Veiledning for utfylling av 5.1.344 Storbritannia, helsesertifikat, fiskeolje NHC, GBHC520

Veiledning om utfylling av del 1.

Veiledning for hva som skal fylles ut i de ulike feltene i del 1 av sertifikatet finnes her: https://www.gov.uk/government/publications/how-to-complete-a-health-certificate-for-imports-to-great-britain/how-to-complete-a-health-certificate-for-imports-to-great-britain

Det er eksportør som er ansvarlig for at innholdet i sertifikatet er korrekt.

Veiledning om utfylling av del 2.

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); information is to be provided in case of unloading and reloading.

Box reference I.19: Use the appropriate HS code: 15.04 or 15.18.

Box reference I.23: For bulk containers, the container number and the seal number (if applicable) should 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: *Manufacturing plant*: provide the registration number of the 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, and in particular Article 10 of that Regulation, and Commission Regulation (EU) No 142/2011, and in particular Chapter II of Annex XIV thereto.

AH/E106 Establishment requirements

The fish plant must be approved, validated and supervised by the competent authority in accordance with Article 24 of Regulation (EC) No 1069/2009.

AH/P101E 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: Aquatic animals, and parts of such animals, except sea mammals, which did not show any signs of diseases communicable to humans or animals.

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

AH/P152 Product requirements

No further notes for completion.

AH/P512 Processing, packaging and labelling

(a) GB processing requirements as set out in Annex X, Chapter II, Section 3 of Regulation (EU) No 142/2011, in order to kill pathogenic agents.