate, I certify that the requirements laid out above and in the accompanying notes for completion have been met.

Name (in capital letters):

Qualification and title:

Date:

Signature:

Stamp:

Part III. Notes for completion

These notes for completion must be read and understood by the certifying officer before signing the certificate. Notes are set out in sections that correspond to the sections in the certificate. By signing this certificate, certifiers are verifying that the consignment meets the requirements set out in the certificate and any relevant corresponding notes for completion.

These notes do not need to be printed as part of a paper certificate that accompanies the consignment or in any electronic copy of the certificate.

References to European Union legislation within this certificate are references to direct EU legislation which has been retained in Great Britain (retained EU law as defined in the European Union (Withdrawal) Act 2018) and can be viewed on the UK legislation website (legislation.gov.uk).

References to Great Britain in this certificate include Channel Islands and Isle of Man.

References to GB requirements refer to the requirement(s) of Great Britain as set out in the accompanying notes for completion.

Part I

Box reference I.6: *Person responsible for the consignment in Great Britain*: this box is to be filled in only if it is a certificate for transit commodity; it may be filled in if the certificate is for import commodity.

Box reference I.12: *Place of destination*: this box is to be filled in only if it is a certificate for transit commodity. Products in transit can only be stored in free zones, free warehouses and custom warehouses.

Box reference I.15: Registration number (railway wagons or container and lorries), flight number (aircraft) or name (ship) is to be provided. In case of unloading and reloading, the consignor must inform the BCP of entry into Great Britain.

Box reference I.19: Use the appropriate Harmonized System (HS) code under the following headings: 15.16 or 15.08.

Box reference I.23: For bulk containers, the container number and the seal number (if applicable) must be included.

Box reference I.25: *Technical use*: any use other than for animal consumption.

Box reference I.26 and I.27: Fill in according to whether it is a transit or an import certificate.

Box reference I.28: *Species*: select from the following: *Ruminantia*, *Other*;

Manufacturing plant: provide the registration number of treatment/processing establishment.

Part II

Animal Health

By signing this certificate, you, the official veterinarian, are certifying that you have read and understood Regulation (EC) No 1069/2009 of the European Parliament and of the Council and in particular Article 10 thereof, and Commission Regulation (EU) No 142/2011, and in particular Chapter II of Annex XIV thereto.

AH/E105 Establishment requirements

GB Requirements

The plant where the products have been prepared and stored must be approved and supervised by the competent authority in accordance with Article 24 of Regulation (EC) No 1069/2009 in order to kill pathogenic agents.

AH/P107 Product requirements

(a) One or more options can be selected.

A: The following material:

(i) specified risk material;

(ii) entire bodies or parts of dead animals containing specified risk material at the time of disposal.

B: Animal by-products which have been derived from animals which have been submitted to illegal treatment as defined in Article 1(2)(d) of Directive 96/22/EC or Article 2(b) of Directive 96/23/EC.

C: Animal by-products containing residues of other substances and environmental contaminants listed in Group B (3) of Annex I to Directive 96/23/EC, if such residues exceed the permitted levels laid down by retained EU law or, by UK legislation.

(b) One or more options can be selected.

D: Animal by-products containing residues of authorised substances or contaminants exceeding the permitted levels referred to in Article 15(3) of Directive 96/23/EC.

E: Products of animal origin which have been declared unfit for human consumption due to the presence of foreign bodies in those products.

F: Animals and parts of animals, other than those referred to in Articles 8 and 10 of Regulation (EC) No 1069/2009, that died other than being slaughtered or killed for human consumption, including animals killed for disease control purposes.

(c) One or more options can be selected.

G: Carcases and parts of animals slaughtered or, in the case of game, bodies or parts of animals killed, and which are fit for human consumption in accordance with retained EU law, but are not intended for human consumption for commercial reasons.

H: Carcases and the following parts originating either from animals that have been slaughtered in a slaughterhouse and were considered fit for slaughter for human consumption following an ante-mortem inspection or bodies and the following parts of animals from game killed for human consumption in accordance with retained EU law:

(i) carcases or bodies and parts of animals which are rejected as unfit for human consumption in accordance with retained EU law, but which did not show any signs of disease communicable to humans;

(ii) heads of poultry;

(iii) hides and skins, including trimmings and splitting thereof, horns and feet, including the phalanges and the carpus and metacarpus bones, tarsus and metatarsus bones;

(iv) pig bristles;

(v) feathers.

I: Blood of animals which did not show any signs of disease communicable through blood to humans or animals, obtained from animals that have been slaughtered in a slaughterhouse after having been considered fit for slaughter for human consumption following an ante-mortem inspection in accordance with retained EU law.

J: Animal by-products arising from the production of products intended for human consumption, including degreased bone, greaves and centrifuge or separator sludge from milk processing.

K: Products of animal origin, or foodstuffs containing products of animal origin, which are no longer intended for human consumption for commercial reasons or due to problems of manufacturing or packaging defects or other defects from which no risk to public or animal health arises.

L: Petfood and feedingstuffs of animal origin, or feedingstuffs containing animal by-products or derived products, which are no longer intended for feeding for commercial reasons or due to problems of manufacturing or packaging defects or other defects from which no risk to public or animal health arises.

M: Blood, placenta, wool, feathers, hair, horns, hoof cuts and raw milk originating from live animals that did not show signs of any disease communicable through that product to humans or animals.

N: Aquatic animals, and parts of such animals, except sea mammals, which did not show any signs of diseases communicable to humans or animals.

O: Animal by-products from aquatic animals originating from plants or establishments manufacturing products for human consumption.

P: The following material originating from animals which did not show any signs of disease communicable through that material to humans or animals:

(i) shells from shellfish with soft tissue or flesh;

(ii) the following originating from terrestrial animals;

- hatchery by-products,
- eggs,
- egg by-products, including egg shells;

(iii) day-old chicks killed for commercial reasons.

AH/P156 Product requirements

No further notes for completion.

AH/P706 Product requirements

No further notes for completion.

AH/P707 Product requirements

The fat derivatives must have been produced in accordance with one of the processing methods [1]-[2]-[3]-[4]-[5]-[6]-[7] referred to in Chapter III of Annex IV to Regulation (EU) No 142/2011.