

HAZARD ANALYSIS CRITICAL CONTROL POINT (HACCP) SYSTEM: AN OVERVIEW

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HACCP is an acronym that stands for Hazard Analysis Critical Control Point. HACCP is a systematic approach to the identification, evaluation, and control of food safety hazards. It is a proactive strategy where hazards are identified and assessed, and control measures are developed to prevent, reduce, or eliminate a hazard. The goal of HACCP is to produce food products that are 'SAFE' for human consumption. The result of the implementation of the HACCP Plan is known as the HACCP System. Use of HACCP system will move a food producing company from sole retrospective end product testing approach towards a preventive quality assurance approach.

HACCP Definitions

- ▶ **HACCP:** A systematic approach to the identification, evaluation, and control of food safety hazards.
- ▶ **Prerequisite Programs:** Procedures, including Good Manufacturing Practices, that address operational conditions providing the foundation for the HACCP system.
- ▶ **Risk:** an estimate of the likelihood of the occurrence of a hazard
- ▶ **Severity of risk:** the seriousness of a hazard if not properly controlled
- ▶ **Hazard:** any biological, chemical or physical property that may be expected to cause an unacceptable health risk to consumers if present in the product
- ▶ **Hazard Analysis:** The process of collecting and evaluating information on hazards associated with the food under consideration to decide which are significant and must be addressed in the HACCP plan.
- ▶ **HACCP Team:** The group of people who are responsible for developing, implementing and maintaining the HACCP system.

- ▶ **Critical control point (CCP)** – a specific point in a process where control can be applied to eliminate or reduce the risk of a hazard to an acceptable level
- ▶ **CCP Decision Tree:** A sequence of questions to assist in determining whether a control point is a CCP.
- ▶ **Control Measure:** Any action or activity that can be used to prevent, eliminate or reduce a significant hazard.
- ▶ **Monitor:** To conduct a planned sequence of observations or measurements to assess whether a CCP is under control and to produce an accurate record for future use in verification.
- ▶ **Validation:** That element of verification focused on collecting and evaluating scientific and technical information to determine if the HACCP plan, when properly implemented, will effectively control the hazards.
- ▶ **Verification:** Those activities, other than monitoring, that determine the validity of the HACCP plan and that the system is operating according to the plan.
- ▶ **HACCP Plan:** The written document which is based upon the principles of HACCP and which delineates the procedures to be followed.

History of HACCP:

The concept of HACCP was first used in the US space program to ensure safe food for astronauts without relying on end-product testing. It was developed in 1960s by a team from Pillsbury company to produce 'zero defects' food products for the NASA astronauts. HACCP was adopted in 1973 by the USFDA for low acid canned food regulations ($\text{pH} < 4.6$). The HACCP standard developed by the U.S. National Advisory Committee on the Microbiological Criteria for Food (NACMCF) was later adopted by the Codex Alimentarius Commission and published as the first international HACCP standard in 1992. In 1997, the USFDA made HACCP mandatory for all seafood processors in the US as well as for those foreign plants exporting to

the US. The Canadian Food Inspection Agency (CFIA) made HACCP mandatory for all Canadian seafood processors in 1998. Similar legislation has been imposed in the EU for food processors within the EU and those exporting to EU countries. The Codex Alimentarius Commission – an organization linked to the WHO (World Health Organization) and FAO (Food and Agriculture Organization of the United Nations), with representatives from more than 180 countries – has published the HACCP approach for managing food safety risks. This approach is being adopted by government agencies and industry organizations globally. Although specific regulations in different parts of the world continue to evolve, they typically feature some common elements from HACCP:

Seven fundamental principles of HACCP

Principle 1 - Conduct hazard analysis for each product.

