# Identification of requirements for implementing Food Safety Management System (ISO 22000:2005) to the carbonated beverage process at Shaa Cola Beverages (Pvt) Ltd.

By

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- Thesis submitted in partial fulfillment of the requirements for the

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Petruny, 2009

# DECLARATION

The work described in this thesis was carried out by me at the Department of Food Science & Technology, Faculty of Applied Sciences, Sabaragamuwa University of Sri Lanka, under the supervision of Mr. A.L.C.J. Liyanage, Lecturer, Department of Food Science & Technology, Faculty of Applied Sciences, Sabaragamuwa University of Sri Lanka & Mr. A. Wanasinghe, Operational Manager, Shaa Cola Beverages (Pvt) Ltd, Moragahahena, Horana. A report on this has not been submitted to any other university for another degree.

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My Parents, Teachers

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My Alma mater

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## ABSTRACT

As food organizations which thrive in competitive market context, product quality is one of the most important factor to be considered at strategic decision making process. Thus the food safety management system has become necessary rather than the quality management system which is widely implemented in all sectors but does not itself specifically address food safety. ISO 22000:2005 is therefore designed to allow all types of organizations within the food chain to implement a food safety management system.

The present study was focused on the setting up of preliminary requirements for the implementation of food safety management system (ISO 22000:2005) to the carbonated beverage process at Shaa cola beverages (Pvt) Ltd.

As the first step; scope, food safety policy and related objectives were defined and Prerequisite Programs (PRPs) were developed by studying the carbonated beverage production flow beginning from raw material reception to transportation of the finished product. Critical Points (CPs) and Critical Control Points (CCPs) were determined by identifying all the food safety hazards associated with production process and assessing those hazards using risk assessment matrix and decision tree. Then Operational Prerequisite Programs (OPRPs) and Hazard Analysis Critical Control Point (HACCP) plan were established and documented. Process capability analysis was done for validation of food safety management system. Verification and continuous improvement procedures of food safety management system were also developed.

PRPs were developed for calibrations, cleaning and sanitizing, personal hygiene, pest control, supplier assessment, waste and sewage disposal and water, air and energy supply. Raw material receiving, ingredient weighing, sugar dissolving, sugar filtration, water filtration, carbonation and finished product storage were identified as CPs and raw material receiving and inspection, syrup room management, carbonated process management, filtration process management and finished product inspection were established as OPRPs to monitor the CPs. UV sterilization and bottle capping were identified as points that need to be critically controlled and HACCP plan was developed. Addressing those according to the process barely acceptable processes and soda as well as lemonade production processes should be improved according to their specifications. Food safety plan and internal audit plan were developed as verification procedures of food safety management system.

it is further expected to conduct a gap analysis to evaluate the successful implementation of **NO 22000**:2005 to the carbonated beverage process.

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# LIST OF ABBREVIATIONS

ADI	Acceptable Daily Intake
APC	Aerobic Plate Count
CPs	Critical Points
CCPs	Critical Control Points
CIP	Cleaning In Place
CLs	Critical Limits
Codex	Codex Alimentarius Commission, an FAO/WHO Organization
EEC	European Economic Community
ESCF	European Standing Committee on Food
FAO	Food and Agricultural Organization
FDA	Food and Drug Administration
FIFO	First In First Out
FSMS	Food Safety Management System
GRAS	General Recognize As Food
HACCP	Hazard Analysis Critical Control Points
ISO	International Organization for Standardization
ISO/TS	International Organization for Standardization/Technical Specification
JECFA	Joint Expert Committee on Food Additives
MR	Management Representative
MRM	Management Review Meeting
NCR	Non Conformity Records
OPRPs	Operational Prerequisite Programs
PET	Poly Ethylene Tetraptheline
PRPs	Prerequisite Programs
QAM	Quality Assurance Manager
SLS	Sri Lankan Standards
SQA	Supplier Quality Assurance
ŬV	Ultra Violet
WHO	World Health Organization

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# CHAPTER 01 Introduction

#### 1.1 Introduction

In order to complete the global picture, food organizations in competitive market context, the product quality is one of the most important factor to be consider at strategic decision making process. Food reaches consumers via supply chains that may link many different types of organization and that may stretch across multiple borders. One weak link can result in unsafe food that is dangerous to health and when this happens, the hazards to consumers can be serious and the cost to food chain suppliers considerable. As food safety hazards can enter the food chain at any stage, adequate control and communication throughout is essential. Food safety is a joint responsibility of all the actors in the food chain and requires their combined efforts.

Further development of Socio-economical, Educational and changes in consumer habits forces the food companies to produce safe products that give minimum harm effect to its consumers. The standard has become necessary because of the significant increase of illnesses caused by infected food in both developed and developing countries. In addition to the health hazards, food borne illnesses can give rise to considerable economic costs covering medical treatment, absence from work, insurance payments and legal compensation.

Since industry develops, the International Organization of Standardization (ISO) standards become an essential tool to reach the consumers as they more aware of those standards and may tend to choose them more. The food safety management system (ISO 22000:2005 standard) has become necessary rather than the quality management system (ISO 9001:2000) which is widely implemented in all sectors but does not itself specifically address food safety.

ASO 22000 is therefore designed to allow all types of organization within the food chain to implement a food safety management system. These range from feed producers, primary producers, food manufacturers, transport and storage operators and subcontractors to retail and food service outlets together with related organizations such as producers of equipment, packaging material, cleaning agents, additives and ingredients.

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ISO 22000 combines the Codex Alimentarius HACCP (Hazard Analysis and Critical Control Points) principles and application steps, developed by Codex Alimentarius, with prerequisite programmes. The development of ISO 22000 was based on the assumption that the most effective food safety systems are designed, operated and continually improved within the framework of a structured management system, and incorporated into the overall management activities of the organization. It is applicable to all organizations, regardless of size, which are involved in any aspect of the food chain and want to implement systems that consistently provide safe products (Faergemand and Jespersen, 2004).

Carbonated beverage is an effervescent drink that releases carbon dioxide under conditions of normal atmospheric pressure. Production of carbonated soft drinks started with pure source of water and carbon dioxide, flavours, preservatives, colours are use for complete the production. Most of the people tend to consume carbonated beverages due to its thirst quenching property (Mitchell, 1990).

Shaa Cola Beverages (Pvt) Ltd is one of the subsidiary company of Jaycey group and it was set up in year 2006. Company engaged in the process of manufacturing range of carbonated beverages including soda, cream soda, cola, mandarin, tamarind, lemonade, etc. which are marketed under the brand name "Shaa". They incorporate some of the fully automated modern machinery with high technology, such as carbonation, filling and capping machines.

At present, Shaa Cola Beverages (Pvt) Ltd is trying to become a Sri Lanka's market leader in the carbonated beverage industry by supplying safe food products with an affordable price to the market. To achieve the company goal, implementing food safety management system (ISO 22000:2005) is a mandatory requirement due to its combination of HACCP (Hazard Analysis Critical Control Points) principles and application steps with prerequisite programmes.

This project work attempts to set up preliminary requirements for implementation of ISO 22000:2005; food safety management system to the carbonated beverage process in line with its existing processes.

## 1.2 Overall objective:

Setting up of preliminary requirements for the implementing Food Safety Management System (ISO 22000:2005) to the carbonated beverage process.

## 1.3 Specific Objectives:

1. To define the scope, food safety policy and related objectives associated with the company.

2. To develop Prerequisite programs (PRPs) for the company.

3. To identify of all potential food safety hazards associated with the carbonated beverage process and establish Operational prerequisite program (OPRPs) and Hazard Analysis Critical Control Point (HACCP) plan and there by specifying control measures.

4. To prepare documentation for the implementation of Food safety management system (ISO22000:2005) to the carbonated beverage process.

# **CHAPTER 02**

## Literature Review

## 2.1 General Overview

### 2.1.1 International Organization for Standardization

The International Organization for Standardization (ISO) is a worldwide federation of national standards bodies (ISO member bodies). The work of preparing International standards is normally carried out through ISO technical committees. Draft International Standards adopted by the technical committees are circulated to the member bodies for voting. Publication as an International standard requires approval by at least 75% of the member bodies casting a vote (ISO 22000:2005(E)).

#### Increased demand for International Standards due to;

- Globalization of trade in products and services.
- Outsourcing and foreign investment.
- Deregulation/privatization of public services.
- The climate change challenge and energy efficiency mandates.
- Public demand for consumer safety, environmental protection, corporate social responsibility.
- Need for international solidarity to face global terrorism, pandemics and natural disasters.
- Pace of innovation and convergence of new technologies.

