codex alimentarius commission

FOOD AND AGRICULTURE ORGANIZATION OF THE UNITED NATIONS

WORLD HEALTH ORGANIZATION

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Agenda Item 5

CX/FFP 02/5 PART I

JOINT FAO/WHO FOOD STANDARDS PROGRAMME CODEX COMMITTEE ON FISH AND FISHERY PRODUCTS

Twenty-fifth Session Ålesund, Norway, 3 - 7 June 2002

CODE OF PRACTICE FOR FISH AND FISHERY PRODUCTS: DRAFT SECTIONS (1, 2.1, 2.2, 2.9, 3 TO 6 AND 13) AND PROPOSED DRAFT SECTIONS (OTHER SECTIONS)¹

PART I - DRAFT SECTIONS AT STEP 6

The 24th Session of the Committee on Fish and Fishery Products considered the Proposed Draft Code of Practice for Fish and Fishery Products and advanced to Step 5 the following sections: Section 1. *Scope*; Section 2. *Definitions (General Definitions, Fresh, Frozen and Minced Fish, Canned Fish)*; Section 3. *Prerequisite Programme*; Section 4. *General Consideration for the Handling of Fresh Fish and Shellfish* Section 5. *HACCP*; Section 6. *Processing of Fresh Frozen and Minced Fish*; and Section 13. *Processing of Canned Fish and Shellfish*. The other sections of the Code were returned to Step 3 for further redrafting by the Working Group including the countries responsible for the development of specific sections under the general coordination of the United Kingdom, Canada and France (ALINORM 01/18, paras. 79-82).

The 49th (Extraordinary) Session of the Executive Committee adopted the above sections at Step 5 (ALINORM 03/3, para. 9 and 21, Appendix II) and referred the comments of Malaysia on the Scope to the CCFFP for further consideration. These comments and the comments submitted to the Committee on Food Hygiene will be presented in document CX/FFP 02/5-Add.2. Comments on the present document will be presented in CX/FFP 02/5-Add.1.

The Draft Sections of the Code of Practice for Fish and Fishery Products were circulated for comments at Step 6 in CL 2001/50-FFP (December 2001). They are reproduced in Part I - Annex 1.

Due to time constraints, there was no meeting of the Working Group including the coordinating countries. The United Kingdom, Canada and France integrated the revised sections prepared by the coordinating countries into the final text and also proposed amendments to the adopted Draft Sections (Part II - Annex 2).

The current document is presented in two parts, as follows:

Part I - Annex 1: Draft Sections (1, 2.1, 2.2, 2.9, 3 to 6 and 13)

Part II - Annex 2: Proposed Draft Sections and other amendments to the Code

Part II - Annex 3: Report of the Drafting Group Meeting (United Kingdom, Canada and France)

Governments and international organizations wishing to comment on the Draft Sections at Step 6 and on the Proposed Draft Sections at Step 3 should do so in writing, preferably by email, to the Secretary, Codex Alimentarius Commission, Joint FAO/WHO Food Standards Programme, FAO, Viale delle Terme di Caracalla 00100 Rome, Italy, Fax: +39 (06) 5705 4593, E-mail: codex@fao.org, with a copy to the Codex Contact Point for Norway, Norwegian Food Control Authority, P.O. Box 8187 Dep. 0034 Oslo, Norway, Fax: +47.23.21.70.01, E-mail: ccffp@snt.no, **no later than 10 May 2002**.

The document is presented in two parts due to its size

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DRAFT CODE OF PRACTICE FOR FISH AND FISHERY PRODUCTS DRAFT SECTIONS (1, 2.1, 2.2, 2.9, 3 TO 6 AND 13)

(At Step 6 of the Procedure)

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INTRODUCTION

This Code of Practice for Fish and Fishery Products has been developed by the Codex Committee on Fish and Fishery Products from the merging of the individual codes of listed in Appendix XII plus a section on aquaculture and a section on frozen surimi. These codes were primarily of a technological nature offering general advice on the production, storage and handling of fish and fishery products on board fishing vessels and on shore. It also deals with the distribution and retail display of fish and fishery products.

This combined Code of practice has been further modified to incorporate the Hazard Analysis Critical Control Point (HACCP) approach described in the *Recommended International Code of Practice – General Principles of Food Hygiene* (CAC/RCP 1-1969, Rev.3 1997), Annex: *HACCP System and Guidelines for its Application* (Supplement to Codex Volume 1B). A pre-requisite programme is described in the Code covering technological guidelines and the essential requirements of hygiene in the production of fish and fishery products, which are safe for human consumption, and otherwise meets the requirements of the appropriate Codex product standards. The Code also contains guidance on the use of HACCP, which is recommended to ensure the hygienic production of fish and fishery products to meet health and safety requirements.

Within this Code a similar systematic approach has been applied to essential quality, composition and labelling provisions of the appropriate Codex product standards. Throughout the code this is referred to as "Defect Action Point (DAP) Analysis".

The Codex Committee on Fish and Fishery Products recommended at its Twentieth Session that defects of a commercial nature, i.e. workmanship defects, which had been removed from Codex fish product standards, be transferred to the appropriate Codex Code of practice for optional use between buyers and sellers during commercial transactions. The Committee further recommended that this detail should be described in a section on Final Product Specifications, which now appear as Appendices II - XI of this document. A similar approach to HACCP has been incorporated into the Code as guidelines for the control of defects (DAP Analysis).

This Code will assist all those who are engaged in the handling and production of fish and/or fishery products, or are concerned with their storage, distribution, export, import and sale in attaining safe and wholesome products which can be sold on national or international markets and meet the requirements of the Codex Standards (see Appendix XII).

HOW TO USE THIS CODE

The aim of this Code is to provide a user-friendly document as background information and guidance for the elaboration of fish and shellfish process management systems which would incorporate Good Management Practice (GMP) as well as the application of HACCP in countries where these, as yet, have not been developed. In addition, it could be used for training of fishermen and employees of the fish processing industry.

The practical application of this *international* Code, with regard to *national* fisheries, would therefore require some modifications and amendments, taking into account local conditions and specific consumer requirements. This Code, therefore, is not intended to replace the advice or guidance of trained and experienced technologists regarding the complex technological and hygienic problems which might be unique to a specific geographical area or specific fishery and, in fact, is intended to be used as a supplement in such instances.

This Code is divided into separate, though interrelated, Sections. It is intended that in order to set up a HACCP or DAP programme these should be consulted as appropriate:

- (a) Section 2 Definitions Being acquainted with the definitions is important and will aid the overall understanding of the Code.
- (b) Section 3 Pre-requisite Programme Before HACCP or a similar approach can properly be applied to a process it is important that a solid foundation of good hygienic practice exists. This Sections covers the

- groundwork which should be regarded as the minimum requirements for a processing facility prior to the application of hazard and defect analyses.
- (c) Section 4 General Considerations for the Handling of Fresh Fish and Shellfish This Section provides an overall view of the potential hazards and defects which may have to be considered when building up a HACCP or DAP plan. This is not intended to be an exhaustive list but is designed to help a HACCP or DAP team to think about what hazards or defects should be considered in the fresh fish or shellfish, and then it is up to the team to determine the significance of the hazard or defect in relation to the process.
- (d) Section 5 Hazard Analysis Critical Control Point (HACCP) and Defect Action Point (DAP) Analysis Only when the groundwork in Section 3 has been satisfactorily achieved should the application of the principles outlined in Section 5 be considered. This Section uses an example of the processing of a canned tuna product to help illustrate how the principles of HACCP should be applied to a process.
- (e) Section 6 Processing of Fresh, Frozen and Minced Fish This Section forms the foundation for most of the subsequent fish and shellfish processing Sections. It deals with the major process steps in the handling of raw fish through to cold storage and gives guidance and examples on the sort of hazards and defects to expect at the various steps. This Section should be used as the basis for all the other fish processing operations (Sections 7-14) which give additional guidance specific to the appropriate product sector.

Although potential hazards and potential defects are listed for most steps in the Processing Sections, it should be noted that this is only for guidance and the consideration of other hazards and/or defects may be appropriate. Also, the format in these Processing Sections has been designed for maximum 'ease of use' and therefore the 'potential hazards' or 'potential defects' are listed only where they may be introduced into a product or where they are controlled, rather than repeating them at all the intervening processing steps.

Additionally, it must be stressed that hazards and defects, and their subsequent control or action points, are product and line specific and therefore a full critical analysis based on *Section 5* must be completed for each individual operation.

- (f) Sections 7 to 15 Specific Fish and Shellfish Processing Sections Processors operating in particular sectors will need to consult the appropriate Section to find additional information specific to that sector.
- (g) Section 16 Aquaculture Production deals with aquaculture production.
- (h) Sections 17 and 18 Transportation and Retail cover general transportation and retail issues.
- (i) Additional information will be found in the *Appendices*.

SECTION 1 SCOPE

This Code of practice applies to the growing, harvesting, handling, production, processing, storage transportation and retail of fish and fishery products from marine and freshwater sources, which are intended for human consupmtion.

SECTION 2 DEFINITIONS

For the purpose of this Code:

2.1 GENERAL DEFINITIONS

Chilled Water is clean water in which the temperature is maintained at approximatelly 0°C (32°F) or

slightly colder by the addition of ice;

Chilling is the process of cooling fish to a temperature approaching that of melting ice;

Clean Water means water from any source where harmfull microbiological contamination, substances

and/or toxic plankton are not present in such quantities as may affect the health quality of

fishery products;

Cleaning means the removal of soil, food residues, dirt, grease or other objectionable matter;

Contaminant means any biological or chemical agent, foreign matter, or other substances not

intentionally added to food which may compromise food safety or suitability;

Contamination the introduction or occurence of a contaminant in fish;

Control Measure means any action and activity that can be used to prevent or eliminate a food safety

hazard or reduce it to an acceptable level. For the purposes of this Code a control

measure is also applied to a defect.

Corrective Action means any action to be taken when the results of monitoring at the CCP indicate a loss of

control. For the purposes of this Code this also applies to a DAP.

Critical Control

Point (CCP)

a step at which control can be applied and is essential to prevent or eliminate a food

safety hazard or reduce it to an acceptable level.

Critical Limit is a criterion, which separates acceptability from unacceptability. For the purpose of this

Code this also applies to a DAP;

Decision Tree a sequence of questions applied to each process step with an identified hazard to identify

which process steps are CCPs. For the purpose of this Code this also applies to a DAP;

Decomposition is the deterioration of fish including texture breakdown and causing a persistant and

distinct objectionable odour or flavour;

Defect means a condition found in a product which fails to meet essential quality, composition

and/or labelling provisions of the appropriate Codex product standards;

Defect Action Point (DAP)

a step at which control can be applied and a defect can be prevented, eliminated or

reduced to acceptable level, or a fraud risk eliminated;

Disinfection means the reduction, by means of chemical agents and/or physical methods, the number

of micro-organisms in the environment, to a level that does not compromise food safety or

suitability;

Dressed means that portion of fish remaining after heading and gutting;

Fish means any of the cold-blooded aquatic vertebrates and aquatic invertebrates. Aquatic

mammals and amphibians are not included;

Hazard a biological, chemical or physical agent in, or condition of, food with the potential to

cause an adverse health effect;

Hazard Analysis the process of collecting and evaluating information on hazards and conditions leading to

their presence to decide which are significant for food safety and therefore should be

addressed in the HACCP plan;

Hazard Analysis Critical Control Point (HACCP) a system which identifies, evaluates, and controls hazards which are significant for food

safety;

Biotoxins means poisonous substances accumulated by fish and shellfish feeding on toxin

producing algae, or in water containing toxins produced by such organisms;

Monitor the act of conducting a planned sequence of observations or measurements of control

parameters to assess whether a CCP is under control. For the purpose of this Code this

also applies to a DAP;

Potable Water is fresh water fit for human consumption. Standards of potability should not be lower

than those contained in the latest edition of the "International Standards for Drinking

Water", World Health Organisation;

Pre-Requisite Programme is a programme that is required prior to the application of the HACCP system to ensure that a fish processing facility is operating according to the Codex Principles of Food Hygiene, the appropriate Code of Practice and appropriate food safety legislation;

Processing Facility

means any premises where fishery products are prepared, processed, chilled, frozen, packaged or stored. For the purposes of this Code, premises also includes vessels;

Raw Material are fresh and frozen fish and/or parts of fish which may be utilised to produce fish and

fishery products intended for human consumption;

Refrigerated Water is clean water cooled by a suitable refrigeration system;

Shelf-Life the period during which the product maintains its microbiological and chemical safety

and sensory qualities at a specific storage temperature. It is based on identified hazards for the product, heat or other preservation treatments, packaging method and other

hurdles or inhibiting factors that may be used;

Shellfish means those species of molluscs and crustaceans including cephalopods that are usually

used for food;

Step is a point, procedure, operation or stage in the food chain including raw materials, from

primary production to final consumption;

Validation means obtaining evidence that the elements of the HACCP plan are effective;

Verification the application of methods, procedures, tests and other evaluations, in addition to

monitoring to determine compliance with the HACCP plan. For the purposes of this

Code this also applies to a DAP;

Whole Fish are fish as captured, ungutted.

2.2 FRESH, FROZEN AND MINCED FISH

Candling Is passing fillets of fish over a translucent table illuminated from below to detect

parasites and other defects

Dehydration Is the loss of moisture from frozen products through evaporation. This may occur if the

products are not properly glazed, packaged or stored. Deep dehydration adversely affects the appearance and surface texture of the product and is commonly known as "freezer burn";

Fillet Is a slice of fish of irregular size and shape removed from the carcase by cuts made

parallel to the backbone;

Freezer Is equipment designed for freezing fish and other food products, by quickly lowering the

temperature so that after thermal stabilisation the temperature in the thermal centre of the

product is the same as the storage temperature;

Freezing Process Is a process which is carried out in appropriate equipment in such a way that the range of

> temperature of maximum crystallisation is passed quickly. The quick freezing process shall not be regarded as complete unless and until the product temperature has reached -18°C (0°F)

or lower at the thermal centre after thermal stabilisation;

Fresh Fish are fish or fishery products which have received no preserving treatment other than

chilling;

Frozen Fish re fish which have been subjected to a freezing process sufficient to reduce the temperature of

> the whole product to a level low enough to preserve the inherent quality of the fish and which have been maintained at this low temperature, as specified in the Standard for Quick Frozen Finfish, Eviscerated and Uneviscerated during transportation, storage and distribution up to and including the time of final sale. For the purpose of this Code the terms "frozen", "deep

frozen", "quick frozen", unless otherwise stated, shall be regarded as synonymous;

Glazing The application of a protective layer of ice formed at the surface of a frozen product by

spraying it with, or dipping it into, clean sea water, potable water, or potable water with

approved additives, as appropriate;

Minced Fish Is comminuted flesh produced by separation from skin and bones;

Modified means packaging in which the atmosphere surrounding the fish is different from the

normal composition of air; **Atmosphere**

Packaging (MAP)

Separation Is a mechanical process for producing minced fish whereby the skin and bone is

substantially removed from the flesh;

Is a mechanical device used for separation; **Separator**

Steak Is a section of fish, removed by cutting approximately at right angle to the backbone.

2.9 **CANNED FISH**

For the purpose of this Code, only the definitions of the main terms related to canning industry and used in section 13 are given. For an overall set of definitions; please refer to the Recommended International Code of Hygienic Practice for Low-Acid and Acidified Low-Acid Canned Food (CAC/PRC 23-1979, Rev. 2 (1993)).

