



SANITARY CERTIFICATE

for processed fish meal and hydrolyzed fish proteins intended for human consumption

NORWAY

Reference number: _____

| | | | |
|-------------------------|--|-----------------------------------|--|
| Country of dispatch: | NORWAY | | |
| Competent authority: | NORWEGIAN FOOD SAFETY AUTHORITY, N-2381 BRUMUNDDAL, NORWAY | | |
| Inspection body: | NORWEGIAN FOOD SAFETY AUTHORITY, DISTRICT OFFICE | | |
| Phone: + 47 22 40 00 00 | Facsimile: + 47 23 21 68 01 | E-mail: postmottak@mattilsynet.no | |

I. Identification of protein or product

| Product description: | Type of packaging: | Number of packages: | Net weight: |
|----------------------|--------------------|---------------------|-------------|
| | | | |
| | | | |
| Sum: | | | |

II. Origin of protein or product

Address and approval number of preparation or processing establishment: _____

Name and address of consignor: _____

III. Destination of protein or product

The product is to be dispatched from: _____

(Place of dispatch)

to: _____

(Country and place of destination)

by the following means of transport: _____

Name of consignee and address at place of destination: _____

IV. Attestation

The undersigned official inspector hereby certifies that the product described above:

- 1) has been handled, prepared, processed, marked, packaged, stored and transported in accordance with the relevant provisions of Regulations (EC) No 178/2002, (EC) No 852/2004, (EC) No 853/2004 and (EC) No 854/2004;
- 2) contains exclusively non-mammalian and non-ruminant protein;
- 3) was entirely derived from fresh fish material which have been found fit for human consumption;
- 4) was produced in a processing establishment dedicated to fish meal production. Material of other animals, including ruminants, or poultry is not received, stored or used in this establishment;
- 5) the end product was packaged in new packing material¹ or in the case of dispatch as bulk transport: container or any other means of transport was thoroughly cleaned and disinfected with a disinfectant approved by the competent authority before use;
- 6) the end product was stored in enclosed storages;
- 7) the end product has undergone all precautions to avoid recontamination with pathogenic agents after the treatment.

Done at _____ on _____
(Place) (Date)

Seal² _____
(Signature of official inspector) (Name and qualifications in capitals)

¹ Delete as appropriate

² The signature and the stamp must be in a colour different to that of the printing.