### 2.1.2 ISO 22000

ISO 22000 does not follow an exhaustive and prescriptive checklist approach; instead the standard allows an organization to develop a food safety management system tailored to its particular suppliers, customers and relevant parties.

### 2.1.3 ISO 22000 family of standards

Four Standards so far:

ASO 22000:2005 - Food safety management system - Requirements

ISO 22003:2007 - Requirements for bodies providing audit and certification of food safety management systems

ISO TS 22004:2005 - Guidance on the application of ISO 22000

ISO 22005:2007 - Traceability in the feed and food chain

(Makialey and Chambers, 2007).

#### 2.1.4 ISO 22000:2005 - Food Safety Management System - Requirements

ISO 22000:2005 specifies requirements for a food safety management system where an organization in the food chain needs to demonstrate its ability to control food safety hazards in order to ensure that food is safe at the time of human consumption. The standard builds on generally accepted principles for food safety in the food chain and focuses on identifying and preventing food safety hazards through PRPs (Pre-Requisite Programs) and HACCP (Hazard Analysis and Critical Control Points) plans. It promotes communication, continual improvement and management and updating of the food safety management system (ISO 22000:2005(E)).

#### 2.1.5 Benefits of ISO 22000:2005

**Obtaining ISO 22000:2005** compliance will yield the following benefits to the company.

- A systematic approach to identification of food safety hazards, which in turn aids in the development and implementation of proactive control measures.
- Marketing advantage increased business
- Improved efficiency and profitability
- Increased customer satisfaction
- Documented system provides useful reference
- Improved records in case of litigation
- Responsibilities of personnel clearly defined
- Improved control during periods of change or growth
- Improved communication on food safety issues
- Improved performance from suppliers
- Increased customer satisfaction
- Closes the gap between ISO 9001 and HACCP
- Meets food industry expectations
- Based on system management, not on inspections or a product approach
- Provides an opportunity for international recognition through 3<sup>rd</sup> party registration
- Makes savings from fewer customer audits possible
- Provides a baseline for the entire food chain (Lab alert: ISO 22000)

#### 2.2 Key elements of ISO 22000:2005

ISO 22000 identifies the following four 'pillars' of a food safety management system.

#### 2.2.1 Interactive communication

Communication along the food chain is essential to ensure that all relevant food safety hazards are identified and adequately controlled at each step within the food chain. This implies communication between organizations both upstream and downstream in the food chain. Recognition of the organization's role and position within the food chain is essential to ensure effective interactive communication throughout the chain in order to deliver safe food products to the final consumer (ISO 22000:2005(E)).

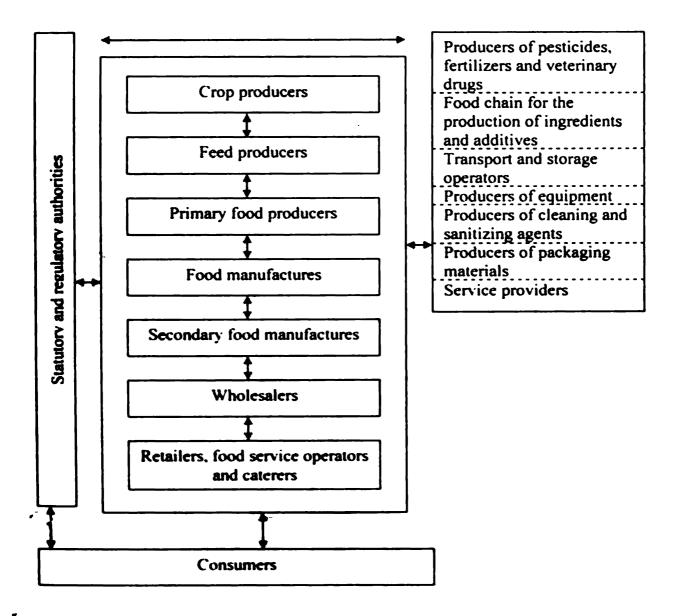


Fig. 2.1: Model of communication within the food chain (Source: ISO 22000:2005(E))

#### 2.2.2 System management

The most effective food safety systems are established, operated and updated within the framework of a structured management system and incorporated in to the overall management activities of the organization. This provides maximum benefit for the organization and interested parties. Food safety management system has been aligned with ISO 9001 quality management system in order to enhance the compatibility of two standards.

#### 2.2.3 Prerequisite programs

Prerequisite programs (PRPs) must be in place and fully operational before the HACCP is in place .The PRPs needed depend on the segment of the food chain in which the organization operates and the type of organization. These program must be implement adequate and effectively in the process. By application of those able to control the number of CPs and CCPs occur in the process unnecessarily. Some examples are personal hygiene, cleaning and sanitation, waste and sewage disposal etc (ISO 22000:2005(E)).

#### 2.2.4 HACCP Principles

**ISO 22000:2005** is based on the Hazards Analysis and Critical Control Point (HACCP) principles that have international acceptance and details of the approach have been published by the Codex Alimentarius Commission to identify, prevent and control food safety hazards.

Principle 1: Conduct a hazard analysis

Principle 2: Identify Critical Control Points (CCPs)

**Principle 3: Establish critical limits** 

Principle 4: Establish CCP monitoring requirements

Principle 5: Establish corrective actions

Principle 6: Establish record keeping

Principle 7: Establish procedures for verification to determine the effectiveness of HACCP system (Mortimore and Wallace, 1998).

#### 2.3 ISO 22000:2005 as a process approach

For an organization to function effectively and efficiently, it has to identify and manage numerous linked activities. An activity using resources, and managed in order to enable the transformation of inputs in to outputs, is considered as a process. Often the output from one process directly forms the input to the next.

The application of a system of processes within an organization, together with the identification of interactions and the management of these processes can be referred to as the "process approach". An advantage of the process approach is the ongoing control that it provides over the linkage between the individual processes within the system of processes as well as their combination and interaction.

When used within a food safety management system, such an approach emphasizes the importance of

- Understanding and fulfilling the requirements
- The need to consider processes in terms of food safety and traceability
- Obtaining results of process performance and effectiveness and
- Continual improvement of processes based on objective measurement

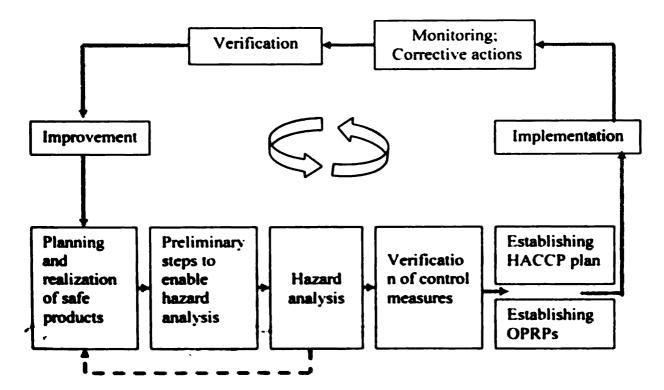


Fig. 2.2: The model of a process-based food safety management system (Source: ISO/TS 22004:2005(E))

#### 2.4 ISO 22000:2005 - Food safety management system

#### 2.4.1 Requirements

#### 2.4.1.1 General requirements

The organization defines the scope of the food safety management system that is specify the product or product categories, process and production sites that are address by the food safety management system. Also ensure that food safety hazards are identified, evaluated, controlled in such a manner, communicate appropriate information.

#### 2.4.1.2 Documentation requirements

A food safety management system needs to be documented. The organization must have, as a minimum, a written food safety policy and related objectives, the procedures and records required by ISO 22000 and any other documents that need to ensure the effective development, implementation and updating of the system.



Fig. 2.3: Documentation requirements hierarchy of food safety management system (Source: ISO 22000: UNCTAD/WTO 2007)

#### 2.4.2 Management responsibility

#### 2.4.2.1 Management commitment

Top management's commitment is critical to the successful implementation of the food safety management system. The system costs time and money to develop, implement and maintain. It will directly involve top management it self in allocating resources and in decision making at key steps in the process, its maintenance and updating.

#### 2.4.2.2 Food safety policy

The overall food safety policy should be short and tightly focused, because it is a key benchmark for food safety management system. So top management needs to be involved in its development and clearly committed to its implementation. Food safety policy should speak directly to employees, as a vital part of training.

#### 2.4.2.3 Food safety management system planning

The plan should identify the business objectives to be met by the system, set out the resources required and identify measurable objectives. Therefore planning is critical to the development, implementation and updating of a food safety management system by top management.