Canned Food means commercially sterile food in hermetically sealed containers;

Commercial sterility of thermally processed food

means the condition achieved by application of heat, sufficient, alone or in combination with other appropriate treatments, to render the food free from micro-organisms capable of growing in the food at normal non-refrigerated conditions at which the food is likely to be held during distribution and

storage;

Hermetically **Containers**

Sealed are containers which are sealed to protect the content against the entry of micro-organisms during and after heat processing;

Retort means a pressure vessel designed for thermal processing of food packed in

hermetically sealed containers;

Scheduled Process (or Sterilisation schedule)

means the thermal process chosen by the processor for a given product and

container size to achieve at least commercial sterility;

Sterilisation means the temperature maintained throughout the thermal process as specified

Temperature in the scheduled process;

Sterilisation time means the time between the moment sterilisation temperature is achieved and

the moment cooling started;

Thermal Process means the heat treatment to achieve commercial sterility and is quantified in

terms of time and temperature;

Venting means thorough removal of the air from steam retorts by steam prior to a

scheduled process;

SECTION 3 PRE-REQUISITE PROGRAMME

Prior to the application of HACCP to any segment of the fish processing chain, that segment must be supported by pre-requisite programmes based on good hygienic practice or as required by the competent authority.

The establishment of pre-requisite programmes will allow the HACCP team to focus on the HACCP application to food safety hazards which are directly applicable to the product and the process selected, without undue consideration and repetition of hazards from the surrounding environment. The pre-requisite programmes would be specific within an individual establishment or for an individual vessel and will require monitoring and evaluation to ensure their continued effectiveness.

Reference should be made to the *International Recommended Code of Practice – General Principles of Food Hygiene* (CAC/RCP 1-1969, Rev.3 1997), Annex: *HACCP System and Guidelines for its Application* for further information to assist with the design of the pre-requisite programmes for a processing facility or vessel.

It should be noted that some of the issues listed below, e.g. those related to damage, are designed to maintain quality rather than food safety and are not always essential to a pre-requisite programme for a food safety oriented HACCP system.

HACCP principles can also be applied to defect action points.

3.1 FISHING AND HARVESTING VESSEL DESIGN AND CONSTRUCTION

There are many different types of fishing vessel used throughout the world which have evolved in particular regions to take account of the prevailing economics, environment and types of fish caught or harvested. This Section attempts to highlight the basic requirements for cleanability, minimising damage, contamination and decomposition to which all vessels should have regard to the extent possible in order to ensure hygienic, high quality handling of fresh fish intended for further processing and freezing.

The design and construction of a fishing vessel and vessels used to harvest farmed fish should take into consideration the following:

3.1.1 For Ease of Cleaning and Disinfection

- vessels should be designed and constructed to minimise sharp inside corners and projections to avoid dirt traps;
- construction should facilitate ample drainage;
- a good supply of clean water or potable water² at adequate pressure.

WHO Guidelines for Drinking Water Quality, 2nd edition, Geneva, 1993

3.1.2 To Minimise Contamination

- all surfaces in fish handling areas should be non-toxic, smooth impervious and in sound condition, to minimise the build-up of fish slime, blood, scales and guts and to reduce the risk of physical contamination;
- where appropriate, adequate facilities should be provided for the handling and washing of fish and should have an adequate supply of cold potable water or clean water for that purpose;
- adequate facilities should be provided for washing and disinfecting equipment, where appropriate;
- the intake for clean water should be located to avoid contamination;
- all plumbing and waste lines should be capable of coping with peak demand;
- non-potable water lines should be clearly identified and separated from potable water to avoid contamination;
- objectionable substances, which could include bilge water, smoke, fuel oil, grease, drainage and other solid or semi-solid wastes should not contaminate the fish;
- where appropriate, containers for offal and waste material should be clearly identified, suitably constructed with a fitted lid and made of impervious material;
- separate and adequate facilities should be provided to prevent the contamination of fish and dry materials, such as packaging, by:
 - poisonous or harmful substances;
 - dry storage of materials, packaging etc.;
 - offal and waste materials;
- adequate hand washing and toilet facilities, isolated from the fish handling area, should be available where appropriate;
- prevent the entry of birds, insects, or other pests, animals and vermin, where appropriate.

3.1.3 To Minimise Damage to the Fish

- in fish handling areas, surfaces should have a minimum of sharp corners and projections;
- in boxing and shelving fish storage areas, the design should preclude excessive pressure being exerted on the fish;
- chutes and conveyors should be designed to prevent physical damage caused by long drops or crushing;
- the fishing gear and its usage should minimise damage and deterioration to the fish.

3.1.4 To Minimise Damage during Harvesting of Aquacultured Fish

When aquacultured fish are harvested using seines or nets and are transported live to processing facilities:

- seines, nets and traps should be carefully selected to ensure minimum damage during harvesting;
- harvesting areas and all equipment for harvesting, catching, sorting, grading, conveying and transporting of live fish should be designed for rapid and efficient handling of live fish without causing mechanical damage;
- conveying equipment for live and slaughtered fish should be constructed of suitable corrosion-resistant material which does not transmit toxic substances and should not cause mechanical injuries to fish;
- where fish are transported live, care should be taken to avoid overcrowding and to minimise bruising.
- Ice should be provided for handling of harvested fish not allived.

3.2 PROCESSING FACILITY DESIGN AND CONSTRUCTION

The processing facility should include a product flow-through pattern that is designed to prevent potential sources of contamination, minimise process delays which could result in further reduction in fish quality, and prevent cross-contamination of finished product from raw materials. Fish are highly perishable foods and should be handled carefully and chilled without undue delay. The fish processing facility, therefore, should be designed for the rapid processing and storage of fish and fishery products.

The design and construction of a fish processing facility should take into consideration the following:

3.2.1 For Ease of Cleaning and Disinfection

- the surfaces of walls, partitions and floors should be made of impervious, non-toxic materials;
- all surfaces with which fish might come in contact should be of corrosion resistant, impervious material which is light-coloured, smooth and easily cleanable;
- walls and partitions should have a smooth surface up to a height appropriate to the operation;
- floors should be constructed to allow adequate drainage;
- ceilings and overhead fixtures should be constructed and finished to minimise the build-up of dirt and condensation, and the shedding of particles;
- windows should be constructed to minimise the build-up of dirt and, where necessary, be fitted with removable and cleanable insect-proof screens. Where necessary, windows should be fixed;
- doors should have smooth, non-absorbent surfaces;
- joints between floors and walls should be constructed for ease of cleaning (round joints).

3.2.2 To Minimise Contamination

- processing facility layout should be designed to minimise cross-contamination and may be accomplished by physical or time separation;
- all surfaces in fish handling areas should be non-toxic, smooth impervious and in sound condition, to minimise the build-up of fish slime, blood, scales and guts and to reduce the risk of physical contamination;
- working surfaces that come into direct contact with fish should be in sound condition, durable and easy to maintain. They should be made of smooth, non-absorbent and non-toxic materials, and inert to fish, detergents and disinfectants under normal operating conditions;
- adequate facilities should be provided for the handling and washing of fish and should have an adequate supply of cold potable water for that purpose;
- suitable and adequate facilities should be provided for storage and/or production of ice;
- ceiling lights should be covered or otherwise suitably protected to prevent contamination by glass or otherwise suitably protected to prevent contamination by glass or other material;
- ventilation should be sufficient to remove excess steam, smoke and objectionable odours and cross contamination through aerosols should be avoided;
- adequate facilities should be provided for washing and disinfecting equipment, where appropriate;
- non-potable water lines should be clearly identified and separated from potable water to avoid contamination;
- all plumbing and waste lines should be capable of coping with peak demands;

- accumulation of solid, semi-solid or liquid wastes should be minimised to prevent contamination of fish;
- where appropriate, containers for offal and waste material should be clearly identified, suitably constructed with a fitted lid and made of impervious material;
- separate and adequate facilities should be provided to prevent the contamination of fish by:
 - poisonous or harmful substances;
 - dry storage of materials, packaging etc.;
 - offal and waste materials;
- adequate hand washing and toilet facilities, isolated from the fish handling area, should be available;
- prevent the entry of birds, insects, or other pests and animals;
- water supply lines should be fitted with back flow devices, where appropriate.

3.2.3 To Provide Adequate Lighting

to all work surfaces.

3.3 DESIGN AND CONSTRUCTION OF EQUIPMENT AND UTENSILS

The equipment and utensils used for the handling of fishery products on a vessel or in a processing facility will vary greatly depending on the nature and type of operation involved. During use, they are constantly in contact with the fish. The condition of the equipment and utensils should be such that it minimises the build-up of residues and prevents them becoming a source of contamination.

The design and construction equipment and utensils should take into consideration the following:

3.3.1 For Ease of Cleaning and Disinfection

- equipment should be durable and movable and/or capable of being disassembled to allow for maintenance, cleaning, disinfection and monitoring;
- equipment, containers and utensils coming into contact with fish should be designed to provide for adequate drainage and constructed to ensure that they can be adequately cleaned, disinfected and maintained to avoid contamination;
- equipment and utensils should be designed and constructed to minimise sharp inside corners and projections and tiny crevices or gaps to avoid dirt traps;
- a suitable and adequate supply of cleaning utensils and cleaning agents, approved by the official agency having jurisdiction, should be provided.

3.3.2 To Minimise Contamination

- all surfaces of equipment in fish handling areas should be non-toxic, smooth, impervious and in sound condition, to minimise the build-up of fish slime, blood, scales and guts and to reduce the risk of physical contamination;
- accumulation of solid, semi-solid or liquid wastes should be minimised to prevent contamination of fish;
- adequate drainage should be provided in storage containers and equipment;
- drainage should not be permitted to contaminate fish.

3.3.3 To Minimise Damage

surfaces should have a minimum of sharp corners and projections;

- chutes and conveyors should be designed to prevent physical damage caused by long drops or crushing;
- fish storage equipment should be fit for the purpose and not lead to crushing of the fish.

3.4 HYGIENE CONTROL PROGRAMME

The potential effects of harvesting and handling of products, on-board vessel handling or in-plant production activities on the safety and suitability of fish should be considered at all times. In particular this includes all points where contamination may exist and taking specific measures to ensure the production of a safe and wholesome product. The type of control and supervision needed will depend on the size of the operation and the nature of its activities.

Schedules should be implemented to:

- prevent the build up of waste and debris;
- protect the fish from contamination;
- dispose of any rejected material in a hygienic manner;
- monitor personal hygiene and health standards;
- monitor the pest control programme;
- monitor cleaning and disinfecting programmes;
- monitor the quality and safety of water and ice supplies.

The hygiene control programme should take into consideration the following:

3.4.1 A Permanent Cleaning and Disinfection Schedule

A permanent cleaning and disinfection schedule should be drawn up to ensure that all parts of the vessel, processing facility and equipment therein are cleaned appropriately and regularly. The schedule should be reassessed whenever changes occur to the vessel, processing facility and/or equipment. Part of this schedule should include a 'clean as you go' policy.

A typical cleaning and disinfecting process may involve as many as seven separate steps:

Pre-cleaning	Preparation of area and equipment for cleaning. Involves steps such as removal of all fish or fish products from area, protection of sensitive components and packaging materials from water, removal by hand or squeegee of fish scraps, etc.
Pre-rinse	A rinsing with water to remove remaining large pieces of loose soil.
Cleaning	Treatment of surfaces with an appropriate detergent to loosen and remove any remaining soil.
Rinse	A rinsing with potable water or clean water, as appropriate, to remove all soil and detergent residues.
Disinfection	Application of chemicals, approved by the official agency having jurisdiction and/or heat to destroy most microorganisms on surface
Post-rinse	As appropriate a final rinse with potable water or clean water to remove all disinfectant residues
Storage	Cleaned and disinfected equipment, container and utensils should be stored in a fashion which would prevent its contamination
Check of the eficiency of the cleaning	The efficiency of the cleaning should be controlled as appropriate

Fish handlers or cleaning personnel as appropriate should be well trained in the use of special cleaning tools and chemicals, methods of dismantling equipment for cleaning and should be knowledgeable in the significance of contamination and the hazards involved.

3.4.2 Designation of Personnel for Cleaning

In each processing plant or vessel a trained individual should be designated to be responsible
for the sanitation of the processing facility or vessel and the equipment within.

3.4.3 Maintenance of Premises, Equipment and Utensils

- buildings, materials, utensils and all equipment in the establishment including drainage systems should be maintained in a good state and order;
- equipment, utensils and other physical facilities of the plant or vessel should be kept clean and in good repair;
- procedures for the maintenance, repair, adjustment and calibration, as appropriate, of apparatus should be established. These procedures should specify for each equipment, the methods used, the persons in charge of their application, and their frequency.

3.4.4 Pest Control Systems

- good hygienic practices should be employed to avoid creating an environment conducive to pests;
- pest control programmes could include preventing access, eliminating harbourage and infestations, and establishing monitoring detection and eradication systems;
- physical, chemical and biological agents should be properly applied by appropriately qualified personnel.

3.4.5 Supply of Water, Ice and Steam

3.4.5.1 Water

- an ample supply of cold and hot potable water³ and/or clean water under adequate pressure should be provided where appropriate;
- potable water should be used wherever necessary to avoid contamination to the fish and shellfish.

3.4.5.2 Ice

- ice should be manufactured using potable water or clean water;
- ice should be protected from contamination.

3.4.5.3 Steam

- for operations which require steam, an adequate supply at sufficient pressure should be maintained;
- steam used in direct contact with fish or shellfish or food contact surfaces should not constitute a threat to the safety or suitability of the food.

3.4.6 Waste Management

- offal and other waste materials should be removed from the premises of a processing facility or vessel on a regular basis;
- facilities for the containment of offal and waste material should be properly maintained;
- vessel waste discharge should not contaminate vessel water intake system or incoming product.

WHO Guidelines for Drinking Water Quality, 2 nd edition, Geneva, 1993

3.5 PERSONAL HYGIENE AND HEALTH

Personal hygiene and facilities should be such to ensure that an appropriate degree of personal hygiene can be maintained to avoid contaminating fish

3.5.1 Facilities and Equipment:

Facilities and equipment should include:

- adequate means of hygienically washing and drying hands;
- adequate toilet and changing facilities for personnel should be suitably located and designated.

3.5.2 Personnel Hygiene

- no person who is known to be suffering from, or who is a carrier of any communicable disease or
 has an infected wound or open lesion should be engaged in the preparation, handling or
 transporting of fish or fishery products;
- where necessary, adequate and appropriate protective clothing, headcovering and footwear should be worn;
- all persons working in a processing facility should maintain a high degree of personal cleanliness and should take all necessary precautions to prevent the contamination of the fish or their products or ingredients;
- 0 hand-washing should be carried out by all personnel working in a processing area:
 - at the start of fish handling activities and upon re-entering a processing area;
 - immediately after using the toilet;
 - the following should not be permitted in fish handling and processing areas:
 - smoking
 - spitting
 - chewing or eating
 - sneezing or coughing over unprotected food
 - the adornment of personal effects such as jewellery, watches, pins or other items that, if dislodged, may pose a threat to the safety and suitability of the fish.

3.6 TRANSPORTATION

Vehicles should be designed and constructed:

- such that walls, floors and roofs, where appropriate, are made of a suitable corrosion-resistant material with smooth non-absorbent surfaces. Floors should be adequately drained;
- where appropriate with chilling equipment to maintain chilled fish during transport to a temperature as close as possible to 0°C or, for frozen fish and fishery products, to maintain a temperature of -18°C or colder;
- to provide the fish with protection against contamination, exposure to extreme temperatures and the drying effects of the sun or wind;
- to permit the free flow of chilled air around the load when fitted with mechanical refrigeration means.

