(HACCP)

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Agenda

- HACCP definition and history
- * Prerequisite programs
- * HACCP components
 - Plan description
 - Process flow diagram
 - * Seven principles
 - Conduct hazard analysis
 - Determine critical control points
 - * Establish critical limits
 - Establish monitoring procedures
 - * Establish corrective action
 - Establish verification procedures
 - Establish record keeping and documentation procedures
- * Q&A

What is HACCP?

- * HACCP is a systematic approach to the identification, evaluation, and control of food safety hazards (biological, chemical, and physical) throughout the production process to prevent the risk of foodborne illness or other safety concerns
- HACCP is based on seven key principles
- * It should be used in conjunction with other food protection programs
- Programs are mandatory for meat & poultry (USDA), seafood (FDA), and juice (FDA) and voluntary for all other food and beverage industries

History

- HACCP was developed in the early 1960's as a collaborative effort between Pillsbury, NASA, and US Army Labs (Natick) to provide safe food for space expeditions
- * Its success propelled implementation across all Pillsbury commercial lines and the food industry in the early 70's

Prerequisite Programs – Foundation for HACCP

- Facility and equipment should be located, constructed and maintained according to sanitary design principles. Traffic control to minimize crosscontamination
- Supplier control to ensure they are adhering to GMP's and food safety programs
- Written specifications for all ingredients, products, and packaging materials
- Procedures (including list of chemicals and dilution/use) for cleaning and sanitation of facility and equipment. Implementation of eight key FDA sanitation conditions 21 CFR, Part 123.11.
- GMP's and personal hygiene
- Chemical segregation and proper use of non-food chemicals in the facility
- Proper conditions for receiving, storage and shipping
- Traceability and recall programs
- Pest control program

Description of Product

- General description of the food, ingredients, and processing methods
- Method of distribution and distribution temperatures
- Expected use (intended for general population or particular segment)
- Primary and secondary packaging
- Shelf life and format
- Label instructions

PRODUCT DESCRIPTION

PROCESS CATEGORY: Fermented food pH < 4.6

PRODUCT: Kombucha. A beverage made from black, green, and/or pu-erh tea (Camellia Sinensis), evaporated cane juice solids, and filtered water that has been fermented (wild) by means of added culture and cellulose pellicle containing primarily yeast species and Gluconoactobacter.

COMMON NAME
FORM/INTENDED USE
Ready to drink refrigerated beverage to be consumed by general public

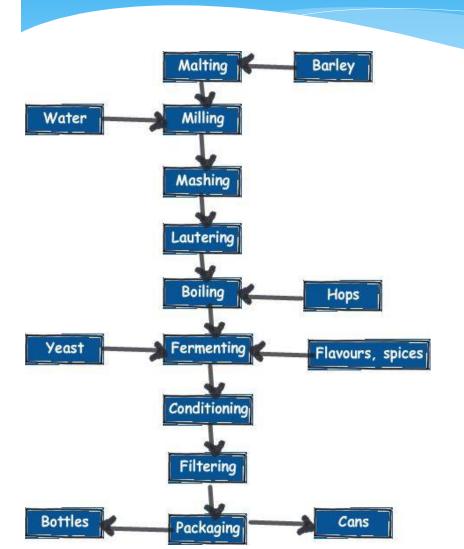
TYPE OF PACKAGE

□ PRIMARY
 □ SECONDARY
 16 fl oz glass bottle; PP closure w/PE foam liner; 2" shrink band
 □ Corrugated shipper (re-shipper) containing 12 bottles

SHELF LIFE 4 months refrigerated

"Best Enjoyed By"; "DD (numeric date) MMM (alpha month) YYYY (numeric year)"

DISTRIBUTION Wholesale to distributors or retailers via refrigerated transport LABELING INSTRUCTIONS Keep refrigerated; do not shake (naturally effervescent)



Process flow diagram (beer example)

Seven Principles

1. Conduct a Hazard Analysis

- * Biological: Bacterial, viral or parasitic pathogens
- * Chemical: Chemical residues
- * Physical: Foreign matter, glass fragments, metal fragments

2. Identify Critical Control Points

* A critical control point (CCP) is a point, step, or procedure in the manufacturing process at which control can be applied and as a result, a food safety hazard can be prevented, eliminated, or reduced to an acceptable level

3. Establish Critical Limits for each Critical Control Point

* A critical limit is the maximum or minimum value to which a biological, chemical or physical hazard must be controlled at a CCP to prevent, eliminate, or reduce to an acceptable level

Seven Principles

4. Establish Critical Control Point monitoring requirements

* Monitoring activities are necessary to ensure that the process is under control at each CCP. Each monitoring procedure and its frequency needs to be listed in the HACCP plan.

5. Establish Corrective Action

Identify actions to be taken when monitoring indicates a deviation from an established critical limit.

6. Establish Verification Procedures and Schedule

 Frequent reviews of plans, verification that its being followed correctly, and review of CCP monitoring and corrective action records

Seven Principles

- 7. Establish record-keeping and documentation procedures
- * Facility must maintain HACCP plan and records documenting the monitoring of CCP's, critical limits, verification activities, and the handling of processing deviations.