#### 2.4.2.4 Responsibility and authority

ISO 22000 statuses that top management must ensure that the specific responsibilities and authorities needed for the establishment of the food safety management system are formally defined and communicate to all employees within the organization. The process of establishing relevant job descriptions contributes to the achievement of specific food safety management system objectives that were set for relevant functions and levels within the organization.

#### 2.4.2.5 Food safety team leader

**ISO 22000 statuses that the responsibility and authority for the food safety management system must be assigned to one person.** ISO 22000 identifies that person **as the food safety team leader, appointed by top management.** The team leader should **be a member of the organization and have a basic knowledge of hygiene management, the application of HACCP principles and the requirement of ISO 22000.** 

#### 2.4.2.6 Communication

Communication should ensure that the necessary interactions occur within the organization and along the food supply chain. External communication should ensure that any relevant hazards are controlled at the appropriate steps in the food chain. Internal communication is just as important as external communication in ensuring ' that the organization's system is effective.

#### 2.4.2.7 Emergency preparedness and response

ISO 22000 says top management must be prepared to respond to potential food emergency situations and accidents that can impact on food safety. These can include incidents such as fire, flooding, bio-terrorism and sabotage, energy failure, vehicle accidents, contamination of the environments and various types of weather related events.

#### 2.4.2.8 Management review

Management must review the food safety management system to ensure its continuing suitability, adequacy and effectiveness. The review also facilitates the assessment of opportunities for improvement and the need for change to the system, including the food safety policy. This review should take place at planned intervals. In the early stages of implementation, these intervals may be shorter than when the system is mature.

#### 2.4.3 Resource management

A variety of resources will be required by every organization establishing, implementing, maintaining and updating a food safety management system. These include human resources, infrastructure and an appropriate work environment.

#### 2.4.4 Planning and realization of safe products

To realize safe food products, an organization needs to plan and develop the necessary processors. These will include prerequisite programmes (PRPs), operational prerequisite programmes (OPRPs) and the HACCP plan.

#### 2.4.4.1 Prerequisite programmes (PRPs)

PRPs are the basic conditions and activities that are necessary to maintain a hygienic environment throughout the food chain suitable for the production, handling and provision of safe end products. When implemented PRPs assist in controlling the likelihood of introducing food safety hazards, biological, chemical or physical contamination of products and any increase in the levels of these hazards in the product or the processing environment. They need to be appropriate to the products and processes of the organization, approved by the food safety team and implemented.

#### 2.4.4.2 Hazard analysis

The development of a food safety management system involves the rigorous application of a hazard analysis. This process involves a number of preliminary steps and rigorous decision making process. The steps and principles involved in this process have been outlined by the Codex Alimentarius Commission (ISO 22000:2005(E)).

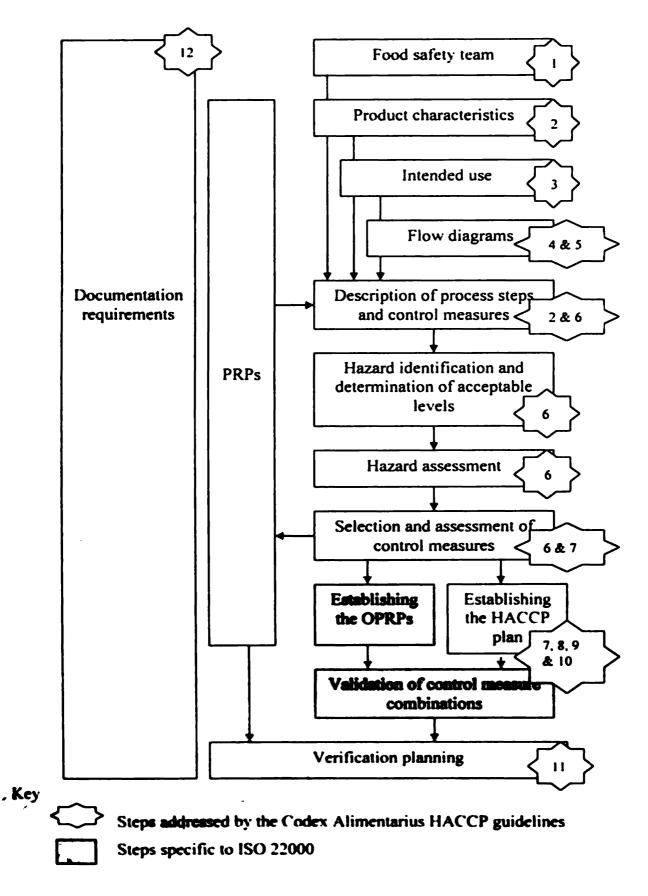


Fig. 2.4: Steps of the ISO 22000 approach to hazard analysis (Source: ISO/TS 22004:2005(E))

#### Preliminary steps to enable hazard analysis

• Food safety team

The food safety team is a combination of multi-disciplinary, knowledge and experience in developing and implementing the food safety management system.

• Product characteristics

All raw materials, ingredients, product contact materials and characteristic of end products are described to the extent needed to conduct the hazard analysis.

Intended use

Intended uses are group of uses or appropriate group of consumers for each product. They are end uses, general public, sensitive population, infant, pregnant mothers, elderly people, sick people, etc.

• Flow diagrams

Flow diagrams are prepared for the products or process categories covered by the food safety management system. It is a clear, accurate and sufficiently detailed diagram providing a basis for evaluating the possible occurrence, increase or introduction of food safety hazards (Boutou, 2008).

#### 2.4.4.3 Hazard identification and determination of acceptable levels

Food safety hazard: A biological, chemical or physical agent in food, or conducting of food with a potential to cause an adverse health effect.

- Biological hazards: pathogenic bacteria, fungi, viruses, parasites, etc.
- Chemical hazards: Intentional additives:- colours, flavours, preservatives
   Unintentional additives: cleaning chemicals
   Natural toxins: seafood, mushrooms

• Physical hazards: foreign matters such as glass, stones, jewelleries, metals, etc Acceptable level means the level of an identified hazard in the end product of the organization that is needed at the next step in the food chain to ensure food safety; it refers to the acceptable level in foods for direct consumption only when the next step is actual consumption (Mortimore and Wallace, 1998).

#### 2.4.4.4 Hazard assessment

A hazard assessment is conducted to determine, for each food safety hazard identified whether its elimination or reduction to acceptable levels is essential to the production of a safe food, and whether its control is needed to enable the defined acceptable levels to be met.

#### 2.4.4.5 Establishing operational prerequisite programmes (OPRPs)

**OPRP is a Prerequisite programme identified by the hazard analysis as essential in order to control the likelihood of introducing food safety hazards to and/or the contamination or proliferation of food safety hazards in the product(s) or in the processing environment.** 

2.4.4.6 Establishing HACCP plan

HACCP plan is a document prepared in accordance with the HACCP principles to ensure the control of hazards, which are significant for food safety in the segment of the food chain under consideration.

# 2.4.5 Validation, verification and improvement of the food safety management system

Validation

Validation is obtaining evidence that the control measures managed by the HACCP plan and by the Operational PRPs are effective and capable of achieving the intended control of the food safety hazard.

#### • Control of monitoring and measuring

It is a procedure to detect any failures in the control measures. Organization provides evidence that the specified monitoring methods and measuring equipments are adequate to ensure the expected performance.

#### Verification

Verification is an assessment carried out during and after the operation, to demonstrate that the intended level of control is actually been achieved. Organizations are carried out internal audit programs at planned intervals for verification process

#### Improvements

Top management ensure that the organization continually improves the effectiveness of the food safety management system through the use of communication, management review, internal audit, etc (ISO 22000:2005(E)).

#### **2.5 Carbonated Beverages**

**Carbonated nonalcoholic beverages are generally sweetened, flavored, acidified, coloured, artificially carbonated and some times chemically preserved beverage that contains the characteristic of carbonation**.

Their origin goes back to Greek and Roman times when naturally occurring mineral waters were prized for 'medicinal' and refreshing qualities. But it was not until about 1767, when the British chemist Joseph Priestley found that he could artificially carbonated water, that the carbonated beverage industry got its start. An early method of obtaining the carbon dioxide was by acidification of sodium bicarbonate or sodium carbonate, and from the use of these sodium salts came the name "soda" which remains today, although most carbon dioxide is no longer generated in this fashion. Generally, fruit juices and extracts were added to carbonated water for improved flavour (Hotchkiss and Potter, 1996).

#### 2.5.1 Raw materials

#### • Water

Water is the principal ingredient in carbonated beverage production, and regular soft drinks contain 90% water while diet soft drinks may contain up to 99% water. Water must be free from suspended matters, colouring matters, objectionable odours and minerals which may interfere with the flavour and colour of the soft drink. The alkalinity of beverage water must be low to prevent neutralization of the acid used in the beverage, which would alter flavour and decrease the preservative property of the beverage.