3.7 TRACEABILITY AND RECALL PROCEDURES

Experience has demonstrated that a system for recall of product is a necessary component of a pre-requisite programme because no process is fail-safe. Traceability, which includes lot identification, is essential to an effective recall procedure.

• managers should ensure effective procedures are in place to effect the complete traceability and rapid recall of any lot of fishery product from the market;

- appropriate records of processing, production and distribution should be kept and retained for a period that exceeds the shelf-life of the product;
- each container of fish or fishery product intended for the final consumer or for further processing should be clearly marked to ensure the traceability of the producer and of the lot;
- where there is an immediate health hazard, products produced under similar conditions, and likely
 to present a similar hazard to public health, may be withdrawn. The need for public warnings
 should be considered;
- recalled products should be held under supervision until they are destroyed, used for purposes other than human consumption, or reprocessed in a manner to ensure their safety.

3.8 TRAINING

Fish hygiene training is fundamentally important. All personnel should be aware of their role and responsibility in protecting fish from contamination and deterioration. Fish handlers should have the necessary knowledge and skill to enable them to handle fish hygienically. Those who handle strong cleaning chemicals or other potentially hazardous chemicals should be instructed in safe handling techniques.

Each fish processing facility should ensure that individuals have received adequate and appropriate training in the design and proper application of a HACCP system and process control. Training of personnel in the use of HACCP is fundamental to the successful implementation and delivery of the programme in fish processing establishments. The practical application of such systems will be enhanced when the individual responsible for HACCP has successfully completed a course given by or certified by a competent authority. Managers should also arrange for adequate and periodic training of every employee in the establishment so that they understand the principles involved in HACCP.

SECTION 4 - GENERAL CONSIDERATIONS FOR THE HANDLING OF FRESH FISH AND SHELLFISH

Unless they can be reduced to an acceptable level by normal sorting and/or processing, no fish and shelfish should be accepted if it is known to contain parasites, undesirable microorganisms, pesticides, veterinary drugs or toxic, decomposed or extraneous substances known to be harmfull to human health. When fish and shellfish determined as unfit for human consumption are found they should be removed and stored separately from the catch and either reworked and/or disposed of in a proper manner. Potential hazards, which have been known to be associated with fresh fish and shellfish, are described in Section 4.1. All fish and shellfish deemed fit for human consumption should be handled properly with particular attention being paid to time and temperature control.

4.1 POTENTIAL HAZARDS ASSOCIATED WITH FRESH FISH AND SHELLFISH

4.1.1 Biological Hazards

4.1.1.1 Parasites

The parasites known to cause disease in humans and transmitted by fish or crustaceans are broadly classified as helminths or parasitic worms. These are commonly referred to as Nematodes, Cestodes and Trematodes. Fish can be parasitised by protozoans, but there are no records of fish protozoan disease being transmitted to man. Parasites have complex life cycles, involving one or more intermediate hosts and are generally passed to man through the consumption of raw, minimally processed or inadequately cooked products that contain the parasite infectious stage, causing foodborne disease. Freezing at [-20°C or below for 7 days or -35°C for about 20 hours] for fish intended for raw consumption will kill parasites. Processes such as brining or pickling may reduce the parasite hazard if the products are kept in the brine

for a sufficient time but may not eliminate it. Candling, trimming belly flaps and physically removing the parasite cysts will also reduce the hazards but may not eliminate it.

Nematodes

Many species of nematodes are known to occur worldwide and some species of marine fish act as secondary hosts. Among the nematodes of most concern are *Anisakis* spp., *Capillaria* spp., *Gnathostoma* spp., and *Pseudoteranova* spp., which can be found in the liver, belly cavity and flesh of marine fish. An example of a nematode causing disease in man is *Anisakis simplex*; as the infective stage of the parasite is killed by heating ([60°C] for 1 minute) and by freezing ([-20°C] for 24 hours) in the fish core.

Cestodes

Cestodes are tapeworms and the species of most concern associated with the consumption of fish is *Diphyllobotrium latum*. This parasite occurs worldwide and marine fish are intermediate hosts. Similar to other parasitic infections, the foodborne disease occurs through the consumption of raw or under-processed fish. Similar freezing and cooking temperatures as applied to nematodes will inactivate the infective stages of this parasite.

Trematodes

Fish-borne trematode (flatworm) infections are major public health problems that occur endemically in about 20 countries around the world. The most important species with respect to the numbers of people infected belong to the genera *Clonorchis* and *Ophisthorchis* (liver flukes), *Paragonimus* (lung flukes), and to a lesser extent *Heterophyes* and *Echinochasmus* (intestinal flukes). The most important definitive host of these trematodes is man or other mammals. Freshwater fish are the second intermediate host in the life cycles of *Clonorchis* and *Ophistorchis*, and freshwater crustaceans in the case of *Paragonimius*. Foodborne infections take place through the consumption of raw, undercooked or otherwise underprocessed products containing the infective stages of these parasites. Freezing fish at -20°C for 7 days or at -35°C for 24 hours will kill the infective stages of these parasites.

4.1.1.2 Bacteria

The level of contamination of fish at the time of capture will depend on the environment and the bacteriological quality of the water in which fish are harvested. Many factors will influence the microflora of finfish, the more important being water temperature, salt content, proximity of harvesting areas to human habitations, quantity and origin of food consumed by fish, and method of harvesting. The edible muscle tissue of finfish is normally sterile at the time of capture and bacteria are usually present on the skin, gills and in the intestinal tract.

There are two broad groups of bacteria of public health importance that may contaminate products at the time of capture - those that are normally or incidentaly present in the aquatic environment, referred to as the indigenous microflora, and those introduced through environmental contamination by domestic and /or industrial wastes. Examples of indigenous bacteria, which may pose a health hazard, are *Aeromonas hydrophyla*, *Clostridium botulinum*, *Vibrio parahaemolyticus*, *Vibrio cholerae*, *Vibrio vulnificus*, and *Listeria monocytogenes*. Non-

indigenous bacteria of public health significance include members of the Enterobacteriaceae, such as *Salmonella* spp., *Shigella* spp., and *Escherichia coli*. Other species that cause foodborne illness and which have been isolated occasionally from fish are *Edwardsiella tarda*, *Pleisomonas shigeloides* and *Yersinia enterocolitica*.

Indigenous pathogenic bacteria, when present on fresh fish, are usually found in fairly low numbers, and where products are adequately cooked prior to consumption, food safety hazards are insignificant. During storage, indigenous spoilage bacteria will outgrow indigenous pathogenic bacteria, thus fish will spoil before becoming toxic and will be rejected by consumers. Hazards from these pathogens can be controlled by heating

seafood sufficiently to kill the bacteria, holding fish at chilled temperatures and avoiding post-process cross-contamination.

Vibrio species are common in coastal and estuarine environments and populations can depend on water depth and tidal levels. They are particularly prevalent in warm tropical waters and can be found in temperate zones during summer months. *Vibrio* species are also natural contaminants of brackish water tropical environments and will be present on farmed fish from these zones. Hazards from *Vibrio* spp. associated with finfish can be controlled by thorough cooking and preventing cross-contamination of cooked products. Health risks can also be reduced by rapidly chilling products after harvest, thus reducing the possibility of proliferation of these organisms.

4.1.1.3 Viral Contamination

Molluscan shellfish harvested from inshore waters that are contaminated by human or animal faeces may harbour viruses that are pathogenic to man. Enteric viruses that have been implicated in seafood-associated illness are the hepatitis A virus, caliciviruses, astroviruses and the Norwalk virus. The latter three are often referred to as small round structured viruses. All of the seafood-borne viruses causing illness are transmitted by the faecal-oral cycle and most viral gastro-enteritis outbreaks have been associated with eating contaminated shellfish, particularly raw oysters.

Generally viruses are species specific and will not grow or multiply in foods or anywhere outside the host cell. There is no reliable marker for indicating presence of the virus in shellfish harvesting waters. Seafood-borne viruses are difficult to detect, requiring relatively sophisticated molecular methods to identify the virus.

Occurence of viral gastro-enteritis can be minimized by controlling sewage contamination of shellfish farming areas and pre-harvest monitoring of shellfish and growing waters as well as controlling other sources of contamination during processing. Depuration or relaying are alternative strategies but longer periods are required for shellfish to purge themselves clean of viral contamination than for bacteria. Thermal processing (85-90°C for 1.5 min.) will destroy viruses in shellfish.

4.1.1.4 Biotoxins

There are a number of important biotoxins to consider. Around 400 poisonous fish species exist and, by definition, the substances responsible for the toxicity of these species are biotoxins. The poison is usually limited to some organs, or is restricted to some periods during the year.

For some fish, the toxins are present in the blood; these are ichtyohaemotoxin. The involved species are eels from the Adriatic, the moray eels, and the lampreys. In other species, the toxins are spread all over the tissues (flesh, viscera, skin); these are ichtyosarcotoxins. It concerns tetrodotoxic species responsible for several poisonings, often lethal.

Biotoxins are often heat-stable and the only possible control measure is to check the identity of the used species.

Scombrotoxin

Scombroid intoxication, sometimes referred to as histamine poisoning, results from eating fish that have been incorrectly chilled after harvesting. Scombrotoxin is attributed mainly to *Enterobacteriaceae* which produce high levels of histamine in the fish muscle when products are not immediately chilled after catching. The main susceptible fish are the scombroids such as tuna, mackerel, and bonito, although it can be found in other species. The intoxication is rarely fatal and symptoms are usually mild. Rapid refrigeration after catching and a high standard of handling during processing should prevent the development of the toxin. The toxin is not inactivated by normal cooking temperatures or by canning. In addition, fish may contain toxic levels of histamine without exhibiting any of the usual sensory parameters characteristic of spoilage.

Phycotoxins

Ciguatoxin

And the other important toxin to consider is ciguatoxin, which can be found in a wide variety of mainly carnivorous fish inhabiting shallow waters in or near tropical and subtropical coral reefs. The source of this toxin is dinoflagellates and over 400 species of tropical fish have been implicated in intoxication. The toxin is known to be heat stable. There is still much to be learnt about this toxin and the only control measure that can reasonably be taken is to avoid marketing fish that have a known consistent record of toxicity.

PSP/DSP/ASP/NSP

These toxins concern especially the bivalve shellfish; the toxicity is due to the ingestion by the shellfish of phytoplanktonic species, which are able to synthesise toxic substances. The shellfish concentrates the toxin to a level such as it becomes potentially toxic. The principal toxins are the Paralytic Shellfish Poison (PSP) produced by dinoflagellates genus *Alexandrium*, the Diarrheic Shellfish Poison (DSP) produced by other dinoflagellates genus *Dinophysis* Amnesic Shelfish Poisoning (ASP) in which Nitzchi spp produce domoic acid or Neurotoxic Shelfish Poisoning (NSP) produced by Gymnodium spp.

All these toxins are known to keep in general their toxicity through processing, even in canned fish products, so the knowledge of the species identity and/or origin of fish or shellfish intended for processing is important.

Tetrodotoxin

Fish mainly belonging to the family Tetradontidea ("puffer fishes") may accumulate this toxin which is resposible for several poisonings, often lethal. The toxin is generally found in the fish liver, roe and guts, and less freequent in the flesh. Diffrently from most other fish biotoxins that accumulate in the live fishor shelfish, algae do not produce this toxin. The mechanism of toxin production is still not clear, however, apparently there are often indications of the involvement of symbiotic bacteria.

4.1.2 Chemical hazards

Fish may be harvested from coastal zones and inland habitats that are exposed to varying amounts of environmental contaminants. Of greatest concern are fish harvested from coastal and estuarine areas rather than fish harvested from the open seas. Chemicals, organochloric compounds and heavy metals may accumulate in products that can cause public health problems. Veterinary drug residues can occur in aquaculture products when correct withdrawal times are not followed or when the sale and use of these compounds are not controlled. Fish can also be contaminated with chemicals such as diesel oil, when incorrectly handled and detergents or disinfectants when not properly rinsed out.

4.1.3 Physical Hazards

These can include material such as metal or glass fragments, shell, bones, etc.

4.2 TIME AND TEMPERATURE CONTROL

Temperature is the single most important factor affecting the rate of fish deterioration and multiplication of micro-organisms. For species prone to scombroid toxin production, time and temperature control may be the most effective method in controlling food safety. It is therefore essential that fresh fish, fillets and other and shellfish and their products which are to be chilled should be held at a temperature as close as possible to 0°C.

4.2.1 Minimise the Deterioration of Fish - Time

To minimise the deterioration of fish, it is important that:

- chilling of fish should commence as soon as possible;
- fresh fish should be kept chilled, processed and distributed with care and minimum delay.

4.2.2 Minimise the Deterioration of Fish - Temperature Control

Where temperature control is concerned:

- sufficient and adequate icing, or chilled or refrigerated water systems where appropriate, should be employed to ensure that fish are kept chilled at a temperature as close as possible to 0°C;
- fish should be stored in shallow layers and surrounded by finely divided melting ice;
- chilled or refrigerated water systems and/or cold storage systems should be designed and maintained to provide adequate cooling and/or freezing capacities during peak loads;
- fish should not be stored in refrigerated water systems to a density which impairs its working efficiency;
- monitoring and controlling the time and temperature and homogeneity of chilling should be performed regularly.

4.3 MINIMISE THE DETERIORATION OF FISH - HANDLING

Poor handling practices can lead to damage of fresh fish which can accelerate the rate of decomposition and increase unnecessary post-harvest losses. Handling damage can be minimised by:

- fish should be handled and conveyed with care particularly during transfer and sorting in order to avoid physical damage such as puncture, mutilation, etc.;
- where fish are held or transported live, care should be taken to maintain factors that can influence fish health (e.g. CO₂, O₂, temperature, nitrogenous wastes, etc.);
- fish should not be trampled or stood upon;
- where boxes are used for storage of fish they should not be overfilled or stacked too deeply;
- while fish are on deck, exposure to the adverse effects of the elements should be kept to a minimum in order to prevent unnecessary dehydration;
- finely divided ice should be used where possible, which can help minimise damage to fish and maximise cooling capacity;
- in refrigerated water storage areas, the density of the fish should be controlled to prevent damage.

SECTION 5 HAZARD ANALYSIS CRITICAL CONTROL POINT (HACCP) AND DEFECT ACTION POINT (DAP) ANALYSIS

The Hazard Analysis Critical Control Point (HACCP) is a science-based system which is aimed to prevent food safety problems from occurring rather than reacting to non-compliance of the finished product. The HACCP system accomplishes this by the identification of specific hazards and the implementation of control measures. An effective HACCP system should reduce the reliance on traditional end-product testing. Section 5 explains the principles of HACCP as it applies to the handling and processing of fish and fishery products, but the Code can only provide guidance on how to use these principles and offer suggestions as to the type of hazards which may occur in the various fish and fishery products. The HACCP plan, which should be incorporated into the food management plan should be well documented and be as simple as possible. This section will demonstrate one format, which may be considered in the development of the HACCP plan.

Section 5 also explains how a similar approach involving many of the principles can apply to the broader application covering the essential quality, composition and labelling provisions of Codex standards or other non-safety requirements which in this case are referred to as **Defect Action Point Analysis**. This approach for defect analysis is optional and other techniques, which achieve the same objective, may be considered.

Figure 5.1 summarises how to develop a HACCP and Defect Analysis system.