Example - Hazard Analysis

| Process Step | Food Safety Hazard | Reasonably | Basis | If yes in Column 3, what | Critical |
|--------------------------|--------------------------------|------------|-------------------------------|---------------------------------|----------------------|
| | | Likely to | | measures could be applied | Control Point |
| | | Occur? | | to prevent, eliminate, or | |
| | | | | reduce the hazard to an | |
| | | | | acceptable level? | |
| Batch Fermentation | Biological - Pathogens | Yes | Pathogens could potentially | Product is maintained at or | 3B |
| | | | grow in this product if pH is | below pH 4.5 at all times; | |
| | | | not maintained below 4.6 | finished product is ≤ pH 3.3 | |
| | Chemical - None | | | | |
| | Physical - None | | | | |
| Filtering | Biological - None | | | | |
| | Chemical - None | | | | |
| | Physical – Foreign matter | No | Gaskets to be visually | | |
| | (stainless steel) from in-line | | inspected before and after | | |
| | screen gasket | | every tank is filtered | | |
| Packaging Materials – | Biological - None | | | | |
| Receiving & Storage | Chemical - None | | | | |
| (glass bottles, | Physical – Foreign matter | Yes | Damage to glass bottles | 100% visual inspection of | 4P |
| closures, shrink bands, | (glass) | | could result in glass | bottles prior to filling during | |
| re-shippers | | | inclusion | label application | |
| Bottle filling & capping | Biological – None | | | | |
| | Chemical – None | | | | |
| | Physical – Foreign matter | Yes | Damage to glass bottles | 100% visual inspection of | 5P |
| | (glass) | | could result in glass | bottles prior to filling and | |
| | | | inclusion | capping during filling step | |
| Shrink banding & case | Biological – None | | | | |
| packing | Chemical - None | | | | |
| | Physical - None | | | | |
| Refrigerated storage | Biological - None | | | | |
| and distribution | Chemical - None | | | | |
| | Physical - None | | | | |

Example - CCP's

| | Monitoring | HACCP Records | Verification Procedures and | Corrective Action |
|----------------------------|--|---|--|---|
| | Procedures and | | Frequency | |
| | Frequency | | | |
| | | | | |
| | | | | |
| Must maintain pH < 4.6 | Test pH of each batch | · · | | If pH exceeds pH 4.5, or |
| | - | product record | | discard contents of tank |
| | prior to bottling | | after each batch is bottled | |
| | | Corrective action log | | |
| | | | · · · · · · · · · · · · · · · · · · · | |
| | · · | • | - | Dispose of all broken or |
| cracks in glass containers | · · | record | | compromised glass |
| | | Causatius astisulas | bottles for each production run | bottles. Inspect others |
| | application | Corrective action log | | for fragments or similar |
| Nie byseliese skies von | 1000/ | Class in an astion | | defects. |
| | - | · · | • | Dispose of all broken or |
| cracks in glass containers | | record | | compromised glass bottles. Inspect others |
| | and manual capping | Corrective action log | | for fragments or similar |
| | | Corrective action log | Tull | defects. Inspect bottling |
| | | | | area of production room |
| | | | | and remove any glass |
| | | | | fragments. |
| (| Must maintain pH < 4.6 No breakage, chips, nor cracks in glass containers No breakage, chips, nor cracks in glass containers | Must maintain pH < 4.6 Must maintain pH < 4.6 Test pH of each batch of finished product prior to bottling No breakage, chips, nor cracks in glass containers 100% visual inspection of bottles during manual label application No breakage, chips, nor 100% visual inspection | Must maintain pH < 4.6 Test pH of each batch of finished product prior to bottling No breakage, chips, nor cracks in glass containers No breakage, chips, nor cracks in glass containers Test pH of each batch of finished product record Corrective action log Glass inspection record Corrective action log Corrective action log Corrective action log Glass inspection record Corrective action log Obstreakage, chips, nor cracks in glass containers Obstreakage, chips, nor cracks in glass containers Obstreakage, chips, nor cracks in glass containers | Must maintain pH < 4.6 Must maintain pH < 4.6 Test pH of each batch of finished product prior to bottling No breakage, chips, nor cracks in glass containers No breakage, chips, nor cracks in glass containers Test pH of each batch of finished product product record product record will be verified and filed after each batch is bottled Corrective action log Glass inspection record will be verified and filed after labeling bottles for each production run Corrective action log No breakage, chips, nor cracks in glass containers Obstiles during manual label application Glass inspection record will be verified and filed after labeling bottles for each production run Glass inspection record will be verified and filed after labeling bottles for each production record will be verified and filed after filling and case packing for each production |

Example – CCP's Logs

pH Meter Calibration Log

| Date | pH Meter | Calibrate to pH 2.0 | Calibrate to pH 4.0 | Technician Initials |
|------|----------|---------------------|---------------------|---------------------|
| | | | | |
| | | | | |
| | | | | |
| | | | | |
| | | | | |
| | | | | |

CCP 4P and 5P - Glass Inspection Record

| Date | Labeling 4P | Filling & Case Packing 5P | Quantity | Comments and/or Corrective Action |
|------|----------------|---------------------------------|----------|-----------------------------------|
| | | | | |
| | | | | |
| | | | | |
| | | | | |

Helpful guides and examples:

Key sanitation conditions

http://www.qualityassurancemag.com/Article.aspx?article_id=101570

HACCP plans

www.fda.gov/Food/GuidanceRegulation/HACCP/ucm2006801.htm

www.beercanada.com/beer-canadahaccp-food-safety-program

http://www.apiservices.com/articles/us/haccp_en.htm