CHAPTER 2, STANDARD 7

GENERAL CANNED FINFISH STANDARD

1. INTRODUCTION

This general standard for canned finfish derives its authority from the Fish Inspection Regulations. It defines minimum acceptability for taint, decomposition, unwholesomeness and other requirements, other than weight, as defined in the Fish Inspection Act and Regulations and describes methods for determining that acceptability.

2. SCOPE

This standard applies to canned finfish in hermetically sealed containers. It is intended to be used for the inspection of canned finfish species for which specific Canadian product standards have not been elaborated.

Canned finfish shall be prepared from sound, wholesome raw material processed using good manufacturing practices.

Documents used to determine good manufacturing practice and compliance include:


2) Metal Can Defects Identification and Classification Manual, Canadian Food Inspection Agency.


3. NOMENCLATURE

The name of the product shall be that recognized in common usage in Canada, and in accordance with the Fish Inspection Regulations and any requirements of the applicable Codex Alimentarius Recommended International Standard.
Descriptive terms shall be used where necessary to accurately describe the contents of the can.

4. FORMS OF PRODUCT PRESENTATION

Canned finfish may be prepared from fish which is fresh, frozen, cooked or smoked.

4.1 Packing Media

The product may be presented in one of the following packing media as appropriate to the species and style of pack, with or without permitted optional ingredients:

a) Own juice
   Fish packaged without added liquid;

b) Potable water
   In conformity with the requirements of the Fish Inspection Regulations for water used in registered establishments;

c) Spring water or mineral water
   Potable water from an underground source but not obtained from a public community water supply and which meets the requirements of Section B12.001 of the Food and Drug Regulations;

d) Vegetable broth
   The liquid arising from the cooking of sound wholesome vegetables in water and which may be prepared from one or more types of vegetables. Vegetable broth may also be prepared from hydrolysed vegetable protein but a broth so prepared requires its components to be declared in a list of ingredients;

e) Olive oils
   In conformity with:
   
   the relevant sections of Division 9 of the Canadian Food and Drug Regulations; or
   
   the Recommended International Standard for Olive Oil, Virgin and Refined, and for Refined Olive-Residue Oil (Ref. CAC/RS 33-1970);

f) Other vegetable oils
   Clear, refined, deodorized, edible vegetable oil in conformity with:
   
   the relevant sections of Division 9 of the Canadian Food and Drug Regulations; or
the relevant recommended International standards adopted by the Codex Alimentarius Commission;

g) Sauces
A thickened liquid made from acceptable food ingredients giving a characterizing flavour and odour to the product;

h) Marinades
A thin liquid made from acceptable food ingredients, usually containing a sweetener, an acid solution or an alcoholic solution, with or without spices, herbs, seasonings, vegetables, and other condiments;

i) Fish oils
Clear, refined, edible fish (marine) oil. The species from which the oil is derived should be noted on the product label.

4.2 Optional Ingredients

a) salt;

b) natural starches; and

c) other optional ingredients provided that all ingredients are suitable for human consumption, are free from abnormal taste, flavour, or odour, and are permitted in Division 21 of the Food and Drug Regulations. Examples of such ingredients are spices, herbs, vegetable seasonings, vinegar and wine, and vegetables and fruits for decorative and flavouring purposes only.

4.3 Other Presentations

Any other presentation of the product may be permitted provided that it:

a) is sufficiently distinctive from the forms of presentation set out above;

b) meets all other Canadian Regulatory requirements; and

c) is adequately described on the label and in accordance with all regulatory labelling requirements.

5. SAMPLING

The sampling and tolerance plans at the front of this manual shall be used to determine the acceptability of the lot. The sampling plans dictate the minimum sample size to be taken. If necessary, in the opinion of the inspector, more than the minimum sample size specified may
be taken.