#### • Sugar

Sugars used in carbonated soft drinks can be divided in to those in a dry, granular form and those in a liquid or syrup form or in certain circumstances, wheat (glucose syrup/high fructose glucose syrup). Granulated sugar is a dry crystallized extracted from sugar beet and sugar cane called sucrose that gives the caloric value to the bryerage.

#### Carbon dioxide

Carbon dioxide  $(CO_2)$  is a colourless gas with a slightly pungent odour; when dissolved in water the resultant carbonic acid mixture has an acidic and biting taste which is not unpleasant. Also  $CO_2$  is one of the very few gasses suitable for providing the effervescence in soft drinks. It is non-toxic, inert, virtually tasteless, and readily available at moderate cost and may be liquefied at reasonable temperatures and pressures, allowing convenient bulk transportation and storage.

#### Flavourings and emulsions

Natural flavourings, nature identical flavourings and artificial flavourings are used as flavouring agents for soft drink production. The type of flavouring agent used will depend upon the finished drink required. Although flavourings are normally used in extremely small quantities in a carbonated soft drink, their impact can make the difference between a tasty product and one that is bland and uninteresting. If follows, therefore, that many flavourings are highly concentrated and their application dose rate must be optimized with great care.

As emulsion is a dispersion of one liquid in another in which it would not normally be miscible. The main use of emulsion in carbonated soft drinks is as a clouding agent. When considering the organoleptical features of a soft drink, the visual appearance is extremely important as the 'eye appeal' can often be the determining factor in deciding which drink is actually purchased.

#### • Acids, colours, preservatives and other additives

The main reasons for acidification are, to enhance beverage flavours (sharpness to the background taste), to enhance to the thirst quenching experience by stimulating saliva flow and to act as a preservative against microbial growth. The principle acids used are phosphoric, citric, fumaric, tartaric and malic acids. Among this citric acid being the most widely used.

Some important colouring agents for carbonated beverages are the synthetic colours which have been approved by FAD or natural colours. Colours affect the psychological impression of beverage. Caramel colour (E150) from heated sugar (natural colour) is commonly used in dark beverages such as colas. Sunset yellow (E110), tartezine (E102) like synthetic colours use for orange, lemon like carbonated beverage production.

A preservative can be defined as any substance that is capable of inhibiting, retarding or arresting the growth of micro-organisms or any deterioration of food due to microorganisms or of masking the evidence of any such deterioration. Most commonly use sodium benzoate, sodium citrate, potassium sorbate, ascorbic acid like preservatives

for beverage production.

PERMANENY REFERENCE Sabaragamuwa University Library

#### • PET bottles

PET bottles have provided the carbonated beverage industry with a package which is able to deliver to the consumer volumes of product above the glass bottle's practical limit of 1 litre in a single container and have therefore contributed strongly to the overall growth in the industries volume in the 1980s. It is rare for PET bottles to break and even when breakage does occur, no sharp fragments are produced. The light weight of PET bottles is also an obvious attraction (Mitchell, 1990).

#### 2.5.2 Production process

#### 2.5.2.1 Water treatments

Beverage manufacturer pay particular attention to the quality of the water used in their formulation, to ensure that when the product arrives at the consumer it is consistent in taste and appearance and has not deteriorated during storage. Some of the impurities found in water can have a significant effect upon the quality of the final carbonated product.

Impurity	Effect on product	Treatment process
Suspended particles	Foaming at the filler.	Filtration or coagulation
	Loss of carbonation.	and filtration
	Visible particles in product.	
Organic matter	Deposits or precipitation in	Coagulation and
	product. Formation of neck	chlorination.
	rings during storage.	
Bacteria	Spoilage by creation of off	Chlorination or Ultraviolet
	tastes. Great danger of health	radiation
	risks.	
Taste and odour due	Off-tastes	Chlorination or absorption
to organic compounds		by activated carbon
High alkalinity	Neutralizes acidity. Creates a	Coagulation, Dealkalization
-	bland taste	by iron exchange
Nitrate	Health hazard	Iron exchange
polysaccharides	Precipitation in storage	Reverse osmosis

Table 2.1: Effect of impurities in water on the quality of the product.

(Source: Mitchell, 1990)

#### • Water filtration

Normally use sand filters and activated carbon filters. Fine particles of floc will be removed by filtration through a sand filter. The activated carbon will be housed in a pressure filter which is similar in construction and arrangement to the sand filter. Using the activated carbon filter will remove all the free chlorine from the water and also remove other organic molecules.

• Sterilization

Most food and beverage industries use ultraviolet (UV) sterilized water that is acceptable for their process applications. The advantage of UV sterilization is that it is immediate, and as it does not require a long incubation period. The wave length of the ultraviolet rays can be selected so that not only are bacteria eliminated but naturally occurring organic matter can also be destroyed. The sterilization reaction is very rapid, meaning that the plant is compact and may be sited close to the point of use, ensuring that the only water that is sterilized is that to be used for beverage production (Mitchell, 1990).

#### 2.5.2.2 Sugar dissolving and filtration

The operation of sugar dissolving system requires a quantity of water to be added to the vessel before the addition of the granulated sugar such that, after dissolution, a brix slightly higher than required is produced. After assessing the precise concentration of the liquid sugar produced, dilution with water is then carried out to give the required level. It is prudent to initially 'over-brix' in this manner because of the practical difficulties involved in adding extra sugar if too dilute a liquid sugar is produced.

The levels of extraneous matter in dissolved sugar are of particular concern because of their effect on the appearance of the final products and the possibility of loss of carbonation if excessive sites of nucleation are present. Filtering before dispatch thus minimizes the above problem. Micro-filtration is a safe and efficient method of removal of unwanted particles and other turbidity components.

#### 2.5.2.3 Mixing ingredients

The dissolved sugar and flavour concentrates are pumped and carefully mixed (too much agitation can cause unwanted aeration). The syrup may be sterilized while in the tanks, using ultraviolet radiation or flash pasteurization which involves quickly heating and cooling. The water and syrup are carefully combined by sophisticated

machines called proportioners which regulate the flow rates and ratios of the liquid. Fixed proportions of syrup and water then pass to the carbonator.

#### 2.5.2.4 Carbonation, filling, capping, labeling and packing

Carbonation is the impregnation of a liquid with carbon dioxide gas  $(CO_2)$ . When applied to a soft drink product, the result is a beverage which sparkles and foams as it is dispends and consumed. This escape of  $CO_2$  during consumption of the drink should complement and enhance the flavour, and will add an exciting tingle that stimulates the palate.

Each soft drink formulation requires a particular degree of carbonation. So that the effervescence is appropriate to the flavour and nature of the beverage. Fruit drinkssuch as orange, bitter lemon, etc. - should contain low levels of carbonation where as juice based drinks- colas, ginger beer and cream soda – should be in the medium to high range of  $CO_2$  content.

Normally filling done by automated filling machine. When filling the product is transferred in to glass bottles or PET bottles or cans at extremely high flow rates. The containers are immediately sealed with pressure resistant closures. Then bottles were sprayed with warm water and dry well because soft drinks are generally cooled during manufacturing process. Finally labels are fixed and containers are packed in to cartons for distribution.

#### 2.6 Micro-organisms and soft drinks

Because of their utilization of sugars to produce alcohols and carbon dioxide, yeasts are of more immediate concern. Yeasts, moulds and bacteria can bring about deterioration in flavour, producing taints, off-notes, differences in mouth feel, etc. most yeast can grow with or without oxygen, where as some bacteria cannot survive in it. Most yeast thrive at temperatures between 25 and 27 °C; some can survive over 70 °C and others can exist, apparently quite comfortably, at 0-10 °C. Bacteria exhibit similar characteristics, with the optimum growth temperature being 37 °C.

Micro-organisms require nutrient media to enable growth. The range depends upon the individual organisms but apart from water, as the environmental necessity, typical requirements are sources of carbon (carbohydrates), nitrogen (amino acids), phosphorus (phosphates), potassium, magnesium, calcium (mineral salts) and traces of other minerals, e.g. sulphur, iron, cobalt and even vitamins. Because of the obvious link with protein formation during cell growth, the presence of combined nitrogen is of particular importance. Also where introduced to beverages via fruit pulp or caramel (colouring), there will be greater susceptibility to spoilage by certain micro-organisms.