5.1 HACCP PRINCIPLES

The HACCP system consists of seven principles⁴, which are:

PRINCIPLE 1

Conduct a hazard analysis.

PRINCIPLE 2

Determine the Critical Control Points (CCPs).

PRINCIPLE 3

Establish critical limit(s).

PRINCIPLE 4

Establish a system to monitor control of the CCP.

PRINCIPLE 5

Establish the corrective action to be taken when monitoring indicates that a particular CCP is not under control.

PRINCIPLE 6

Establish procedures for verification to confirm that the HACCP system is working effectively.

PRINCIPLE 7

Establish documentation concerning all procedures and records appropriate to these principles and their application.

These principles have to be followed in any consideration of HACCP.

HACCP is an important management tool, which can be used by operators for ensuring safe, efficient processing. It must also be recognised that personnel training is essential in order that HACCP will be effective. In following HACCP principles, users are requested to list all of the hazards that may be reasonably expected to occur for each product type at each step or procedure in the process from point of harvest, during unloading, transport, storage or during processing, as appropriate to the process defined. It is important that HACCP principles be considered on a specific basis to reflect the risks of the operation.

5.2 DEFECT ACTION POINT ANALYSIS

Since the Code is intended to cover not only those hazards associated with safety but to include other aspects of production including the essential product quality, composition and labelling provisions as described in product standards developed by the Codex Alimentarius Commission, not only are critical control points (CCP) described but also defect action points (DAP) are included in the Code. The HACCP principles may be applied to the determination of a DAP, with quality instead of safety parameters being considered at the various steps.

5.3 APPLICATION

International Recommended Code of Practice – General Principles of Food Hygiene (CAC/RCP 1-1969, Rev.3 - 1997), Annex: HACCP System and Guidelines for its Application.

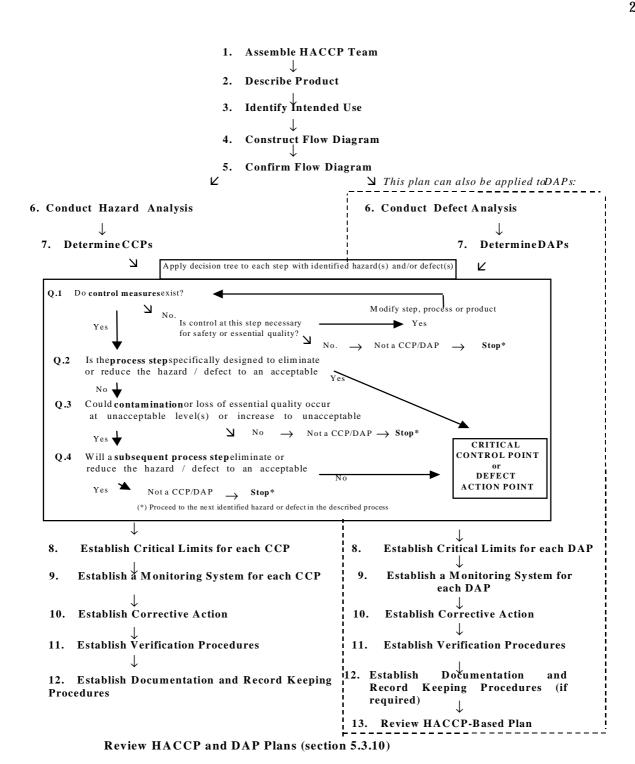


Figure 5.1 Summary of how to implement a HACCP and Defect Analysis

Each aquaculture, molluscan shellfish, and fish processing facility should ensure that the provisions of the appropriate Codex standards are met. To accomplish this, each facility should implement a food safety management system based on HACCP principles and should at least consider a similar approach to defects, both of which are described in this code. Prior to the application of HACCP to any segment of the growing, handling and processing chain of fish and fishery products, that segment must be supported by a prerequisite programme based on good hygienic practice (see Section 3). It should be noted that parts of the pre-requisite programme may be classified as a CCP or DAP within a particular process.

The food management system developed should indicate responsibility, authority and the interrelationships of all personnel who manage, perform and verify work affecting the performance of such systems. It is important that the collection, collation and evaluation of scientific and technical data should be carried out by a multi-disciplinary team. Ideally, a team should consist of people with the appropriate level of expertise together with those having a detailed knowledge of the process and product under review. Examples of the type of personnel to include on the team are the processing facility manager, a microbiologist, a quality assurance/quality control specialist, and others such as buyers, operators, etc., as necessary. For small-scale operations, it may not be possible to establish such a team and therefore external advice should be sought.

The scope of the HACCP plan should be identified and should describe which segments of the food chain is involved and the general classes of hazards to be addressed.

The design of this programme should identify critical control points in the operation where the processing facility or product will be controled, the specification or standard to be met, the monitoring frequency and sampling plan used at the critical control point, the monitoring system used to record the results of these inspections and any corrective action when required. A record for each critical control point that demonstrates that the monitoring procedures and corrective actions are being followed should be provided. The records should be maintained as verification and evidence of the plant's quality assurance programme. Similar records and procedures may be applied to DAPs with the necessary degree of record keeping. A method to identify, describe, and locate the records associated with HACCP programmes should be established as part of the HACCP programme.

Verification activities include the application of methods; procedures (review/audit) and tests in addition to those used in monitoring to determine:

- the effectiveness of the HACCP or DAP plan in delivering expected outcomes i.e. validation;
- compliance with the HACCP or DAP plan, e.g. audit/review;
- whether the HACCP or DAP plan or its method of application need modification or revalidation."

Table 5.1 A product description for Canned Tuna in Salted Water

	Objective	Example
Product name(s)	Identify the species and method of processing.	Canned tuna in salted water
Source of raw material	Describe the origin of the fish	Yellowfin tuna caught by purse seine in the Gulf of Guinea Whole brine frozen
Important final product characteristics	List characteristics that affect product safety and essential quality, especially those that influence microbial flora.	Compliance with Codex Standard Canned Tuna and Bonito; 'low-acid' food; can seal integrity.
Ingredients	List every substance added during processing. Only ingredients approved by the official agency having jurisdiction may be used.	water, salt
Packaging	List all packaging materials. Only materials approved by the official agency having jurisdiction may be used.	Container in coated chromium steel, capacity: 212 ml, total net weight: 185 g, fish weight: 150 g Traditional opening
How the end product is to be used	State how the final product is to be prepared for serving, especially whether it is ready to eat.	Ready to eat
Shelf life (if applicable)	State the date when the product can be expected to begin to deteriorate if stored according to instructions.	3 years
Where the product will be sold	Indicate the intended market. This information will facilitate compliance with target market regulations and standards.	Domestic retail market.
Special labelling instructions	List all instructions for safe storage and preparation	"Best before the date shown on label."
Special distribution control	List all instructions for safe product distribution.	None

The implementation of HACCP principles is better identified in the Logic Sequence for implementation of HACCP (Figure 5.1).

This flow chart is for illustrative purposes only. For in-factory HACCP implementation a complete and comprehensive flow chart has to be drawn up for each process.

References correspond to relevant Sections of the Code.

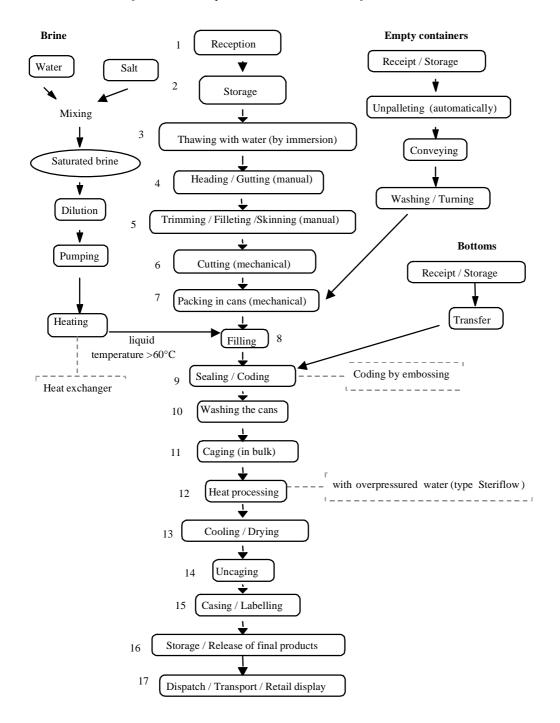


Figure 5.2 Example of a flow diagram for a canned tuna fish processing line

5.3.1 Describe Product

In order to gain a greater understanding and knowledge of the product under review, a thorough product description evaluation should be carried out. This exercise will facilitate in the identification of potential hazards or defects. An example of the type of information used in describing a product is given in Table 5.1.

5.3.2 Flow Diagram

For Hazard and Defect Analysis, it is necessary to carefully examine both the product and the process and produce a flow diagram(s). Any flow diagram should be as simple as possible. Each step in the process, including process delays from the selection of raw materials through to the processing, distribution, sale and customer handling, should be clearly outlined in sequence with sufficient technical data to avoid ambiguity. If a process is too complex to be easily represented by a single flow diagram, then it can be sub-divided into constituent parts, provided the relationship between each of the parts is clearly defined. It is helpful to number and label each processing step for ease of reference. An accurate and properly constructed flow diagram will provide the multi-disciplinary team with a clear vision of the process sequence. Once CCPs and DAPs have been identified they can be incorporated into the flow diagram specific for each processing facility. Figure 5.2 represents an example of a flow diagram for a canned tuna fish processing line. For examples of different processes see Figures 6.1 to 15.1 in the individual processing sections of the code.

5.3.3 Conduct Hazard and Defect Analysis

The purposes of hazard analysis are to identify all such food safety hazards at each Step, to determine their significance and to assess whether control measures for those hazards are available at each Step. Defect analysis serves the same purpose for potential quality defects.

5.3.3.1 Identification of Hazards and Defects

It cannot be stressed enough that where practical and feasible each individual facility should gather sound scientific and technical data relevant to the businesses for each step, from primary production, processing, manufacture, storage and distribution until the point of consumption. The assembly and nature of this information should be such to ensure that the multi-disciplinary team is able to identify and list, at each step of the process, all relevant potential hazards and defects that, in the absence of control measure(s), may likely result in the production of an unacceptable food. Table 5.2 summarises possible pre-harvest and harvest safety hazards in incoming fish & molluscan shellfish and Table 5.3 summarises possible safety hazards introduced in the post harvest and further processing of fish & molluscan shellfish.

It is important to identify potential hazards and defects in the operation from the point of view of plant construction, equipment used in the plant and hygienic practices, including those which may be associated with the use of ice and water. This is covered by the pre-requisite programme and is used to denote hazards that are common to almost any point in the process.

Table 5.2 Examples of Pre-harvest and Harvest Hazards in Incoming Fish & Shellfish

Biological		Chemical		Physical	
Parasites:	Parasites of public health significance: Trematodes, Nematodes, Cestodes	Chemicals:	Pesticides, herbicides, algicides, fungicides, anti-oxidants (added in feeds);		sh ooks
Pathogenic bacteria:	Salmonella, Shigella, E.coli, Vibrio cholerae, Vibrio parahaemolyticus, Vibrio vulnificus,	Veterinary drug residues:	Antibiotics, growth promoters (hormones), other feed additives		
Enteric Viruses:	Norwalk virus	Heavy metals:	Metals leached from marine sediments and soil, from industrial wastes, from sewage or animal manures		
Biological toxins:	Tetrodotoxin; Ciguatoxin, [Paralytic Shellfish Poisoning (PSP), Diarrheic Shellfish Poisoning (DSP), Neurotoxic Shellfish Poisoning (NSP), Amnesic Shellfish Poisoning (ASP)]				
		Miscellaneous:	Petroleum		

Table 5.3 Examples of Hazards Introduced in the Post Harvest and Further Processing of Fish & Shellfish*.

Biological		Chemical		Physical	
Pathogenic bacteria:	Listeria monocytogenes, Clostridium botulinum, Staphylococcus aureus	Chemicals:	Disinfectants, Sanitizers Lubricants (Misapplication)	or Foreign Matter	Metal fragments; hard or sharp objects
Enteric Viruses:	Hepatitis A, Rotovirus		Disinfectants, Sanitizers Lubricants (no approved)	or on-	
Biological toxins:	Scombrotoxin, Staph. Enterotoxin, botulinum toxin				
		Ingredients and Additives:	Misapplication a non-approved	and	

Note: For biological hazards, environmental factors (for example: temperature, oxygen availability, pH and A_w) play a major role in their activity and growth, therefore the type of processing the fish will undergo, and its subsequent storage, will determine their risk to human health and inclusion in a food safety management plan. In addition, some hazards may show a certain degree of overlap between the two levels of operation through their existence and manifestation into the water supply.

For the example on canned tuna developed in this section, the following essential potential hazards can be identified:

Table 5.4: An example of potential hazards for canned tuna

	In raw materials (frozen	During processing or storage or transportation		
	tuna)			
Biological	Presence of Cl. botulinum,	Contamination by Cl. Botulinum, Growth of Cl. Botulinum,		
	Presence of histamine	Survival of spores of Cl. Botulinum, Contamination and		
		growth of Staphylococcus aureus		
		Microbial recontamination after heat processing		
		Production of histamine during processing, Production of		
		staphylotoxin		
Chemical	Presence of heavy metals	Recontamination by metals coming from the cans		
		Recontamination by cleaning agents, by the brine, by		
		mechanical grease,		
Physical	Presence of foreign material	Recontamination during processing (pieces of knives, by the		
		cans,)		

For the example on canned tuna developed in this section, the following potential defects can be identified:

^{*} For hazards relating to specific products see the relevant processing section.

Table 5.5 An example of potential defects of canned tuna

	In raw materials (frozen	During processing or storage or transportation of cans			
	tuna)				
Biological	Decomposition	Decomposition, survival of micro-organisms responsible of			
		decomposition,			
Chemical		oxidation during storage,			
Physical		Objectionable matters (viscera, scales, skin,), formation of			
		struvite crystals, container defects (panelled container,)			
Others	species substitution	abnormal flavours, incorrect weight, incorrect coding,			
		incorrect labelling			

5.3.3.1.1 Hazards

It is equally important to consider, naturally occurring food safety hazards in the environment from which fish are harvested. In general, risks to consumer health from seafoods captured in unpolluted marine environments are low, provided these products are handled in line with principles of Good Manufacturing Practice. However, as with all foods, there are some health risks associated with the consumption of certain products, which may be increased when the catch is mishandled after harvest. Fish from some marine environments, such as tropical reef fish, can pose a consumer risk from natural marine toxins, such as ciguatera. The risk of adverse health effects from certain hazards might be increased under certain circumstances in products from aquaculture when compared with fish from the marine environment. The risks of foodborne disease associated with products from aquaculture are related to inland and coastal ecosystems, where the potential of environmental contamination is greater when compared to capture fisheries. In some parts of the world, where fish are consumed either raw or partially cooked, there is an increased risk of foodborne parasitic or bacterial disease. In order to perform a hazard analysis as part of the process of developing a HACCP plan, fish processors must have scientific information on potential hazards associated with raw material and products for further processing.

5.3.3.1.2 Defects

Potential defects are outlined in the essential quality, labelling and composition requirements described in the Codex Standards listed in Appendix XII. Where no Codex Standard exists regard should be made to national regulations and/or commercial specifications.

End product specifications outlined in Appendices II - XI, describe optional requirements which are intended to assist buyers and sellers in describing those provisions which are often used in commercial transactions or in designing specifications for final products. These requirements are intended for voluntary application by commercial partners and not for application by governments.