Principle 2 - Identify Critical Control Points

Principle 3 - Establish critical limits

Principle 4 - Establish CCP monitoring procedures

Principle 5 - Establish Corrective action when critical limits have been exceeded

Principle 6 - Establish HACCP verification procedures

Principle 7 - Establish effective record keeping and documentation

Twelve Steps to Implement HACCP

1. Assemble HACCP team
2. Describe product
3. Identify intended use
4. Construct process Flow Diagram and Plant Schematic
5. On-site verification of Flow Diagram and Plant Schematic
6. List hazards associated with each step – (HACCP principle 1)

7. Apply HACCP decision tree to determine CCP – (HACCP principle 2)
8. Establish critical limits – (HACCP principle 3)
9. Establish monitoring procedures - (HACCP principle 4)
10. Establish deviation procedures - (HACCP principle 5)
11. Establish verification procedures - (HACCP principle 6)
12. Establish record keeping/documentation for steps 6-11 - (HACCP principle 7)

Step 1 - Assemble HACCP Team: A multi-disciplinary HACCP team comprising of individuals with different specialties is integral for the successful implantation of HACCP in any seafood processing unit. The members of the team include people from maintenance, quality control, production, cleaning and sanitation, people involved in the day- to-day plant operations. The HACCP team has to be led by a team leader who must be well-trained, have a reasonable scientific background and have the ability to motivate and work well with others. The team must have access to reliable technical information. The main duties of the HACCP team are to develop the HACCP plan, verify the HACCP plan, implement and continually revise the HACCP plan to accommodate changes. The team approach is preferred as a single person cannot be an expert on all operations of the processing unit. Moreover, team approach encourages 'ownership' of the decisions.

Steps 2 & 3- Description and Intended Use of Product: The fish products that are to be manufactured have to be described in details with its distribution chain. This provides information on the ingredients, processing methods, and distribution methods (frozen, refrigerated, or at ambient temperature). Identify how the fish are stored after receipt, how the finished product will be shipped, how the finished product will be packaged, how the products are intended to be used and finally identify the intended consumer. The consumers of the food also have to be described thoroughly. These may be the general public or a segment of the population such as infants or the elderly.

Step 4-Construct a process flow diagram: A flow diagram describes the process for each product. This diagram provides a clear, simple outline of the steps involved in the making the product. A block flow diagram is usually sufficient. The flow diagram details all the process activities including inspections, transportation, storage and delays in the process as they “flow” from receipt to distribution.

Step 5- On site verification of the process flow diagram: The HACCP team should perform an on-site review of the operation to verify the accuracy and completeness of the flow diagram. Perform a walk-through of the process to make sure all process steps are covered. It should be done by all members of the HACCP team during all stages and hours of operation

Step 6-Hazard Analysis (HACCP Principle 1): The HACCP team reviews the ingredients used, activities at each processing step, and then makes a list of food safety hazards that are reasonably likely to cause injury or illness if not controlled. Hazard is any biological, chemical or physical agent that may cause an unacceptable health risk to consumers if present in the product. Examples of biological hazards are pathogenic bacteria, viruses, parasites; examples of chemical hazards are heavy metals, pesticides, antibiotic residues, dyes; examples of physical hazards are metal pieces, glass pieces. The hazards may be species related or process related. These food safety hazards may already be present in the raw material or, on the other hand, these may be introduced during the subsequent processing stages, which thus adversely affect the hygiene and safety status of the product. The HACCP team must identify steps in the process where significant hazards may occur, identify the type and nature of contamination, the stage at which it might occur (either during or after processing). The team must estimate both risk and severity of hazards. The risk assessment is based upon experience, epidemiological data and technical information.

Five steps are involved in a hazard analysis:

- i. List process steps
- ii. Identify potential food safety hazards

- iii. Determine if the hazard is significant
- iv. Justify the decision
- v. Identify control measure

The HACCP team should identify the preventive measures for each hazard to reduce probability of risk. Biological hazards are controlled through time/temperature controls, cooking, freezing, fermentation and/or pH controls, adding salt or other preservatives, drying or other processing techniques. Chemical hazards (natural toxins, pesticides, drug residues, unapproved food and colour additives, histamine) are mainly controlled through source controls, time/temperature controls, production controls and labelling controls. Physical hazards (metal, glass) are controlled through source controls and production controls.

Step 7 - Identify the Critical Control Points (CCP's) (HACCP Principle 2): A CCP is a point, step or procedure which can be applied to prevent, eliminate or reduce hazards to an acceptable level. Every significant hazard must have a corresponding CCP and a “decision tree” is used to determine if a process point is a CCP or not.

The decision tree follows a series of questions.

Q1: Does this step involve a hazard of sufficient risk and severity to warrant its control?

Yes – go to Q2

No – Not a CCP

Q2: Does a control measure for the hazard exist at this step?