It is useful to consider the individual status of a bottled drink. It exists as a unique system that can enhance or inhibit the growth of micro-organisms. Micro-flora, which may be present, will enter a dormant stage while the chances of survival are 'assessed'. Following this lag stage, during which specific micro-flora adapt to their new environment and start to grow, there is a burst of activity during which the population will double it self repeatedly at a steady rate. Waste products and diminishing nutrients will act to slow down this growth.

However, in practical terms the drink, while not necessarily a health hazard, no longer exhibits those qualities intended by its formulation (Mitchell, 1990).

#### 2.7 Carbonated Beverages and Food safety issues

The commercial success of a soft drink formulation depends upon a number of factors such as appearance, taste like things. Not only that but also safety of the drink is very important factor. To produce safe carbonated beverage product, each and every ingredient should have a typical use level (Table 2.2).

Component	Typical use level	Contribution
Water	Up to 94% v v	Bland carrier solvent for other ingredients
Sugars	7 - 12 % m/v	Sweetness, body, balance to flavour
Carbon dioxide	0.3 - 0.6 % m/v	Provides mouth-feel and 'sparkle' to drink
Acids	0.05 - 0.30 % m/v	Sourness, sharpness background to flavour; increases thirst quenching effect
Flavours Nature identical	0.1 - 0.28 % m/v	Provides flavour character and identity to the drink
& artificial	-	
Natural	up to 0.5% m/v	
Emulsion	0.1% v/v	Produces cloudiness
Colour	0-70 ppm	Standardizes and identifies colour tone of

		drink	
Prescrvatives	Statutory limits	Restrict microbial attack and prevent	
	generally apply,	destabilization	
	e.g.; SO <sub>2</sub> 70 ppm,		
	benzoic 160 ppm		
Anti-oxidants	< 100 ppm	Limits flavour and colour deterioration owing to oxidation	
Vitamins/minerals	ADI applies	Nutritional requirements	

(Source: Mitchell, 1990)

Both EEC and FDA have published list of colours that are under regular review. Some colours are shown in Table 2.3.

Table 2.3: Some colours used in soft drinks
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Colour	E number
Carmine	E120
Caramel	E150
Quinoline Yellow	E104
Tartrazine	E102
Sunset Yellow	E110
Ponceau 4R	E124

(Source: Mitchell, 1990)

Defined maximum levels for named preservatives are given according to the food substrate; in the case of soft drinks for consumption without dilution, the UK regulations read as follows (Table 2.4).

 Table 2.4: maximum levels of preservatives

mg/kg	E number
70	E220
160	E210
160	E214
160	+E216
300	E200
	70 160 160 160

(Source: Mitchell, 1990)

#### 2.7.1 The safety of food additives

To ensure the carbonated product reaches the consumer in a satisfactory state, it is necessary to create a balanced blend of component parts that will remain predictably sound during retail shelf-life for at least twelve months, under recommended storage conditions. To some, 'component parts' are synonymous with 'additives', a term which often provokes concern, but it must be emphasized that food additives receive official approval only if toxicological testing in the light of current knowledge yields no signs of harmful effects, nor any reason to suspect that harmful effects may arise.

The safety of food additives is tested according to guidelines issued by the joint committee of JECFA, WHO and FAO. Knowledge of health safety is gained primarily in animal experiments. In later stages of testing, humans may also be included in the studies to ascertain that their physiological reactions are similar to those found in animals.

Because of potentially longer term ingestion, the standards applied to food additives in feed trials are significantly higher than those applied to pharmaceuticals where, to treat successfully some conditions of illness, certain side effects caused by the treatments can be tolerated. The ADI is an estimate of the amount of a food additive that can be consumed safely over an entire lifetime. ADI levels are set by JECFA after considering the results of various feeding trials. As a general rule, the ADI value is set at one hundredth of the intake, which produces virtually no toxicological effects in long term animal feeding trials (Mitchell, 1990).

#### 2.8 Process Capability

Process capability is the long-term performance level of the process after it has been brought under statistical control. In other words, process capability is the range over which the natural variation of the process occurs as determined by the system of common causes.

Process capability, for a stable manufacturing process, is the capacity of the process to reach a certain level of quality. It can be shown graphically using histograms and process capability charts

## 2.8.1 Process Capability Index (Cp)

The Process Capability Index (Cp) is expressed as a ratio to the specified value. It is used to quantitatively evaluate the adequacy of the of the process capability – whether the variation in the process is within the limits of the specifications (SLSI, 2006).

Ср	Evaluation	Assessment
Cp > 1.33	Good	Process capability completely meets specifications.
1.33 >= Cp > 1.0	Acceptable	Process capability does not completely meet specifications; process control should be continued.
1.0 >= Cp	Inadequate	Process capability inadequate; improvements should be made.

 Table 2.5: Process capability assessment table

(Source: SLSI, 2006)

# CHAPTER 03

# Methodology

#### 3.1 Define the scope, food safety policy and related objectives

Scope of the food safety management system was defined by identifying the company profile as a entering to the food safety management system study. Food safety policy and related objectives were recorded as a preliminary step to implementation of food safety management system and to identify the commitment of the company for its safe food production.

#### 3.2 Identification of a food safety team

Food safety team was designated to consist of appropriate number of members with specific knowledge and responsibilities that each person involved in the company activities. The team was appointed team leader who had a good knowledge about the factory activities and experienced in implementation of food safety management system.

#### 3.3 Appoint a responsibilities and authorities

**Responsibilities and authorities related to each person was defined and communicated** within the organization to ensure the effective operation and maintenance of the food safety management system.

# 3.4 Establishment of effective arrangement for communication, emergency preparedness, management review and training programs

Effective arrangement for communication through out the food chain, emergency preparedness programs, management review meetings at planned intervals and training programs were scheduled according to the organization activities.

#### 3.5 Product description and intended use of carbonated beverages

Product features and attributes were understood and description of the product such as composition, structure, processing, packaging, storage and distribution conditions, and expected shelf life as well as type of raw materials used, ingredients used, and quantities of ingredients were identified and recorded. Product's intended usages were identified as considering its characteristics and observing catered target market.

### 3.6 Construction of process flow diagram

**Process flow for each product were identified and constructed the flow diagram by covering all the stages from raw materials to end products for evaluate the possible occurrence, increase or introduction of food safety hazards.** 

Factory layout was identified with potential delay stages. Simultaneously all possible cross-contamination risks that would happen were identified. Then the process flow diagrams were confirmed with the discussion of the food safety team.

### 3.7 Hazard assessment by identifying significant hazards

All food safety hazards such as biological, chemical and physical hazards that are reasonably expected to occur in relation to the type of product and process were identified and recorded. The identification was based on the preliminary information, experiences, external information and information from the food chain.

A hazard assessment was conducted to determine, for each food safety hazard identified, whether its elimination or reduction to acceptable levels was essential to the production of safe food, and whether its control is needed to enable the defined acceptable levels to be met. Decision tree was used to find out answers for above questions.

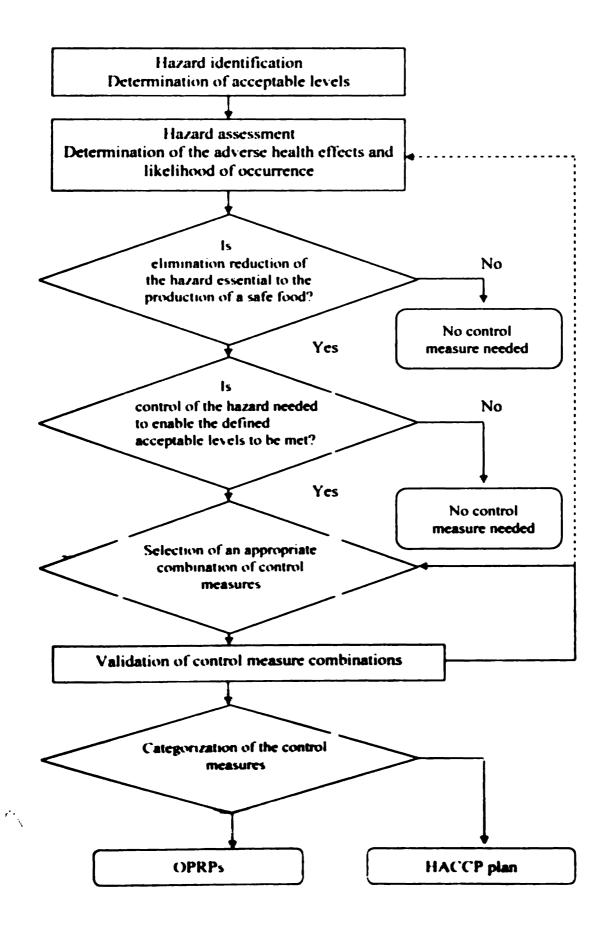


Fig. 3.1: Hazard assessment decision tree (ISO/TS 22004:2005(E))