5.3.3.2 Significance of Hazards and Defects

One of the most important activities, which must be performed in a processing facility as part of the food safety management system is to determine if an identified hazard or defect is significant. The two primary factors that determine whether a hazard or defect is significant for HACCP purposes are probability of occurence of an adverse health effect and the severity of the effect. A hazard that has a high severity of effect, such as death from *Clostridium botulinum* toxin, may impose a societallu unacceptable risk at very low probability of occurence, and thus warrant the application of HACCP controls (i.e., be a significant hazard for purposes of HACCP). Thus, in the processed canned tuna, *Clostridium botulinum* should be considered a significant hazard to be controlled through the application of a validated thermal process schedulle. On the other hand, a hazard with a relatively low severity, such as mild gastroenteritis, might not warrant the HACCP controls at the same very low probaility of occurence, and thus not be significant for purposes of HACCP.

Information gathered during the product description exercise (refer to Section 5.3.1 – Describe Product) could also help facilitate the detrmination of significance since the likelihood of occurence of hazard or defect can be affected by factors such as how the consumer will likely use the product (e.g., to consumed or cooked raw);

the types of consumers who will likely consume it (e.g., immuno-compromised, eldery, chlidren, etc.) and the method of storage and distribution (e.g., refrigeratyed or frozen).

Once significant hazard and defects have been identified, consideration needs to be given to assess their potential to be introduced or controlled at each step of the process. The use of a flow diagram (refer to Section 5.3.2 – Flaw Diagram) is beneficial for this purpose. Control measures must be considered for significant hazard(s) or defect(s) associated with each step with the aim of eliminating its possible occurence or to reduce it to an acceptable level. A hazard or defet may be controlled by more that one control measure. For illustrative purposes, tables 5.6 and 5.7 demonstrate an approach to listing significant hazards and defets and the related control measures for the processing step, "Heat Processing".

Table 5.6 An example of the significant hazard survival of Cl. Butulinum at the step of heat processing for canned tuna

Processing step	Potential hazard	Is the potential	Justification	Control measures
		hazard significant?		
12. Heat	Cl. botulinum	Yes	A non-efficient heat	Ensure adequate
processing	viable spores		processing may result in	heat applied for
			survival of C. botulinum	proper time at retort
			spores and therefore,	
			possibility of toxin	
			production. A product	
			must be commercially	
			sterile	

Table 5.7: An example of the significant defect rancidity during the storage of frozen tuna for canned tuna

Processing step	Potential defect	Is the potential defect significant?	Justification	Control measures
2. Storage of frozen tuna	Persistent and distinct objectionable odours or flavours indicative of rancidity	Yes		Controlled temperature in the storage premises Stock management procedure Maintenance procedure of the refrigeration system Personnel training and qualification

Table 5.8 A schematic example of a hazard analysis with corresponding control measures and the application of the Codex decision tree for the determination of a critical control point at processing step 12 of the example process as set out in Figure 5.2.

Processing Step N° 12		Application	of Codex Decisi	on Tree		
Heat process	sing					
Potential	Control Measures					
Hazards						
Cl.	Ensure adequate	heat	Q1 : Do	Q2: Is the step	Q3: Could	Q4 : Will a
botulinum	applied for proper	time	control	specifically	contamination	subsequent step
viable	at retort		measures	designed to	occur in excess	eliminate or
spores			exist?	eliminate or	of acceptable	reduce the
			70	reduce the	levels or could	hazard to an
			If yes – go to	likely	these increase to	acceptable level?
			Q2.	occurrence of	unacceptable	T6
			If no -	Cl. botulinum	levels?	If yes – not a CCP.
			If no – consider	to an acceptable	If was go to	CCP.
			whether	level?	If yes – go to Q4.	If no – CCP.
			control	icvci:	Q4.	What about
			measures are	If yes – this	If no – not a	consideration of
			available or	step is a CCP.	CCP.	a previous step?
			necessary	stop is a c ci .		or provious step.
			within the	If no – go to		
			process.	Q3.		
			Proceed to			
			next			
			identified			
			hazard.			
			A: Yes: a	A : Yes, this		
			heat .	step was		
			processing	specifically		
			procedure	designed to		
			(schedule,	eliminate		
			method) is clearly	spores.		
			defined.			
				essing sten N°12 /	Heat processing »	
			is a Critical Co		· Heat processing »	

5.3.4 Determine Critical Control Points and Defect Action Points

A thorough and concise determination of Critical Control Points and Defect Action Points in a process is important in ensuring food safety and compliance with elements related to essential quality, composition and labelling provisions of the appropriate Codex standard. The Codex decision tree (Figure 5.1, step 7) is a tool, which can be applied, to the determination of CCPs and a similar approach may be used for DAPs. Using this decision tree, a significant hazard or defect at a step can be assessed through a logical sequence of questions. Where CCPs and DAPs have been identified at a step, that point in the process must be controlled to prevent, reduce or eliminate the likely occurrence of the hazard or defect to an acceptable level. For illustrative purposes, an example of the application of the Codex decision tree to a hazard and defect using the canned tuna fish processing line, are shown in Tables 5.4 & 5.5, respectively.

Table 5.9 A schematic example of a defect analysis with corresponding control measures and the application of the Codex decision tree for the determination of a defect action point at processing step 2 of the example process as set out in Figure 5.2.

Processing Steps N °	22	Application	n of Codex Decision	n Tree	
Storage of frozen tu	ına				
Potential Defects	Control Measures				
Persistent and distinct objectionable odours or flavours indicative of rancidity	Controlled temperature in storage premises. Stock managemnt procedure.	Q1: Do control measures exist? If yes – go to Q2. If no – consider whether control measures are available or necessary within the process. Proceed to next identified hazard.	Q2: Is the step specifically designed to eliminate or reduce the likely occurrence of rancidity to an acceptable level? If yes – this step is a DAP. If no – go to Q3.	Q3: Could rancidity occur in excess of acceptable levels or could it increase to unacceptable levels? If yes – go to Q4. If no – not a DAP.	Q4: Will a subsequent step eliminate rancidity or reduce its likely occurrence to acceptable level? If yes – not a DAP. If no – DAP. What about consideration of a previous step?
		A: Yes, the storage temperature is controlled,	A: No	A: Yes, if the storage time is too long and/or	A : No
		procedures exist		the storage temperature is too high	
		Decision: Processing Step N°2 «Storage of frozen			
		is a Defect Act	•	2 "Storage of	frozen tuna »

5.3.5 Establish Critical Limits

For each CCP and DAP, critical limits for the control of the hazard or defect must be specified. For any given hazard or defect, it may be necessary to have more than one critical limit designated for each control measure. The establishment of critical limits should be based on scientific evidence and validated by appropriate technical experts to ensure its effectiveness in controlling the hazard or defect to the determined level. Table 5.10 illustrates critical limits for a CCP and a DAP using a canned tuna fish processing line as an example.

5.3.6 Establish Monitoring Procedures

Any monitoring system developed by the multi-disciplinary team should be designed to detect loss of control at a CCP or DAP relative to its critical limit. The monitoring activity of a CCP or DAP should be documented in a concise fashion providing details regarding the individual responsible for the observation or measurement,

the methodology used, the parameter(s) being monitored and the frequency of the inspections. The complexity of the monitoring procedure should also be carefully considered. Considerations include optimising the number of individuals performing the measurement and selection of appropriate methods, which will produce rapid results (for example: time, temperature, pH). For CCPs, records of monitoring should be acknowledged and dated by a responsible person for verification.

Because each process is unique for each fish product, it is possible only to present, for illustrative purposes, an example of a monitoring approach for a CCP and DAP using the canned tuna fish processing line. This example is shown in Table 5.10.

5.3.7 Establish Corrective Action

An effective HACCP or DAP plan is anticipatory by nature and it is recognised that corrective action may be necessary from time to time. A documented corrective action program should be established to deal with instances where the critical limit has been exceeded and loss of control has occurred at a CCP or DAP. The goal of this plan is to ensure that comprehensive and specific controls are in place and can be implemented to prevent the affected lot(s) from reaching the consumer. Of equal importance, is an assessment by plant management and other appropriate personnel to determine the underlying reason(s) why control was lost. For the latter, a modification to HACCP and DAP plans may be necessary. A record of investigation results and actions taken should be documented by a responsible person for each instance where loss of control occurred at a CCP or DAP. The record should demonstrate that control of the process has been re-established. An example of a corrective action approach for a CCP and DAP using a canned tuna fish processing line is illustrated in Table 5.10.

5.3.8 Establish Verification Procedures

A processing facility should establish a verification procedure carried out byqualified individuals, to periodically assess if the HACCP and DAP plans are adequate, implemented and working properly. This step will help determine if CCPs and DAPs are under control. Examples of verification activities include: validation of all components of the HACCP plan including: a paper review of HACCP system, its procedures and records; review of corrective actions and product disposition actions when critical limits are not met and validation of established critical limits. The latter is particularly important when an unexplained system failure has occurred, when a significant change to the process, product or packaging is planned or when new hazards or defects have been identified. Observation, measurement and inspection activities within the processing facility should also be incorporated as a part of the verification procedure, where applicable. Verification activities should be carried out by qualified competent individuals. The verification frequency of the HACCP and DAP plans should be sufficient to provide assurance that their design and implementation will prevent food safety problems as well as issues associated with essential quality, composition and labelling provisions of the appropriate Codex standard to enable problems to be detected and dealt with in a timely manner. For illustration purposes, an example of a verification procedure approach for a CCP and DAP using the canned tuna fish processing line is shown in Table 5.10.

5.3.9 Establish Documentation and Record Keeping Procedures

Documentation may include Hazard Analysis, CCP determination and verification procedures.

A current, accurate and concise record keeping system will greatly enhance the effectiveness of a HACCP program and facilitate in the verification process. Examples of the elements of a HACCP plan that should be documented have been provided in this section for illustrative purposes. Inspection and corrective action records should be practical and collect all the appropriate data necessary to demonstrate "real-time" control or deviation control of a CCP. Records are recommended but not required for a DAP except where a loss of control occurred. For illustration purposes, an example of a record keeping approach for a CCP and DAP using the canned tuna fish processing line is shown in Table 5.10.

5.3.10 Review of HACCP and DAP Plans

Upon completion of all the steps for the development of HACCP and DAP plans as outlined in Figure 1 a full review of all components should be conducted. The purpose of these reviews is to verify that the plans are capable of meeting their objectives.

Table 5.10 An example of the results of the application of HACCP principles to the two specific steps in the canned tuna process (Tables 5.8 & 5.9), for a CCP & a DAP, respectively.

ССР	CCP						
Processing Ste	p No. 12: Heat Process	sing					
Hazard: Closti	ridium botulinum viabl	e spores					
Critical Limit	Monitoring Procedure	Corrective Action	Records	Verification			
Those specific parameters associated with heat processing.	Who: Qualified person assigned to heat processing What: All parameters Frequency: every batch How: Checks of sterilisation schedule and other factors	Who: qualified personnel What: Personnel retraining New heat processing or batch destruction Corrective maintenance of equipment Hold product until safety can be evaluated. Who: Appropriate trained personnel	Monitoring records, corrective action records, product evaluation records, validation records, audit records, HACCP plan review record	Validation, finished product evaluation, internal audit, review of records, calibration of machinery (may be a prerequsite), review of HACCP plan, external audit			

DAP				
Processing Step No. 2 : Storage of frozen tuna				
Defect: Persistent and distinct objectionable odours or flavours indicative of rancidity				
Critical Limit:	Monitoring Procedure	Corrective Action	Records	Verification
Number of rancid sample units cannot exceed acceptance number of established sampling plan. Storage temperature and time.	Who: Appropriate trained personnel How: Organoleptic examination Chemical tests Checking of the storage premise temperature Checking of stock forms What: fish quality and acceptability based on product Codex standard. Frequency: as required	What: Application of an intensified monitoring According to the results of this intensified inspection, immediate processing, sorting or reject of frozen tuna exceeding the critical limits. Adjust storage temperature. Personnel retraining Who: Appropriate trained personnel	Analysis results Stock forms Temperature records	On-site audit Review of monitoring and corrective action reports

Conclusion

Section 5 has demonstrated the principles of HACCP and how they should be applied to a process to ensure safe product. The same principles can be used to determine the points in a process where it is necessary to control defects. Since every processing facility and each processing line is different it is possible within this Code only to demonstrate the types of potential hazards and defects that must be considered. Furthermore, because of the nature of the significance of hazards and defects it is not possible to categorically determine which steps in a process will be CCPs and/or DAPs without actually assessing the process, the objectives of the process, its environment and expected outcomes. The example of the canned tuna fish processing line is intended to illustrate how to apply the principles, given the outcome of a commercially sterile product, and why a HACCP and DAP plan will be unique to each operation.

The remaining Sections in the Code concentrate on fish and fishery product processing and attempt to illustrate the potential hazards and defects at the various stages in a wide range of processes. In developing a HACCP or DAP plan it will be necessary to consult Sections 3 & 5 before turning to the appropriate processing section for specific advice. It should also be noted that Section 6 refers to processing of fresh, frozen and minced fish and will provide useful guidance for most fish processing operations.

SECTION 6 PROCESSING OF FRESH, FROZEN AND MINCED FISH

In the context of recognising controls at individual processing steps, this section provides <u>examples</u> of potential <u>hazards</u> and <u>defects</u> and <u>describes</u> technological guidelines, which can be used to develop <u>control</u> <u>measures</u> and <u>corrective action</u>. At a particular step only the hazards and defects, which are likely to be introduced or controlled at that step, are listed. It should be recognised that in preparing a HACCP and/or DAP plan it is essential to consult Section 5 which provides guidance for the application of the principles of HACCP and DAP analysis. However, within the scope of this Code of Practice it is not possible to give details of critical limits, monitoring, record keeping and verification for each of the steps since these are specific to particular hazards and defects.

In general, the processing of fresh, frozen fish and minced fish, will range in sophistication. In its simplest form, the processing of fresh and frozen fish may be presented in a raw state such as dressed, fillets, and minced to be distributed in markets and institutions or used in processing facilities. For the latter, the processing of fresh, frozen and minced fish is often an intermediate step to the production of value added products (for example, smoked fish as described in section 12, canned fish as described in section 13, frozen breaded or battered fish as described in section 15). Traditional methods often prevail in the design of a process. However, modern scientific food technology is having an increasingly important role in enhancing the preservation and shelf-stability of a product. Regardless of the complexity of a particular process, the fabrication of the desired product relies on the consecutive execution of individual steps. As stressed by this Code, the application of appropriate elements of the pre-requisite program (Section 3) and HACCP principles (Section 5) at these steps will provide the processor with reasonable assurance that the essential quality, composition and labelling provisions of the appropriate Codex standard will be maintained and food safety issues controlled.

The example of the flow diagram (Figure 6.1) will provide guidance to some of the common steps involved in a fish fillet preparation line, and three examples of final product types: modified atmosphere packaging (MAP), minced and frozen fish. As in the further processing of fresh fish in a MAP product, or minced or frozen fish, the section labelled "Fish Preparation" is used as the basis for all the other fish processing operations (Sections 7-15), where appropriate.

References correspond to relevant Sections of the Code.