5.1 Sampling of lots for the sensory examination of the product shall be in accordance with the FAO/WHO Codex Alimentarius Sampling Plan for Prepackaged Foods (AQL 6.5) (CAC/RM42-1969) except that a lower acceptance number for decomposition shall be used as indicated in the sampling tables.

The sampling tables specify the minimum number of sample units to be used for the following types of inspections:

a) Level I - Sensory examination of all products subject to inspection other than lots which are subject to reinspection.

b) Level II - Sensory examination of all products which are under reinspection.

5.2 Size of Sample Unit

The sample unit shall consist of a can of fish and the entire contents thereof.

6. DESCRIPTION OF DEFECTS

6.1 Taint

A unit will be considered tainted when any of the following conditions are found:

a) **Rancid**
   The contents in the container show the following defects:
   Odour characterized by the distinct or persistent odour of oxidized oil; or
   Flavour characterized by that of oxidized oil which leaves a distinct bitter aftertaste.

b) **Abnormal**
   Distinct and persistent uncharacteristic odours or flavours such as burnt or acrid, metallic, or associated with feed and not defined as rancid or decomposed; or
   Flavour or odour resulting from the improper addition and/or mixing of ingredients.
6.2 Decomposition

A unit will be considered decomposed when any of the following conditions are found:

a) Odours and flavours
   Persistent, distinct and uncharacteristic odour or flavour including but not limited to the following:
   fruity, vegetable, stale, musty, yeasty, sour, faecal, ammonia, hydrogen sulphide, bilge-like and putrid.

b) Discolouration
   Discolouration associated with decomposition which is uncharacteristic of the species and type of pack, such as flushed pink, dark brown, green or yellowish to orange colours.

c) Texture
   Breakdown of muscle structure due to decomposition characterized by:
   muscle structure which is very tough, dry, mealy or chalky; or
   muscle structure which is very soft, mushy, or pasty.

6.3 Unwholesome

a) Critical Foreign Material

   A lot will be considered defective when any of the following conditions are found:

   the presence of any material which has not been derived from fish (and packing media) and which poses a threat to human health (such as glass, etc.); or

   distinct and persistent odour or flavour of any material which has not been derived from fish (and packing media) and which poses a threat to human health (such as solvents, fuel oil, etc.).

b) Foreign Material

   A unit will be considered defective when the following condition is found:

   the presence of any material which has not been derived from fish (and packing media) but does not pose a threat to human health (such as insect pieces, sand, etc.).
c) **Other Defects**

A unit will be considered defective when any of the following conditions are found:

1) **Struvite Crystals** *(magnesium ammonium phosphate crystals)*
   Any struvite crystal greater than 5 mm in length.

2) **Sulphide Blackening** *(smut)*
   Staining affecting greater than 5% of the drained contents.

3) **Undesirable Parts**
   Any combination of head parts, heads, tails, scales and viscera exceeding 2% of the drained weight.

7. **EXAMINATION METHODS**

7.1 Complete external can examination. Open can and complete net weight determination, according to defined policies and procedures for these examinations.

7.2 Examine appearance of product in can. Carefully remove fish from can to examination tray. Inspect can contents for presence of foreign material or other undesirable parts, carefully separating fish as necessary.

7.3 Examine can interior for presence of foreign material, smut, struvite, and corrosion or other can interior defects.

7.4 Observe colour of flesh as an indicator of decomposition.

7.5 Assess odour, flavour and texture as required.

7.6 Record any defect for that unit on the appropriate worksheet.

8. **CLASSIFICATION OF "DEFECTIVES"**

A sample unit which contains defects as described in section 6 is classified as a "defective".

9. **LOT ACCEPTANCE**

A lot will be considered unacceptable when:

a) any single instance of critical foreign material occurs; or
b) the total number of sample units found defective for taint, decomposition or unwholesomeness, individually or in combination, exceeds the acceptance number for the sample size designated in the sampling plans; or

c) the total number of sample units found defective for decomposition exceeds the acceptance number shown in parentheses for the sample size designated in the sampling plans.