Each food safety hazard was evaluated according to the possible severity of adverse health effects and the likelihood of their occurrence using risk assessment matrix.

gh	Low severity	Moderate	Serious	High severity
	high	severity high	severity high	high
	occurrence	occurrence	occurrence	occurrence
occurrence	Low severity medium occurrence	Moderate severity medium occurrence	Serious severity medium occurrence	High severity medium occurrence
occur	Low severity	Moderate	Serious	High severity
	rear	severity rear	severity rear	rear
	occurrence	occurrence	occurrence	occurrence
w	Low severity	Moderate	Serious	High severity
	low	severity low	severity low	low
	occurrence	occurrence	occurrence	occurrence

Low

Severity

High

Fig. 3.2: Risk assessment matrix

Criteria for severity

- No or little discomfort
- Diarrhea
- Doctor, hospitalization
- Life threatening (death)

Probability likelihood of occurrence

- Once per year
- Monthly
- Daily
- Constantly

The risk level was differentiated in 5 categories

- I Negligible no need to have control measures
- 2-3 Minor low level of control
- 4-8 Important -- implement require PRPs
- 9-15 Major more stringent OPRPs
- 16 Critical HACCP plan

### 3.8 Selection and assessment of control measures

**Based** on the hazard assessment appropriate combination of control measures was selected which is capable of preventing, eliminating or reducing the food safety hazard to defined acceptable levels.

### 3.9 Development of Prerequisite programs (PRPs)

Prerequisite programs (PRPs) were developed by considering and utilizing appropriate information such as statutory and regulatory requirements, customer requirements, recognized guidelines, Codex principles and codes of practices, and national or international standards.

### 3.10 Establishment of the Operational Prerequisite programs (OPRPs)

By considering the control measures and the monitoring frequency, determined OPRPs were established.

Food safety hazard(s) to be controlled by the program, control measure(s), monitoring procedures that demonstrate that the OPRPs are implemented, corrective actions to be taken if monitoring shows that the OPRPs are not in control, responsibilities and record(s) of monitoring were included to the Operational Prerequisite program.

### 3.11 Establishment of the HACCP plan

According to the hazard assessment, determined HACCP plan was established. Food safety hazard(s) to be controlled at the CCP, control measure(s), critical limit(s), monitoring procedure(s), corrective action(s) to be taken if critical limits are exceeded, responsibilities and record(s) of monitoring were included to the HACCP plan.

### 3.12 Validation of control measure combinations

CO<sub>2</sub> gas volume, brix and pH value in the cola, mandarin, soda, cream soda and lemonade process were measured by using gas volume tester, refractometer and pH meter. Data were collected within three months. Validation of control measure combinations were done by using process capability analysis to the collected data. Microbiological analysis for raw water was done in once a month.

### Methods for microbiological analysis of raw water Aerobic plate count test (Pour plate method)

Materials: water samples, 1.0 ml pipettes, petri dishes. Method:

1 ml of the test samples were transferred to the sterilized petri dishes using sterilized pipettes. 15 ml of the plate count medium at  $45 \pm 0.5$  °C was poured in to each petri dish. (Procedure was repeated with the prepared dilution series). The inoculum with the medium was mixed thoroughly and allowed to solidify placing the petri dishes on a clean horizontal surface. After complete solidification, the dishes were inverted and incubated at  $30 \pm 1$  °C for  $72 \pm 3$  hours. After the specified period of incubation, colonies were counted using the colony counting equipment (SLS 516: part 1: 1991).

### Presumptive coliform test

Materials: water samples, 1.0 ml pipettes, 5 ml of single strength MacConkey broth with inverted Durham tube.

### Method:

The sample bottles were inverted several times. The stopper was removed and flame the mouth of the bottle. Volumes of 1.0 ml of the original samples were inoculated in to 5 ml of single strength MacConkey broth with inverted Durham tube. The inoculated tubes were incubated at 36.0  $^{\circ}$ C ± 0.5  $^{\circ}$ C and examined each tube for the production of acid and gas after 24 to 48 ± 2 hours or earlier.

Acid and gas produced tubes constitute as presumptive coliform positive and the absence of formation of acid and gas constitute a negative test (SLS 614: part 2: 1983).

### Confirmation coliform and E.coli test

Materials: peptone water (pre heated to 44 °C), Kovac's reagent, test tubes, pipettes. Method:

All the presumptive positive coliform tubes were sub cultured in to tubes of peptone water (pre heated to 44 °C). Sub cultured tubes were incubated for 6 to 24 hours in a water bath maintain at 44 ± 0 1 °C. The incubated tubes were immersed in the water at least to the level of broth in the tube. Half of the contents of the peptone culture were transferred to a test tube in aseptic precautions. 0.2 ml = 0.3 ml of Kovac's reagent was added to it and mixed gently.

The appearance of red colour in the upper layer indicates a positive for confirmed coliform and E.coli. If gets positive for confirmed coliform and E.coli; Most Probable Number (MPN) of coliforms and E.coli were estimated by calculating the MPN of coliforms per 100 ml of sample and MPN of E.coli per 100 ml of sample (SLS 614: part 2: 1983, table in appendix C).

### 3.13 Verification and improvement of food safety management system

Internal audit plan and food safety plan were developed to complete the verification procedure of food safety management system.

### 3.14 Documentation of food safety management system

All the records and forms were documented as Pre requisite programs, Operational Pre requisite programs and HACCP plan.

### **CHAPTER 04**

### **Results and discussion**

### 4.1 Scope, food safety policy and related objectives

### 4.1.1Scope

The food safety management system was developed to prevent, eliminate or reduce food safety hazards reflected with carbonated beverage process, products and production sites in order to ensure that food is safe at the time of human consumption through out the food chain.

### **Company** profile

Shaa Cola Beverages (Pvt) Limited is one of the subsidiary company of Jaycey group and it was set up in year 2006.

Company engaged in the process of manufacturing range of carbonated beverages including soda, cream soda, cola, mandarin, tamarind, lemonade, etc. which are marketed under the brand name "Shaa".

The company is situated at Millawa, Moragahahena, in Horana town in Kalutara District and the head office of the company is at 11/7, Baddagana, Pitakotte in Colombo district.

All the raw materials required for production of Sogo Bottled Drinking Water such as PET bottles, Lids, Labels, Corrugated boxes are also manufactured by other subsidiary companies of Jaycey Group.

Water required for production of carbonated beverages, is extracted from more than 120 feet deep tube wells as a main raw material. Other raw materials are purchased form the approved list of suppliers and packaging materials are purchased from fully owned subsidiaries of JC group of companies.

They incorporate some of the fully automated modern machinery with high technology,  $\sim$  such as carbonation, filling and capping machines. There is no any single point in the filling process where humans are directly involved with open product.

Quality Assurance testing of the process is carried out at fully equipped in house laboratory where well qualified personnel are involved.

Marketing, Sales and Distribution are done by Shaa Cola Beverages (Pvt) Ltd and the company use their own studio (jubilant media network) and printing press to advertise & promote their products.

### 4.1.2 Food safety policy and related objectives

### Food safety policy

Shaa cola beverage (pvt) limited is committed to produce safe and superior quality beverages to surpass the customer expectation by continual improvement across our value chain by;

- Developing and implementing a food safety management system towards fulfilling the food safety objectives.
- Ensuring safe and healthy work environment and safe guarding property as per the company requirements.
- Continually training our employees on safe manufacturing practices and vigorously implementing best practices in all areas to protect the health and safety of our employees, customers and consumers.
- Assure the food safety and reliability to all our products.

### Food safety objectives

### Laboratory and Quality Assurance Section

- Maintain the microbiological parameters in finished product by maintaining maximum hygienic conditions in production area.
- Maintain the chemical parameters in finished product by calibrating instruments and correct ingredient weighing.
- HR Section
  - Improve the skills and knowledge of employees for safe food production by continuous training in each section.

### Production

- Increase safe food production by 12% quarterly
- Reduce food wastage incidences by 2% quarterly

Increase customer satisfaction index (product development, product modification)
 by 1% quarterly.