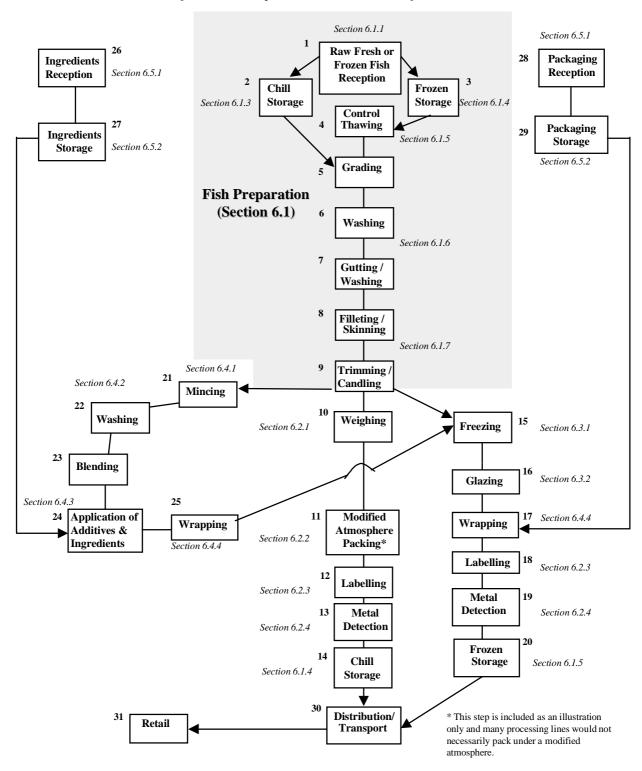


Figure 6.1 Example of a flow chart of a fish fillet preparation line, including MAP, mincing and freezing operations

6.1 FINFISH PREPARATION

The hygienic conditions and technical manner in which fish are prepared is similar and is not influenced greatly by its intended purpose (for direct distribution or for further processing). However, variations will exist in the form in which the fresh fish flesh is to be utilised. The forms may include, but not limited to, dressed, fillets or steaks.

6.1.1 Raw, Fresh or Frozen Fish Reception (Processing Steps 1)

<u>Potential Hazards</u>: Microbiological pathogens, viable parasites, biotoxins, chemicals (including

veterinary drug residues) and physical contamination.

Potential Defects: Decomposition, parasites, physical contamination

Technical Guidance:

for raw fish material, product specifications could include the following characteristics:

- organoleptic characteristics such as appearance, odour, texture, etc;
- chemical indicators of decomposition and/or contamination, for example, TVBN, histamine, heavy metals, pesticide residues, nitrates etc;
- microbiological criteria, in particular for intermediate raw materials, to prevent the processing of raw material containing microbial toxins;
- foreign matter;
- physical characteristics such as size of fish;
- species homogeneity.
- training in species identification and communication in product specification should be provided
 to fish handlers and appropriate personnel to ensure a safe source of incoming fish where written
 protocols exist. Of special consideration, are the reception and sorting of fish species that poses a
 risk of biotoxins such as ciguatoxin in large carnivorous tropical and sub-tropical reef fish or
 scombrotoxin in scombroid species or parasites;
- skills should be acquired by fish handlers and appropriate personnel in sensory evaluation techniques to ensure raw fish meet essential quality provisions of the appropriate Codex standard;
- fish requiring gutting on arrival at the processing facility should be gutted efficiently, without undue delay and with care to avoid contamination (see Section 6.1.5 Washing & Gutting);
- fish should be rejected if it is known to contain harmful, decomposed or extraneous substances, which will not be reduced or eliminated to an acceptable level by normal procedures of sorting or preparation;
- information about the harvesting area.

6.1.1.1 Sensory Evaluation of Fish

The best method of assessing the freshness or spoilage of fish is by sensory evaluation techniques⁵. It is recommended that appropriate sensory evaluation criteria be used to evaluate the acceptability of fish and to eliminate fish showing loss of essential quality provisions of the appropriate Codex standards. As an example, fresh white fish species are considered unacceptable when showing the following characteristics:

Skin / Slime: dull, gritty colours with yellow brown dotting slime.

Eyes: Concave, opaque, sunken discoloured.

Gills: grey – brown or bleached, slime opaque yellow, thick or clotting.

Odour: flesh odour amines, ammonia, milky lactic, sulphide, faecal, putrid, rancid.

⁵ Guidelines for Sensory Evaluation of Fish and Shelfish in Laboratories (CAC/GL 31-1999)

6.1.2 Chilled Storage (Processing Steps 2 & 14)

<u>Potential Hazards</u>: Microbiological pathogens and biotoxins.

<u>Potential Defects</u>: Decomposition, physical damage.

Technical Guidance:

• fish should be moved to the chill storage facility without undue delay;

- the facility should be capable of maintaining the temperature of the fish between $[0^{\circ}\text{C} +4^{\circ}\text{C}]$;
- the chill room should be equipped with a calibrated indicating thermometer. Fitting of a recording thermometer is strongly recommended;
- stock rotation plans should ensure proper utilisation of the fish;
- the fish should be stored in shallow layers and surrounded by sufficient finely divided ice or with a mixture of ice and of water before processing;
- fish should be stored such that damage will be prevented from over-stacking or over-filling of boxes;
- fish should be rejected if known to contain harmful, decomposed or extraneous substances, which
 will not be eliminated or reduced to an acceptable level by normal procedures of sorting or
 preparation. An appropriate assessment should be carried out to determine the reason(s) for loss
 of control and the HACCP or DAP plan should be modified where necessary;
- where appropriate replenish ice supply on the fish or alter temperature of the room.

6.1.3 Frozen Storage (Processing Steps 3 & 20)

Potential Hazards: Unlikely. Microbiological pathogens, toxins, viable parasites

<u>Potential Defects</u>: Dehydration, rancidity, loss of nutritional quality

Technical Guidance:

- the facility should be capable of maintaining the temperature of the fish at or colder than -18°C, and with minimal temperature fluctuations;
- the store should be equipped with a calibrated indicating thermometer. Fitting of a recording thermometer is strongly recommended;
- a systematic stock rotation plan should be developed and maintained;
- product should be glazed and/or wrapped to protect it from dehydration;
- fish should be rejected if known to contain defects, which subsequently cannot be reduced or eliminated to an acceptable level by re-working. An appropriate assessment should be carried out to determine the reason(s) for loss of control and the DAP plan modified where necessary.

6.1.4 Control Thawing (Processing Step 4)

<u>Potential Hazards</u>: Microbiological pathogens and biotoxins

<u>Potential Defects</u>: Decomposition

<u>Technical Guidance</u>:

- the thawing method should be clearly defined and should address the time and temperature of thawing, temperature measuring instrument used and placement of device for measurement. The thawing schedule (time and temperature parameters) should be carefully monitored. Selection of the thawing method should take into account in particular the thickness and uniformity of size of the products to be thawed;
- thawing time and temperature and fish temperature critical limits should be selected so as to control the development of micro-organisms, histamine, where high risk species are concerned or persistent and distinctive objectionable odours or flavours indicative of decomposition or rancidity;

- where water is used as the thawing medium, it should be of potable quality;
- where recycling of water is used, care should be taken to avoid the build up of microorganisms;
- where water is used, circulation should be sufficient to produce even thawing;
- during thawing, according to the method used, products should not be exposed to excessively high temperatures;
- particular attention should be paid to controlling condensation and drip from the fish. An effective drainage should be made;
- after thawing, fish should be immediately processed or refrigerated and kept at the adequate temperature (temperature of melting ice);
- fish should be rejected if known to contain harmful, decomposed or extraneous substances, which
 will not be eliminated or reduced to an acceptable level by normal procedures of sorting or
 preparation. An appropriate assessment should be carried out to determine the reason(s) for loss
 of control and the HACCP or DAP plan should be modified where necessary;
- the thawing schedule should be reviewed as appropriate and amended where necessary.

6.1.5 Washing and Gutting (Processing Steps 6 & 7)

<u>Potential Hazards</u>: Microbiological pathogens and biotoxins

<u>Potential Defects</u>: Presence of viscera, bruising, off-flavours, cutting faults.

Technical Guidance:

- gutting is considered complete when the intestinal tract and internal organs have been removed;
- an adequate supply of clean sea water or potable water should be available for washing of:
 - whole fish to remove foreign debris and reduce bacterial load prior to gutting;
 - gutted fish to remove blood and viscera from the belly cavity;
 - surface of fish to remove any loose scales;
 - gutting equipment and utensils to minimise build-up of slime and blood and offal;
- depending on the vessel or processing facility product flow pattern and where a prescribed critical
 limit for staging time and temperature regime has been established for the control of histamine or
 a defect, the gutted fish should be drained and well iced or appropriately chilled in clean
 containers and stored in specially designated and appropriate areas within the processing facility;
- separate and adequate storage facilities should be provided for the fish roe, milt and livers, if these are saved for later utilisation.

6.1.6 Filleting, Skinning, Trimming and Candling (Processing Steps 8 & 9)

<u>Potential Hazards</u>: Viable parasites, microbiological pathogens and biotoxins, presence of bones.

<u>Potential Defects</u>: Parasites, presence of bones, objectionable matter (e.g. skin, scales, etc.),

decomposition.

Technical Guidance:

- to minimise time delays, the design of the filleting line and candling line, where applicable, should be continuous and sequential to permit the uniform flow without stoppages or slow-downs and removal of waste;
- an adequate supply of clean sea water or potable water should be available for washing of:
 - fish prior to filleting or cutting especially fish that have been scaled;
 - fillets after filleting or skinning or trimming to remove any signs of blood, scales or viscera;
 - filleting equipment and utensils to minimise build-up of slime and blood and offal;

- for fillets to be marketed and designated as boneless, fish handlers should employ appropriate inspection techniques and use the necessary tools to remove bones not meeting Codex standards⁶⁷ or commercial specifications;
- The candling of skinless fillets by skilled personnel, in a suitable location which optimises the illuminating effect, is an effective technique in controlling parasites (in fresh fish) and should be employed when implicated fish species are being used;
- the candling table should be frequently cleaned during operation in order to minimise the microbial activity of contact surfaces and the drying of fish residue due to heat generated from the lamp;
- where a prescribed critical limit for staging time and temperature regime has been established for the control of histamine or a defect, the fish fillets should be well iced or appropriately chilled in clean containers, protected from dehydration and stored in appropriate areas within the processing facility.

6.2 PROCESSING OF MODIFIED ATMOSPHERE PACKED FISH

This section is designed to augment the processing of fresh fish section with additional operation steps pertaining specifically to the modified atmosphere packing of fish (see also Appendix I).

6.2.1 Weighing (Processing Step 10)

<u>Potential Hazards</u>: Unlikely

<u>Potential Defects</u>: Incorrect net weight

Technical Guidance:

• weigh scales should be periodically calibrated with a standardised mass to ensure accuracy.

6.2.2 Modified Atmosphere Packaging (Processing Step 11)

<u>Potential Hazards</u>: Subsequent microbiological pathogens and biotoxins, physical contamination

(metal).

Potential Defects: Subsequent decomposition

Technical Guidance:

The extent to which the shelf-life of the product can be extended by MAP will depend on the species, fat content, initial bacterial load, gas mixture, type of packaging material and, especially important, the temperature of storage. Refer to Appendix I for process control issues in modified atmosphere packaging.

- modified atmosphere packaging should be strictly controlled by:
 - monitoring the gas to product ratio;
 - types and ratio of gas mixtures used;
 - type of film used;
 - type and integrity of the seal;
 - temperature control of product during storage;
- fish flesh should be clear of the seam area;
- packaging material should be inspected prior to use to ensure that it is not damaged or contaminated;

Codex Standard for Quick Frozen Blocks of Fish Fillet, Minced Fish Flesh and Mixtures of Fillets and Minced Fish Flesh (Codex Stan. 165-1989, Rev.1-1995)

Codex Standard for Quick Frozen Fish Fillets (Codex Stan. 190-1995)

- packaging integrity of the finished product should be inspected at regular intervals by an
 appropriately trained personnel to verify the effectiveness of the seal and the proper operation of
 the packaging machine;
- following sealing, MAP products should be transferred carefully and without undue delay to chilled storage.

6.2.3 Labelling (Processing Steps 12 & 18)

<u>Potential Hazards</u>: Unlikely

<u>Potential Defects</u>: Incorrect labelling

Technical Guidance:

- prior to their application, labels should be verified to ensure that all information declared meet, where applicable, the Codex General Standard for the Labelling of Pre-packaged Foods⁸, labelling provisions of the appropriate Codex Standard for products and/or other relevant national legislative requirements;
- in many cases it will be possible to re-label incorrectly labelled products. An appropriate
 assessment should be carried out to determine the reason(s) for incorrect labelling and the DAP
 plan should be modified where necessary;

6.2.4 Metal Detection (Processing Steps 13 & 19)

<u>Potential Hazards</u>: Metal contamination

Potential Defects: Unlikely

Technical Guidance:

- it is important that line speeds are adjusted to allow for the proper functioning of a metal detector;
- routine procedures should be initiated to ensure product rejected by the detector is investigated as to the cause of the rejection;
- metal detectors, if used, should be periodically calibrated with a known standard to ensure proper operation;

6.3 PROCESSING OF FROZEN FISH

This section is designed to augment the processing of fresh fish section with additional operation steps pertaining specifically to the processing of frozen fish.

6.3.1 Freezing Process (Processing Step 15)

<u>Potential Hazards</u>: Viable parasites.

Potential Defects: Texture deterioration, development of rancid odours, freezer burn

Technical Guidance:

The fish product should be subjected to a freezing process as quickly as possible since unnecessary delays before freezing will cause temperature of the fish products to rise, increasing the rate of quality deterioration and reducing shelf-life due to the action of micro-organisms and undesirable chemical reactions.

• a time and temperature regime for freezing should be established and should take into consideration the freezing equipment and capacity; the nature of the fish product including thermal conductivity, thickness, shape and temperature and the volume of production, to ensure that the range of temperature of maximum crystallisation is passed through as quickly as possible;

⁸ Codex General Standard for the Labelling of Pre-packaged Foods (Codex Stan 1-1985, Rev. 2-1999)

- the thickness, shape and temperature of fish product entering the freezing process should be as uniform as possible;
- processing facility production should be geared to the capacity of freezers;
- frozen product should be moved to the cold storage facility as quickly as possible;
- the core temperature of the frozen fish should be monitored regularly for completeness of the freezing process;
- frequent checks should be made to ensure correct operation of freezing;
- accurate records of all freezing operations should be kept.

6.3.2 Glazing (Processing Step 16)

<u>Potential Hazards</u>: Microbiological pathogens, biotoxins

Potential Defects: Subsequent dehydration, incorrect net weight

Technical Guidance:

- glazing is considered complete when the entire surface of the frozen fish product is covered with a suitable protective coating of ice and should be free of exposed areas where dehydration (freezerburn) can occur;
- if additives are used in the water for glazing, care should be taken to ensure its proper proportion and application with product specifications;
- where the labelling of a product is concerned, information on the amount or proportion of glaze applied to a product or a production run should be kept and used in the determination of the net weight which is exclusive of the glaze;
- where appropriate monitoring should ensure that spray nozzles do not become blocked;
- where dips are used for glazing it is important to replace the glazing solution periodically to minimise the bacterial load and build-up of fish protein, which can hamper freezing performance;

6.4 PROCESSING OF MINCED FISH

This section is designed to augment the processing of fresh fish section (prior to mincing) and processing of frozen fish section (after mincing) with additional operation steps pertaining specifically to the processing of minced fish.

6.4.1 Mincing Fish Using Mechanical Separation Process (Processing Step 21)

<u>Potential Hazards</u>: Microbiological pathogens and biotoxins, physical contamination (metal, bones,

rubber from separator belt, etc).

<u>Potential Defects</u>: Incorrect separation (i.e. objectionable matter), decomposition, presence of

defect bones, parasites.

