Model health certificate for egg products not intended for human consumption that could be used as feed (EP-A)

GBHC588 v1.1 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

☐ Animal feedingstuff

☐ 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

Approval number of establishments / 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 egg products referred to in Part I of this certificate comply with the following health requirements:

AH/E107 Establishment requirements

have been prepared and stored in a plant, approved, validated and supervised by the competent authority in accordance with GB regulations, in order to kill pathogenic agents;

AH/P007 Product requirements (segregation)

has undergone all precautions to avoid contamination with pathogenic agents after treatment;

AH/P101F Product requirements (composition)

have been prepared (derived) exclusively with the following animal by-products as set out in the notes for completion: (*)[A] (*)[B] (*)[C];

AH/P154 Product requirements

consist exclusively of egg products not intended for human consumption;

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

AH/P506 Packaging and labelling

the end product was:

(*)***EITHER*** [packaged in new or sterilised bags;]

(*)***OR*** [transported in bulk in containers or other means of transport that were thoroughly cleaned and disinfected with a disinfectant approved by the competent authority before use;]

and which bear labels indicating 'NOT FOR HUMAN CONSUMPTION';

AH/P550B Storage

the end product was stored in enclosed storage;

AH/P614 Product requirements

meet GB standards on residues of substances that are harmful or might alter the organoleptic characteristics of the product or make its use as feed dangerous or harmful to animal health;

AH/P704 Product requirements

have been subjected to processing in accordance with GB requirements:

(*)***EITHER*** [**(a)** with processing method (*)[1] (*)[2] (*)[3] (*)[4] (*)[5] (*)[7];]

(*)***OR*** [**(b)** to a method and parameters which ensure that the products comply with the GB microbiological standards;]

(*)***OR*** [**(c)** as fit for human consumption;]

AH/P800B Testing

the competent authority examined a random sample of the products immediately prior to dispatch and found it to comply with microbiological GB requirements for *Salmonella* and *Enterobacteriaceae;*

(*) 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. The products in transit can only be stored in free zones, free warehouses and custom warehouses.
Box reference I.15:	Registration number (railway wagons or container and lorries), flight number (aircraft) or name (ship) is to be provided. In case of unloading and reloading, the consignor must inform the border control post of the point of entry into Great Britain.
Box reference I.19:	Use the appropriate Harmonized System (HS) code under the following headings: 04.08, 23.09 or 35.02.
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.

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, and in particular Article 10 thereof, and Commission Regulation (EU) No 142/2011, and in particular Chapter I of Annex XIV thereto.

AH/E107 Establishment requirements

GB regulations refers to Article 24 of Regulation (EC) No 1069/2009 or Article 4(2) of Regulation (EC) No 853/2004.

AH/P007 Product requirements (segregation)

No further notes for completion.

AH/P101F Product requirements (composition)

One or more options can be selected.

A: Animal by-products arising from the production of products intended for human consumption.

B: 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.

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

(i) hatchery by-products;

(ii) eggs;

(iii) egg by-products, including egg shells.

AH/P154 Product requirements

No further notes for completion.

AH/P506 Packaging and labelling

No further notes for completion.

AH/P550B Storage

No further notes for completion.

AH/P614 Product requirements

No further notes for completion.

AH/P704 Product requirements

(a) Processing method 1 to 5 or 7 described in Chapter III of Annex IV to Regulation (EU) No 142/2011.

(b) Microbiological standards set out in Chapter I of Annex X, to Regulation (EU) No 142/2011.

(c) Processed as fit for human consumption in compliance with Section X, Chapters I and II of Annex III to Regulation (EC) No 853/2004.

AH/P800B Testing

The products must be examined immediately prior to dispatch under the responsibility of the competent authority by taking a random sample during or on removal from storage which complies with the following standards:

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

Where:

n = number of samples to be tested;

m = threshold value for the number of bacteria; the result is considered satisfactory if the number of bacteria in all samples does not exceed m;

M = maximum value for the number of bacteria; the result is considered unsatisfactory if the number of bacteria in one or more samples is M or more; and

c = number of samples the bacterial count of which may be between m and M, the sample still being considered acceptable if the bacterial count of the other samples is m or less.