### 4.2 Identification of a food safety team

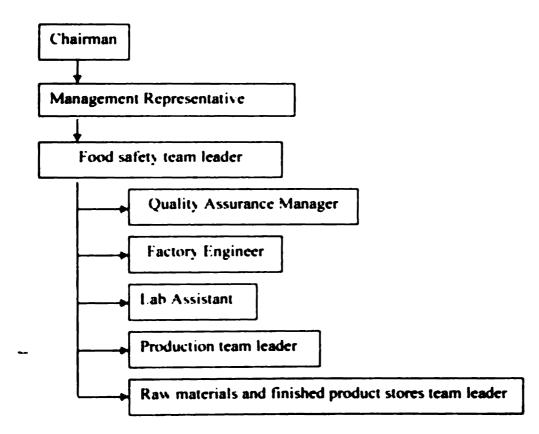


Fig. 4.1: Food safety team

### 4.3 Appoint a responsibilities and authorities

### Chairman

Overall responsibility of the management in the company

### Management Representative

• The food safety management related responsibilities are vested with MR

### Food safety team leader - Operational Manager

- Manage a food safety team and organize its work.
- Update the preliminary information and documents specifying the PRPs, OPRPs and the HACCP plan.
- Ensure a relevant training and education of the food safety team members.
- Ensure that food safety management system is established, implemented, maintained and updated.
- Report to the top management on the effectiveness and suitability of the food safety management system.
- Responsible and authority related to monitoring and evaluation of monitoring results.
- Verification results should be recorded and communicated to the food safety team.
- Responsible for control of non conformities.

### Quality Assurance Manager

- Identify the quality assurance points of the production and other supportive activities and implement documented procedures.
- Identify the resources required to introduce the above quality assurance measures and communicate with food safety team leader.
- Maintaining of environmental hygiene and personal hygiene and training of workers.
- Carry out R&D experiments to upgrade the products and introduce new products.
- The quality related responsibilities given in the interface matrix are also vested with QAM.

### **Factory Engineer**

- Responsible for optimization of machines to achieve the production target.
- Servicing of the machines according to the schedule plan
- Training of machine operators.
- Responsible for maintaining of tool box and spare parts of the machines and other equipments.

- Minimize machines break downs.
- Changing of the machine mould (For changing of production category according to the schedule production plan).

### Lab Assistant

- Responsible for minimum contamination through out the food chain.
- Carry out chemical and microbiology tests according to the schedule plan.
- Take immediate action if having any food safety problem.
- Responsible for changing UV bulb and filters.
- Report to the food safety team leader on the effectiveness of microbiological test results.

### Production team leader - Supervisor

- Implementation of monthly production plan given by the food safety team leader to meet the monthly set target and company food safety objectives.
- Responsible for the fulfilling of daily production target by producing safe products.
- Arrange the production line so that minimize unnecessary delays of production flow, and educate the workers of the same (Time management and work study)
- Identify required resource for the production flow and communicate it with food safety team leader.
- Responsible of maintaining personal and environmental hygiene.
- Maintaining good work practice and discipline in the production flow.

### Raw materials and finished product stores team leader - Store Keeper

- Implementation of monthly raw material receiving plan and finished product dispatching plan given by the food safety team leader to meet the monthly set target and company food safety objectives.
- Arrange the stores as a "FIFO" method so that minimize unnecessary delays of raw materials and finished product supplements.
- Identify required raw materials or any problems related to stores and communicate it with food safety team leader

### 4.4 Establishment of effective arrangement for communication, emergency preparedness, management review and training programs

### 4.4.1 Communication

### External communication

Top management of Shaa Cola beverages (Pvt) Ltd ensures that appropriate communication channels are established to communicate with suppliers, contractors, customers/consumers, statutory and regulatory authorities and other organizations. The following communication channels are used for the purpose.

- Advertisements in Max television and Max radio
- News paper, poster, sticker advertisements
- Label in the product

### **Internal** communication

Top management of Shaa Cola beverages (Pvt) Ltd ensures that appropriate communication channels are established within the company. The following communication channels are used for the purpose.

- Management led communication in work areas
- Team briefings and other meetings
- Notice boards
  - Audio visual and electronic media

### 4.4.2 Emergency preparedness

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Top management of the company should be prepare that response to potential emergency situations and accidents that can impact on food safety include incidents such as fire, flooding, bio-terrorism and sahotage, emergency failure, vehicle accidents, contamination of the environment, various type of whether related events or the impact of the pandemic by; emergency preparedness planning relevant to food safety policy, appointing incident planning coordinator, forming incident planning committee with representatives from all the major parts of the organization, train the staff and carry out exercises of the plan.

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Table 4.1: Emergency	preparedness plan
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Emergency situations	Action taken	Responsibility	Incident planning coordinator
1. Fire	Deploy available fire-fightin facilities. Inform to factory engineer. Isolate products likely to b contaminated by fire extinguishe chemicals.	-	Factory engineer
2. Power cut	Inform to factory engineer. Start generators to meet key power requirements.	Every person in the company	Factory engineer
3. Water pressure drop or stoppage	Inform to factory engineer. Limit water use to vital plant functions only.	Every person in the company	Factory engineer
4. Emergency failure	Inform to operational manager. Take necessary action.	Every person in the company	Operational manager
5. Workers accidents	Inform to medical officer. Immediately supply treatments.	Every person in the company	Medical officer
6. Vehicle accidents	Inform to medical officer. Damage person immediately transport to the hospital	Any person	Medical officer

### 4.4.3 Management review

Management representative (MR) shall organize the management reviews at least once in three months. If need arises due to Non-implementation of the recommended actions agreed during the previous Management Review Meeting, accumulation of a number of customer complaints, accumulation of a number of non-conformities the MR shall convene an extra ordinary Management Review Meeting.

MR shall inform the meeting to the committee members using the minutes of the previous meeting.

The committee shall discuss the review inputs such as follow-up actions from previous management reviews, analysis of results of verification activities, changing circumstances that can affect food safety, emergency situations, customer feedback, external audits or inspections. Review output should be included decisions and actions related to assurance of food safety, improvement of the effectiveness of the food safety management system, resource needs and revisions of the organization's food safety policy and related objectives.

These meetings were minute by indicating recommendations clearly with responsibilities and time frames for implementation and circulate the minute among the participants within one week after the meeting. All the relevant officers shall ensure that recommended actions are implemented by the identified target dates. All records pertaining to the management review meetings shall be maintained by the MR.

### 4.4.4 Training programs

Training programs provide a procedure to maintain & improve the knowledge of the employees in the company to consistently meet and exceed customer requirements. The overall responsibility of training the staff is vested with management representative (MR). MR identifies the training needs at least once a year using information from employee performance evaluation report. The training was carried out as; induction and orientation training for new recruits, on the job training to improve knowledge and operational skills for safe food production by ensuring personal responsibilities.

The detail of the training received is summarized and evaluation of training during the year is carried out by operational manager and MR.

### 4.5 Product description and intended use of carbonated beverages

**Product is a carbonated beverage that will be consumed mainly for the thirst quenching property by the general population including children and elders.** 

### Product name: Shaa cola

- Ingredients:Carbonated waterSugarCaramel colourCola part B (Artificial flavour)Sodium benzoate E 211 (Preservative)
  - Ortho phosphoric acid E 338 (Acidualant)

### Packaging materials:

PET bottles: Colourless Bottle cap (with neck cap): Red in colour

### **Chemical characteristics:**

Gas volume range: 4.0 - 5.0 Brix range: 9.5 - 10.5 pH range: 2.0 - 3.0 Benzoic acid requirement: 160 mg/kg Caffeine content: < 150 mg/kg

### Product name: Shaa cream soda

### Ingredients: Carbonated water

Sugar

Sunset yellow - E 110 (Colour)

Cream soda flavour (Artificial flavour)

Sodium benzoate - E 211 (Preservative)

Citric acid - E 330 (Acidualant)

Xanthen gum - E 415 (Stabilizer)

### **Packaging materials:**

**PET bottles: Colourless** 

Bottle cap (with neck cap): violet in colour

### **Chemical characteristics:**

Gas volume range: 3.0 - 4.0 Brix range: 12.5 - 13.5 pH range: 3.0 - 4.0 Benzoic acid requirement: 160 mg/kg

### Product name: Shaa soda

Ingredients: Carbonated water Sodium bicarbonate

### **Packaging** materials:

PET bottles: Colourless

Bottle cap (with neck cap): Ash in colour

### **Chemical** characteristics:

Gas volume range: 4.0 - 4.5 Brix range: pH range: 4.0 - 4.5 Sodium bicarbonate: < 560 mg/kg

### Product name: Shaa mandarin

Ingredients: Carbonated water Sugar Sunset yellow - E 110 (Colour) Dalandan flavour - nature identical (Artificial flavour) Sodium benzoate - E 211 (Preservative) Citric acid - E 330 (Acidualant) Xanthen gum - E 415