Technical Guidance:

- the separator should be fed continuously but not excessively;
- candling is recommended for fish suspected of high infestation with parasites;
- split fish or fillets should be fed to the separator so that the cut surface contacts the perforated surface;
- fish should be fed to the separator in a size that it is able to handle;
- in order to avoid time-consuming adjustments of the machinery and variations in quality of the finished product, raw materials of different species and types should be segregated and processing of separate batches should be carefully planned;
- the perforation sizes of the separator surface as well as the pressure on the raw material should be adjusted to the characteristics desired in the final product;

- the separated residual material should be carefully removed on a continuous or near-continuous basis to the next processing stage;
- temperature monitoring should ensure undue temperature rises of the product are avoided.

6.4.2 Washing of Minced Fish (Processing Step 22)

<u>Potential Hazards</u>: Microbiological pathogens and biochemical toxins.

<u>Potential Defects</u>: Poor colour, poor texture, excess of water

Technical Guidance:

- if necessary the mince should be washed and should be adequate for the type of product desired;
- stirring during washing should be carried out with care, but it should be kept as gentle as possible
 in order to avoid excessive disintegration of the minced flesh which will reduce the yield due to
 the formation of fines;
- the washed minced fish flesh may be partially de-watered by rotary sieves or centrifugal equipment and the process completed by pressing to appropriate moisture content;
- if necessary, and depending on eventual end-use, the de-watered mince should be either strained or emulsified;
- special attention should be taken to ensure mince being strained is kept cool;
- the resulting waste water should be disposed of in a suitable manner.

6.4.3 Blending and Application of Additives and Ingredients to Minced Fish (Processing Steps 23 & 24)

Potential Hazards: Physical contamination, non-approved additives and/or ingredients.

<u>Potential Defects</u>: Physical contamination, incorrect addition of additives.

Technical Guidance:

- if fish, ingredients and /or additives are to be added, they should be blended in the proper proportions to achieve the desired sensory quality;
- additives should comply with the requirements of the Codex General Standard for Food Additives;
- the minced fish product should be packaged and frozen immediately after preparation; if it is not frozen or used immediately after preparation it should be chilled.

6.4.4 Wrapping and Packing (Processing Steps 17 & 25)

Potential Hazards: Unlikely.

<u>Potential Defects</u>: Subsequent dehydration, decomposition

Technical Guidance:

- packaging material should be clean, sound, durable, sufficient for its intended use and of food grade material;
- the packaging operation should be conducted to minimise the risk of contamination and decomposition;
- products should meet appropriate standards for labelling and weights.

6.5 PACKAGING, LABELS & INGREDIENTS

6.5.1 Reception – Packaging, Labels & Ingredients (Processing Steps 26 & 28)

<u>Potential Hazards:</u> Microbiological pathogens biotoxins, chemical and physical contamination

<u>Potential Defects</u>: Misdescription

Technical Guidance:

• only ingredients, packaging material and labels complying with the processors' specification should be accepted into the processing facility;

- labels which are to be used in direct contact with the fish should be fabricated of a non-absorbent material and the ink or dye used on that label should be approved by the official agency having jurisdiction;
- ingredients and packaging material not approved by the official agency having jurisdiction should be investigated and refused at reception;

6.5.2 Storage - Packaging, Labels & Ingredients (Processing Steps 27 & 29)

Potential Hazards: Microbiological pathogens, biotoxins, chemical and physical contamination.

<u>Potential Defects:</u> Loss of quality characteristics of packaging materials or ingredients.

Technical Guidance:

ingredients and packaging should be stored appropriately in terms of temperature and humidity;

- a systematic stock rotation plan should be developed and maintained to avoid out of date materials;
- ingredients and packaging should be properly protected and segregated to prevent crosscontamination;
- defective ingredients and packaging should not be used.

SECTION 13 PROCESSING OF CANNED FISH AND SHELLFISH

In the context of recognising controls at individual processing steps, this section provides <u>examples</u> of potential <u>hazards</u> and <u>defects</u> and describes technological guidelines, which can be used to develop <u>control</u> <u>measures</u> and <u>corrective action</u>. At a particular step only the hazards and defects, which are likely to be introduced or controlled at that step, are listed. It should be recognised that in preparing a HACCP and/or DAP plan it is essential to consult Section 5 (Hazard Analysis Critical Control Point (HACCP) and Defect Action Point (DAP) Analysis) which provides guidance for the application of the principles of HACCP and DAP analysis. However, within the scope of this Code of Practice it is not possible to give details of critical limits, monitoring, record keeping and verification for each of the steps since these are specific to particular hazards and defects.

This section concerns the processing of heat processed sterilised canned fish and shellfish products which have been packed in hermetically sealed rigid or semi-rigid containers⁹ and intended for human consumption.

As stressed by this Code, the application of appropriate elements of the pre-requisite program (Section 3) and HACCP principles (Section 5) at these steps will provide the processor with reasonable assurance that the essential quality, composition and labelling provisions of the appropriate Codex standard will be maintained

Aseptic filling is not covered by this Code. Reference of the relevant code is made in Appendix XI.

and food safety issues controlled. The example of the flow diagram (Figure 13.1) will provide guidance to some of the common steps involved in a canned fish or shellfish preparation line.

References correspond to relevant Sections of the Code.

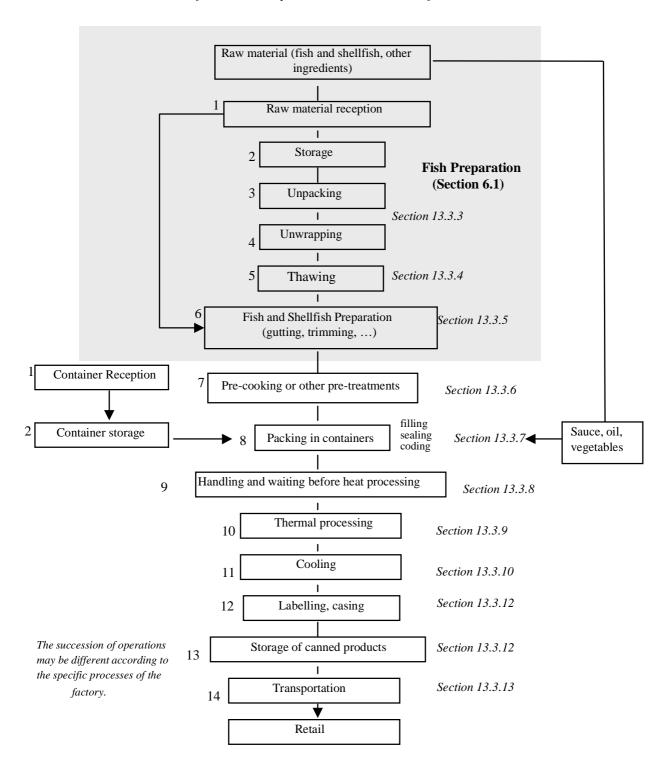


Figure 13.1 Example of a flow chart for the processing of canned fish and shellfish

13.1 GENERAL - ADDITION TO PRE-REQUISITE PROGRAMME

Section 3 (Pre-requisite programme) gives the minimum requirements for good hygienic practices for a processing facility prior to the application of hazard and defect analyses.

For fish and shellfish canneries, additional requirements to the guidelines described in Section 3 are necessary due to the specific technology involved. Some of them are listed below, but reference should also be made to the Recommended International Code of Hygienic Practice for Low-Acid and Acidified Low-Acid Canned Food (CAC/PRC 23-1979, Rev. 2 (1993)) for further information.

- design, working and maintenance of baskets and handling and loading devices aimed at retorting should be appropriate for the kind of containers and materials used. These devices should prevent any excessive abuse to the containers.
- an adequate number of efficient sealing machines should be available to avoid undue delay in processing;
- retorts should have a suitable supply of energy, vapour, water and/or air so as to maintain in it
 sufficient pressure during the heat treatment of sterilisation; their dimensions should be adapted to
 the production to avoid undue delays;
- every retort should be equipped with an indicating thermometer, a pressure gauge and a time and temperature recorder,
- an accurate clearly visible clock should be installed in the retorting room;
- canneries using steam retorts should consider installing automatic steam controller valves;
- Instruments used to control and to monitor in particular the thermal process should be kept in good condition and should be regularly verified or calibrated. Calibration of instruments used to measure temperature should be made in comparison with a reference thermometer. This thermometer should be regularly calibrated. Records concerning the calibration of instruments should be established and kept.

13.2 IDENTIFICATION OF HAZARDS AND DEFECTS

Refer also to Section 4.1 (Potential Hazards Associated with Fresh Fish and Shellfish)
This section describes the main potential hazards and defects specific to canned fish and shellfish.

13.2.1 Hazards

A Biological Hazards

A1 Naturally occurring marine toxins

Biotoxins such as tetrodotoxines or ciguatoxines are known to be generally heat-stable, so the knowledge of the identity of the species and/or the origin of fish intended for processing is important.

Phycotoxins such as DSP, PSP or ASP are also heat stable, so it important to know the origin and the status of the area of origin of molluscan shellfish or other affected species intended for processing.

A2 Microbiological toxins

Histamine

Histamine is heat-stable, and so its toxicity remains practically intact in containers. Good practices for the conservation and handling from capture to heat processing are essential to prevent the histamine production. The Codex Commission adopted in its standards for some fish species maximum levels tolerated for histamine.

Clostridium botulinum

The botulism risk usually appears after an inadequate heat processing and inadequate container integrity. The toxin is heat-sensitive, on the other hand, the destruction of *Clostridium botulinum* spores, in particular from proteolytic strains, requires high sterilisation values. The heat processing effectiveness depends on the contamination level at the time of the treatment. Therefore, it is advisable to limit proliferation and the contamination risks during processing.

Staphylococcus aureus

Toxins from *Staphylococcus aureus* can be present in a highly contaminated raw material or can be produced by bacterial proliferation during processing. These toxins are heat-resistant, so they have to be taken into account in the hazard analysis.

B Chemical Hazards

Care should be taken to avoid contamination of the product from components of the containers (e.g. lead...).

C Physical Hazards

Containers prior to filling may contain materials such as metal or glass fragments.

13.2.2 Defects

Potential defects are outlined in the essential quality, labelling and composition requirements described in the relevant Codex Standards listed in Appendix XII. Where no Codex Standard exists regard should be made to national regulations and/or commercial specifications.

End product specifications outlined in Appendix IX describe optional requirements specific to canned products.

13.3 PROCESSING OPERATIONS

Processors can also refer to the Recommended International Code of Hygienic Practice for Low-Acid and Acidified Low-Acid Canned Foods (CAC/RCP 23-1979, Rev. 2 (1993)) in order to obtain detailed advice on canning operations.

13.3.1 Raw Materials reception

13.3.1.1 Fish and shellfish (Processing step 1)

<u>Potential Hazards:</u> Chemical and biochemical contamination (DSP, PSP, histamine, heavy

metals...)

<u>Potential Defects:</u> Species substitution, decomposition, parasites

Technical Guidance:

Refer to section 6.1.1 (Raw Fresh or Frozen Fish Reception); and also:

• When live shellfish (crustaceans) are received for canning processing, inspection should be carried out in order to discard dead or badly damaged animals.

13.3.1.2 Container and packaging materials (Processing step 1)

<u>Potential Hazards:</u> Subsequent microbiological contamination

<u>Potential Defects:</u> Tainting of the product

Technical Guidance:

Refer to section 6.5.1 (Raw Material Reception – Packaging, Labels & Ingredients); and also:

- containers and packaging materials should be suitable for the type of product, the conditions provided for storage, the filling, sealing and packaging equipment and the transportation conditions;
- the containers in which fish and shellfish products are canned should be made from suitable material and constructed so that they can be easily closed and sealed to prevent the entry of any contaminating substance;
- containers for canned fish and shellfish should meet the following requirements:
 - they should protect the contents from contamination by micro-organisms or any other substance;
 - their inner surfaces should not react with the contents in any way that would adversely affect the product or the containers;
 - their outer surfaces should be resistant to corrosion under any likely conditions of storage;
 - they should be sufficiently durable to withstand the mechanical and thermal stresses encountered during the canning process and to resist physical damage during distribution;

13.3.1.3 Other ingredients (Processing step 1)

Refer to section 6.5.1 (Raw Material Reception – Packaging, Labels & Ingredients).

13.3.2 Storage of raw material, containers and packaging materials

13.3.2.1 Fish and shellfish (Processing step 2)

Refer to sections 6.1.2 (Chilled storage), 6.1.3 (Frozen storage and 7.6.2 Conditioning and storage of molluscan shellfish in sea water tanks, basins, etc.)

13.3.2.2 Containers and packaging (Processing step 2)

<u>Potential Hazards:</u> Unlikely

Potential Defects: Foreign matters

Technical Guidance:

Refer to section 6.5.2 (Raw Material Storage - Packaging, Labels & Ingredients); and also:

- all materials for containers or packages should be stored in satisfactory clean and hygienic conditions;
- during storage, empty containers and covers should be protected from dirt, moisture and temperature fluctuations, in order to avoid condensations on containers and in the case of tin cans, the development of corrosion;
- during loading, stowing, transportation and unloading of empty containers, any shock should be avoided. Containers shouldn't be stepped on. These precautions become more imperative when containers are put in bags or on pallets. Shocks can deform the containers (can body or flange), that can compromise tightness (shocks on the seam, deformed flange) or be prejudicial to appearance.

13.3.2.3 Other ingredients (Processing step 2)

Refer to section 6.5.2 (Raw Material Storage - Packaging, Labels & Ingredients).

13.3.3 Unwrapping, unpacking (Processing steps 3 and 4)

<u>Potential Hazards:</u> Unlikely

<u>Potential Defects:</u> Foreign matters

Technical Guidance:

• During unwrapping and unpacking operations, precautions should be taken in order to limit product contamination and foreign matters introduction into the product. To avoid microbial proliferation, waiting periods before further processing should be minimised.

13.3.4 Thawing (Processing step 5)

Refer to section 6.1.4 (Control Thawing)

13.3.5 Fish and shellfish preparatory processes (Processing step 6)

13.3.5.1 Fish preparation (gutting, trimming...)

<u>Potential Hazards:</u> Microbiological contamination biochemical development (histamine)

<u>Potential Defects:</u> Objectionable matters (viscera, skin, scales, ... in certain products), off flavours,

presence of bones, parasites...

Technical Guidance:

Refer to sections 6.1.5 (Washing and Gutting) and 6.1.6 (Filleting, Skinning, Trimming and Candling); and also:

• when skinning of fish is operated by soaking in soda solution, a particular care should be taken to carry out an appropriate neutralisation.

13.3.5.2 Preparation of molluscs and crustaceans

Potential Hazards: Microbiological contamination, hard shell fragments

<u>Potential Defects:</u> Objectionable matters

Technical Guidance:

Refer to sections 7.7 (Heat Treatment/Heat Shocking of Molluscan Shellfish in Establishment; and also:

- when live shellfish are used, inspection should be carried out in order to discard dead or badly damaged animals;
- particular care should be taken to ensure that shell fragments are removed from shellfish meat.

13.4 PRE-COOKING AND OTHER TREATMENTS

13.4.6 Pre-Cooking

Potential hazards: chemical contamination (polar components of oxidised oils), microbiological or

biochemical (scombrotoxin) growth.

<u>Potential defects</u>: water release in the final product (for products canned in oil), abnormal

flavours.