Yes – go to Q3

No – Is control of step necessary for safety?

Yes – Modify the step process or product

No – Not a CCP – Stop*

Q3: Is control at this step necessary to prevent, eliminate or reduce the risk of the hazard to consumers?

Yes – CCP

No – Not a CCP – Stop*

The examples of CCPs in a fish processing unit

- Reception of raw materials
- Cooking:
- Metal detection point
- Labelling

CCPs are Product and Process Specific. CCPs may change with differences in plant layout, product formulation, process flow, change in equipment, ingredient selection and sanitation and support programs.

Step 8- Establish Critical Limits (HACCP Principle 3): Critical limit is a maximum and/or minimum value to which a biological, chemical or physical parameter must be controlled at a CCP to prevent, eliminate or reduce to an acceptable level the occurrence of a food-safety hazard. Critical limits are boundaries which cannot be exceeded if the hazard is to be prevented, eliminated or minimized. Critical limits are scientifically determined based on information available in journal articles, food science texts, microbiology texts, regulations and guidelines. Examples: Cooking temperature of 99°C for a specified time depending on the product size to attain core temperature of 72°C to control *Listeria monocytogenes*.

Step 9- CCP Monitoring (HACCP Principle 4): Monitoring involves a planned sequence of observations or measurements to assess whether a CCP is under control. These observations are used to determine whether or not corrective action is required. Monitoring facilitates tracking of the operation, indicates when there is a loss of control and a deviation occurs and provides written documentation for use during the verification process. Examples of CCP monitoring are time and

temperature of cooking process, water activity (aw), pH, internal product temperature, salt concentration in brine, metal inclusion screening.

Step 10- Corrective Actions (HACCP Principle 5): Corrective actions are the planned actions that are to be undertaken when monitoring indicates that there is a deviation from an established critical limit. A failure to meet a required critical limit for a critical control point is known as a deviation. There is a need for clear corrective action protocols and chain of command must be emphasized. The corrective actions taken must bring the CCP back under control to ensure that the production process will not cause consumer illness. All the corrective actions taken to fix the problem that caused the deviation and restore process control must be documented and these records will help the firm identify recurring problems. The aim of corrective actions is to re-establish control of the process so that production can start again as soon as possible without further deviations. However, it needs to be emphasized that when critical limit deviations frequently reoccur then, the process and the HACCP plan must be reevaluated. Examples of corrective actions include adding more salt to the brine solution; re-cooking if possible if the internal temperature was not achieved;

Step 11 - Verification (HACCP Principle 6): Verification includes those activities, other than monitoring, that determine the validity of the HACCP plan and that verify the system is operating according to the plan. The purpose of verification is to provide a level of confidence that the plan is based on solid scientific principles and control the hazards associated with the product and process, and is being followed. The type of verifications are Validation, CCP verification activities, HACCP system verification and verification by the regulatory agencies. Validation involves establishing the scientific basis for the HACCP plan. Validation is performed initially and whenever there are changes in raw materials, changes in product or process, recurring deviations, new information on hazards or control measures, new distribution or consumer handling practices. A validation example might be the documentation used to select a cook step to control salmonella in a ready-to-eat product, the minimum time and temperature needed to cook the product,

and the frequency of temperature monitoring to ensure safety. The second part of verification is to “verify” that the validated plan is being followed correctly. Examples of “verifying” include: calibration of process monitoring instruments, direct observation of monitoring activities and corrective actions, and review of records generated and maintained in accordance HACCP Plan. HACCP system is verified annually or whenever there is an occurrence of a system failure or significant change in product or process.

Step 12 - Record Keeping (HACCP Principle 7): Systematic record keeping system with filing system is a vital component of the HACCP system. Records pertaining to HACCP plan and supporting documentation, records of CCP monitoring, records of corrective actions, records of verification activities, sanitation Control records, importer verification records are maintained in fish processing units.

Conclusion

The future of fisheries depends on the production of safe and wholesome products, and this goal can be achieved by the strict enforcement of HACCPbased practices, during primary production /harvest stage, processing stage in fish processing units, distribution to domestic and international markets and storage during retail sale. The aim of quality assurance is to ensure that a product conforms as closely as possible and consistently to that standard at all times.