Technical guidance:

13.4.6.1 General Considerations

- methods used to pre-cook fish or shellfish for canning should be designed to bring about the
 desired effect with a minimum delay and a minimum amount of handling; the choice of method
 is usually strongly influenced by the nature of the treated material. For products canned in oil
 such as sardines or tunas, pre-cooking should be sufficient in order to avoid excessive release of
 water during heat processing;
- means should be found to reduce the amount of handling subsequent to pre-cooking, wherever practical;
- if eviscerated fish is used, then the fish should be arranged in the belly down position for precooking to allow for the drainage of fish oils and juices which may accumulate and affect product quality during the heating process;
- where appropriate, molluscan shellfish, lobsters and crabs, shrimps and prawns and cephalopods should be pre- cooked according to technical guidance laid down in sections 7 (Processing of Molluscan Shellfish), 8 (Processing of Lobsters and Crabs), 9 (Processing of Shrimps and Prawns) and 10 (Processing of Cephalopods);
- care should be taken to prevent temperature abuse of scombrotoxic species before pre-cooking.

13.4.6.1.2 Pre-cooking Schedule

- the pre-cooking method, in particular, in terms of time and temperature, should be clearly defined. The pre-cooking schedule should be checked;
- fish pre-cooked together in batches should be very similar in size. It also follows that they should all be at the same temperature when they enter the cooker.

13.4.6.1.3 Control of Quality of Pre-cooking Oils and Other Fluids

- only good quality vegetable oils should be used in pre-cooking fish or shellfish for canning [refer to relevant Codex Standard for vegetable oils];
- cooking oils should be changed frequently in order to avoid the formation of polar compounds. Water used for pre-cooking should also be changed frequently in order to avoid contaminants;
- care must be taken that the oil or the other fluids used such as vapour or water do not impart an undesirable flavour to the product.

13.4.6.1.4 Cooling

- except for products, which are packed when still hot, cooling of pre-cooked fish or shellfish should
 be done as quickly as possible to bring the product temperatures in a range limiting proliferation
 or toxin production, and under conditions where contamination of the product can be avoided;
- where water is used to cool crustacea for immediate shucking, it should be potable water or clean seawater. The same water should not be used for cooling more than one batch.

13.4.6.2 Smoking

• refer to section 12 (Processing of smoked fish)

13.4.6.3 Use of Brine and Other Dips

Potential hazards: microbiological and chemical contamination by the dip solution

<u>Potential defects</u>: adulteration (additives), abnormal flavours.

Technical guidance:

 Where fish or shellfish are dipped or soaked in brine or in solutions of other conditioning or flavouring agents or additives in preparation for canning, solution strength and time of immersion should both be carefully controlled to bring about the optimum effect;

- dip solutions should be replaced and dip tanks and other dipping apparatus should be thoroughly cleaned at frequent intervals;
- care should be taken to ascertain whether or not the ingredients or additives used in dips would be permitted in canned fish and shellfish by the related Codex Standards and in the countries where the product will be marketed.

13.4.7 Packing in Containers (Filling, Sealing and Coding) (Processing Step 8)

13.4.7.1 Filling

<u>Potential hazards</u>: [microbiological growth (waiting period)], microbiological growth and

recontamination after heat processing due to incorrect filling or faulty

containers.

<u>Potential defects</u>: incorrect weight, foreign matter.

Technical guidance

- a representative number of containers and covers should be inspected immediately before delivery to the filling machines or packing tables to ensure that they are clean, undamaged and without visible flaws:
- if necessary, empty containers should be cleaned. It is also a wise precaution to have all containers turned upside down to make certain that they do not contain any foreign material before they are used;
- care should also be taken to remove faulty containers, because they can jam a filling or sealing machine, or cause trouble during heat processing (bad sterilisation, leaks);
- empty containers should not be left on the packing tables or in conveyor systems during clean up of premises to avoid contamination or splashes;
- where appropriate, to prevent microbial proliferation, containers should be filled with hot fish and shellfish (> 63°C, for example for fish soups) or should be filled quickly (the shortest possible waiting period) after the end of the pre-treatments;
- if the fish and shellfish must be held for a long time before packing into containers, they should be chilled:
- containers of canned fish and shellfish should be filled as directed in the scheduled process;
- mechanical or manual filling of containers should be checked in order to comply with the filling
 rate and the headspace specified in the adopted sterilisation schedule. A regular filling is
 important not only for economical reasons, but also because the heat penetration and the container
 integrity can be affected by excessive filling changes;
- the necessary amount of headspace will depend partly on the nature of the contents. The filling should also take into account the heat processing method. Headspace should be allowed as specified by the container manufacturer;
- furthermore, containers should be filled such as the end product meets the regulatory provisions or the accepted standards concerning weight of contents;
- where canned fish and shellfish is packed by hand, there should be a steady supply of fish, shellfish and eventually other ingredients. Build-up of fish and shellfish, as well as filled containers at the packing table should be avoided;
- the operation, maintenance, regular inspection, calibration and adjustment of filling machines should received particular care. The machine manufacturers' instructions should be carefully followed;
- the quality and the amount of other ingredients such as oil, sauce, vinegar...should be carefully controlled to bring about the optimum desired effect;
- if fish has been brine-frozen or stored in refrigerated brine, the amount of salt absorbed should be taken into consideration when salt is added to the product for flavouring;

- filled containers should be inspected:
 - to ensure that they have been properly filled and will meet accepted standards for weight of
 - and to verify product quality and workmanship just before they are closed;
- manual filled products such as small pelagic fish should be carefully checked by the operators to verify that container flanges or closure surface have not any product residues, which could impede the formation of a hermetic seal. For automatic filled products, a sampling plan should be implemented.

13.4.7.2 Sealing

Sealing the container is one of the most essential processes in canning.

<u>Potential hazards</u>: subsequent contamination due to a bad seam

<u>Potential defects</u>: unlikely

Technical guidance

- the operation, maintenance, regular inspection and adjustment of sealing machines should received particular care. The sealing machines should be adapted and adjusted for each type of container and each closing method which are used. Whatever the type of sealing equipment, the manufacturers or equipment supplier's instructions should be followed meticulously;
- seams and other closures should be well formed with dimensions within the accepted tolerances for the particular container;
- qualified personnel should conduct this operation;
- if vacuum is used during packing, it should be sufficient to prevent the containers from bulging under any condition (high temperature or low atmospheric pressure) likely to be encountered during the distribution of the product. This is useful for deep containers or glass containers. It is difficult and hardly necessary to create a vacuum in shallow containers that have relatively large flexible covers;
- excessive vacuum may cause the container to panel, particularly if the headspace is large, and
 may also cause contaminants to be sucked into the container if there is a slight imperfection in the
 seam;
- to find the best methods to create vacuum, competent technologists should be consulted;
- regular inspections should be made during production to detect potential external defects on containers. At intervals sufficiently close to each other in order to guarantee a closure in accordance with specifications, the operator, the supervisor of the closure or any other competent person should examine the seams or the closure system for the other types of containers, which are used. Inspections should consider for example vacuum measurements and seam teardown. A sampling plan should be used for the checks;
- in particular, at each start of the production line and at each change in container dimensions, after a jamming, a new adjustment or a restarting after a prolonged stop of the sealing machine, a check should be carried out;
- all appropriate observations should be recorded.

13.4.7.3 Coding

<u>Potential hazards</u>: subsequent contamination due to damaged containers

<u>Potential defects</u>: loss of traceability due to an incorrect coding.

Technical guidance

- each container of canned fish and shellfish should bear indelible code markings from which allimportant details concerning its manufacture (type of product, cannery where the canned fish or shellfish was produced, production date, etc.) can be determined
- coding equipment must be carefully adjusted so that the containers are not damaged and the code remains legible;
- coding may sometimes be carried out after the cooling step.

13.4.8 Handling of Containers After Closure - Staging Before Heat Processing (Processing Step 9)

Potential hazards: [microbiological growth (waiting period)], subsequent contamination due to

damaged containers.

<u>Potential defects</u>: Unlikely

Technical guidance

- containers after closure should always be handled carefully in such a way as to prevent every damage capable to cause defects and microbiological recontamination;
- if necessary, filled and sealed metal containers should be thoroughly washed before heat processing to remove grease, dirt and fish or shellfish stains on their outside walls;
- to avoid microbial proliferation, the waiting period should be as short as possible;
- if the filled and sealed containers must be held for a long time before heat processing, the product should be held at temperature conditions which minimise microbial growth;
- every cannery should develop a system, which will prevent non heat-processed canned fish and shellfish from being accidentally taken past the retorts into the storage area.

13.4.9 Thermal Processing (Processing Step 10)

Heat processing is one of the most essential operations in canning.

Canners can refer to the Recommended International Code of Hygienic Practice for Low-Acid and Acidified Low-Acid Canned Foods (CAC/RCP 23-1979, rev. 2 in 1993) in order to obtain detailed advice on heat processing. In this Section, only some essential elements are pointed out.

<u>Potential hazards</u>: survival of spores of Clostridium botulinum.

Potential defects: survival of micro-organisms responsible of decomposition

Technical guidance

13.4.9.1 Sterilisation Schedule

- to determine the sterilisation schedule, at first, the heat process required to obtain the commercial sterility should be established taking into account some factors (microbial flora, dimensions and nature of the container, product formulation, etc.). A sterilisation schedule is established for a certain product in a container of a given size;
- then, competent canning technologist to take into account the sterilisation equipment at disposal
 and the product quality, which is desired, should carry out heat penetration tests. This heat
 penetration in the product must be established in the most unfavourable conditions likely to occur
 during processing. Standard heat processing procedures and experimentally established
 sterilisation schedules should be checked and validated by an expert to confirm that the values are
 appropriate for each product and retort;
- before any changes in operations (initial temperature of filling, product composition, size of containers, fullness of the retort, etc.) are made, competent technologists should be consulted as to the need for re-evaluation of the process.

13.4.9.2 Heat Processing Operation

- only qualified and properly trained personnel should operate retorts. Therefore it is necessary that
 retort operators control the processing operations and ensure the sterilisation schedule is closely
 followed, including meticulous care in timing, monitoring temperatures and pressures, and in
 maintaining records;
- it is essential to comply with the initial temperature described in the schedule process to avoid under-processing. If the filled containers were held at refrigerated temperatures because of a too long waiting period before heat processing, the sterilisation schedule should take into account these temperatures;
- in order that the heat processing is effective and process temperature is controlled, air must be evacuated from the retort through a venting procedure that is deemed efficient by a competent technologist. Container size and type, retort installation and loading equipment and procedures should be considered:
- the timing of the heat processing should not commenced until the specified heat processing temperature has been reached, and the conditions to maintain uniform temperature throughout the retort achieved, in particular, until the minimum safe venting time has elapsed;
- for other types of retorts (water, steam/air, flame, etc.) refer to the Recommended International Code of Hygienic Practice for Low-Acid and Acidified Low-Acid Canned Foods (CAC/RCP 23-1979, rev. 2 in 1993);
- if canned fish and shellfish in different size containers are processed together in the same retort load care must be taken to ensure the process schedule used is sufficient to provide commercial sterility for all container sizes processed;
- when processing fish and shellfish in glass containers, care must be taken to ensure that the initial temperature of the water in the retort is slightly lower than that of the product being loaded. The air pressure should be applied before the water temperature is raised.

13.4.9.3 Monitoring of Heat Processing Operation

- during the application of heat processing, it is important to ensure that the sterilisation process and factors such as container filling, minimal internal depression at closing, retort loading, initial product temperature, etc. are in accordance with the sterilisation schedule;
- retort temperatures should always be determined from the indicating thermometer, never from the temperature recorder;
- permanent records of the time, temperature and other pertinent details should be kept concerning each retort load;
- the thermometers should be tested regularly to ensure that they are accurate. Calibration records should be maintained;
- inspections should be made periodically to ensure that retorts are equipped and operated in a manner that will provide thorough and efficient heat processing, that each retort is properly equipped, filled and used, so that the whole load is brought up to processing temperature quickly and can be maintained at that temperature throughout the whole of the processing period;
- the inspections should be made under the guidance of a competent technologist.

13.4.10 Cooling (Processing Step 11)

<u>Potential hazards</u>: recontamination due to a bad seam and contaminated water <u>Potential defects</u>: formation of struvite crystals, buckled containers, scorch.

Technical guidance:

• after heat processing, canned fish and shellfish should, wherever practical, be water cooled under pressure to prevent deformations, which could result in a loss of tightness. In case of recycling, only potable chlorinated water should be used for this purpose. The residual chlorine level in

- cooling water and the contact time during cooling should be checked in order to minimise the risk of post-processing contamination;
- in order to avoid organoleptic defects of the canned fish and shellfish, such as scorch or overcooking, the internal temperature of containers should be lowered as quickly as possible;
- for glass containers, the temperature of the coolant in the retort should be, at the beginning, lowered slowly in order to reduce the risks of breaking due to thermal shock;
- where canned fish and shellfish products are not cooled in water after heat processing, they should be stacked in such a way that they will cool rapidly in air.
- heat processed canned fish and shellfish should not be touched by hand or articles of clothing unnecessarily before they are cooled and thoroughly dry. They should never be handled roughly or in such a way that their surfaces, and in particular their seams, are exposed to contamination;
- rapid cooling of canned fish and shellfish avoids the formation of struvite crystals;
- every cannery should develop a system to prevent unprocessed containers being mixed with processed containers.

13.4.10.1 Monitoring After Heat Processing and Cooling

- canned fish and shellfish should be inspected for faults and for quality assessment soon after it is produced and before labelling;
- representative samples from each code lot should be examined to ensure that the containers do not
 exhibit external defects and the product meets the standards for weight of contents, vacuum,
 workmanship and wholesomeness. Texture, colour, odour, flavour and condition of the packing
 medium should be assessed;
- if desired, stability tests could be made in order to verify in particular the heat processing;
- this examination should be made as soon as practical after the canned fish and shellfish have been
 produced, so that if there are any faults due to failings on the part of cannery workers or canning
 equipment, these failings can be corrected without delay. Segregating and properly disposing of
 all defective units or lots that are unfit for human consumption should be ensured.

13.4.11 Labelling, Casing and Storage of Finished Products (Processing steps 12 and 13)

<u>Potential hazards</u>: subsequent recontamination due to the damage of containers or to an

exposition to extreme conditions

<u>Potential defects</u>: incorrect labelling

Technical guidance

- the materials used for labelling and casing canned fish and shellfish should not be conducive to corrosion of the container. Cases should have an adequate size in order that the containers fit them and are not damaged by any move inside. Cases and boxes should be the correct size and strong enough to protect the canned fish and shellfish during distribution;
- code marks appearing on containers of canned fish and shellfish should also be shown on the cases in which they are packed;
- the labelling should be made as referred in 6.2.3;
- storage of canned fish and shellfish should be made in order not to damage the containers. In particular, pallets of finished products should not be stacked excessively high and the forklift trucks used for the storage should be used in a proper manner;
- canned fish and shellfish should be so stored that they will be kept dry and not exposed to extremes of temperature.

13.4.12 Transportation of Finished Products (Processing step 14)

<u>Potential hazards</u>: subsequent recontamination due to the damage of containers or to an

exposition to extreme conditions

<u>Potential defects</u>: Unlikely

Technical guidance

Refer to section 17 (Transportation); and also:

- transportation of canned fish and shellfish should be made in order not to damage the containers.
 In particular, the forklift trucks used during the loading and unloading should be used in a proper manner:
- cases and boxes should be completely dry. In fact, moisture has effects on the mechanical characteristics of boxes and the protection of containers against damages during transportation couldn't be sufficient;
- metal containers should be kept dry during transportation in order to avoid corroding and/or rust.