### Packaging materials:

PET bottles Colourless

Bottle cap (with neck cap). Orange in colour

### **Chemical characteristics:**

Gas volume range: 2.5 3.5 Brix range: 12.0 13.0 pH range: 2.5 3.5 Benzoic acid requirement: 160 mg/kg

### Product name: Shaa tamarind

Ingredients: Carbonated water

Sugar

Caramel colour

Tamarind flavour (Artificial flavour)

Sodium benzoate - E 211 (Preservative)

Sodium chloride

Citric acid - E 330 (Acidualant)

Sodium dehydrogen citrate - E 331 (Acidity regulator)

### **Packaging materials:**

PET bottles: Colourless

Bottle cap (with neck cap): Brown in colour

### **Chemical characteristics:**

Gas volume range: 3.0 - 3.5 Brix range: 12.5 - 13.0 pH range: 2.5 - 3.5 Benzoic acid requirement: 160 mg/kg

### Product name: Shaa lemonade

Ingredients: Carbonated water Sugar Natural lemon flavour Sodium benzoate - E 211 (Preservative) Citric acid = E 330 (Acidualant) Sodium dehydrogen citrate = E 331 (Acidity regulator)

### Packaging materials:

PET bottles: Light green Bottle cap (with neck cap): Dark green in colour

### **Chemical characteristics:**

Gas volume range: 3.0 - 4.0 Brix range: 9.5 - 10.5 pH range: 2.5 - 3.5 Benzoic acid requirement: 160 mg/kg

• Below mention characteristics are same for all above beverage products.

### Microbiological specifications:

Standard plate counts, per ml, max.	<100
Coliforms, per ml	absent
Yeast and moulds count per ml	absent

### Heavy metal limits:

	Arsenic, mg/kg, max.	0.2
-	Copper, mg/kg, max.	2
-	Lead, mg/kg, max.	0.2

Packaging size/ Volume: 350 ml, 500 ml, 1000ml

Shelf life: 6 months from manufacture date under ambient temperature or refrigerated conditions and away from sun light.

Label information: Product name, Brand name, Batch code, Date of manufacturing, Date of expiry, Retail price, Barcode, Storage condition, Contact address printed on the bottle.

Intended consumer: Consumers at all ages

Intended use: Ready to drink

### 4.6 Construction of process flow diagram

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### Process flow diagram for Shaa soda

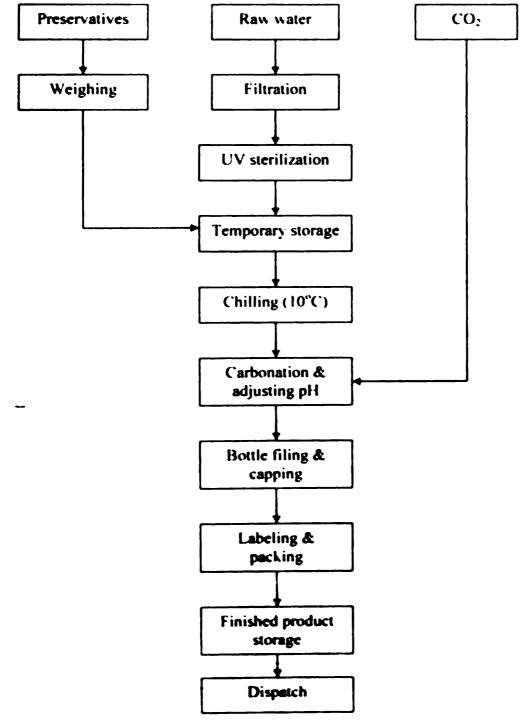
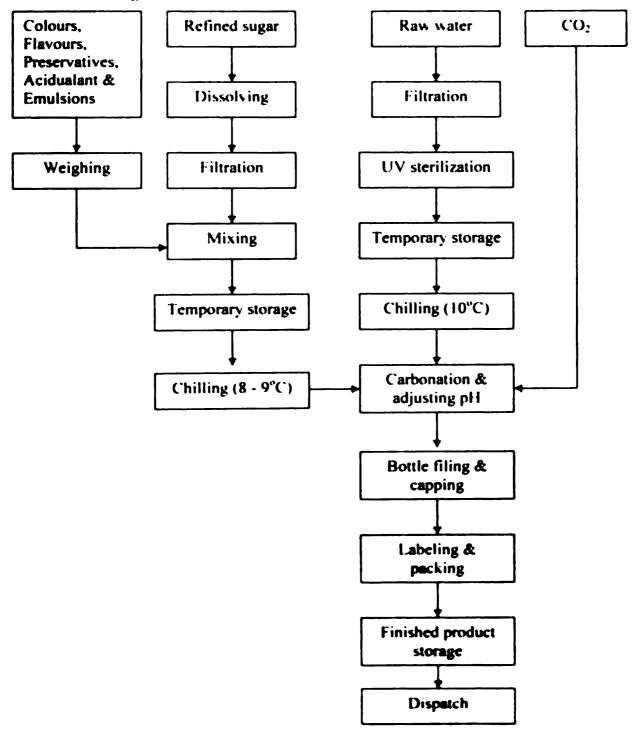


Fig. 4.2: Process flow diagram for Shaa soda



Process flow diagram for Shaa cola, cream soda, mandarin, tamarind and lemonade

Fig. 4.3: Process flow diagram for Shaa cola, cream soda, mandarin, tamarind and lemonade

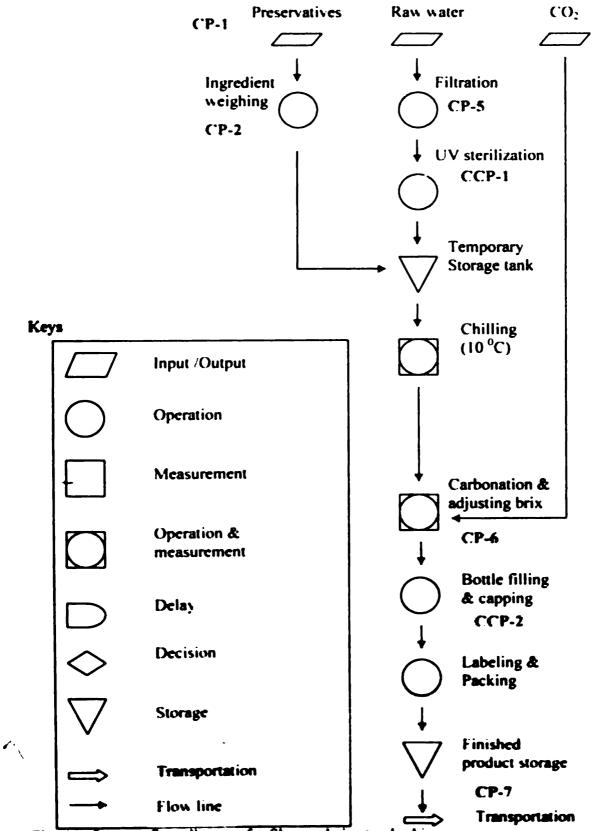


Fig. 4.4: Process flow diagram for Shaa soda in standard icons

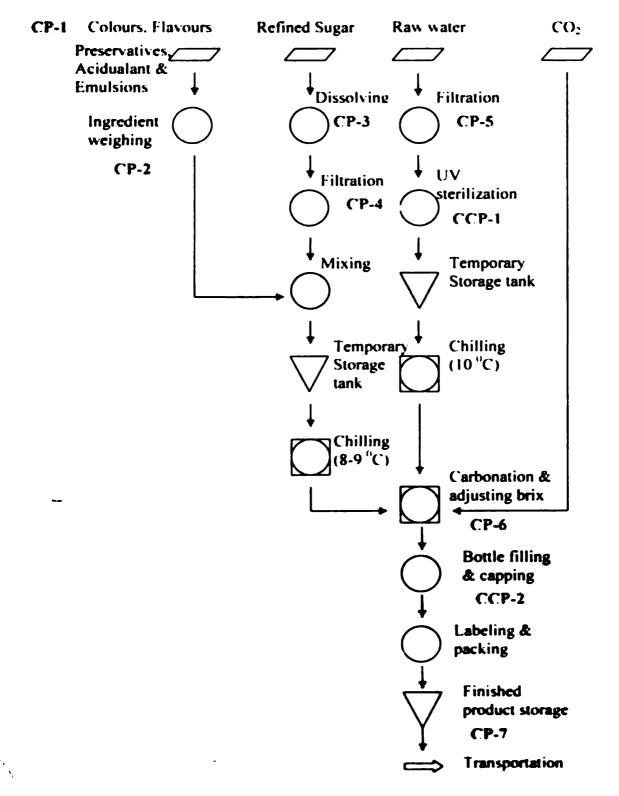
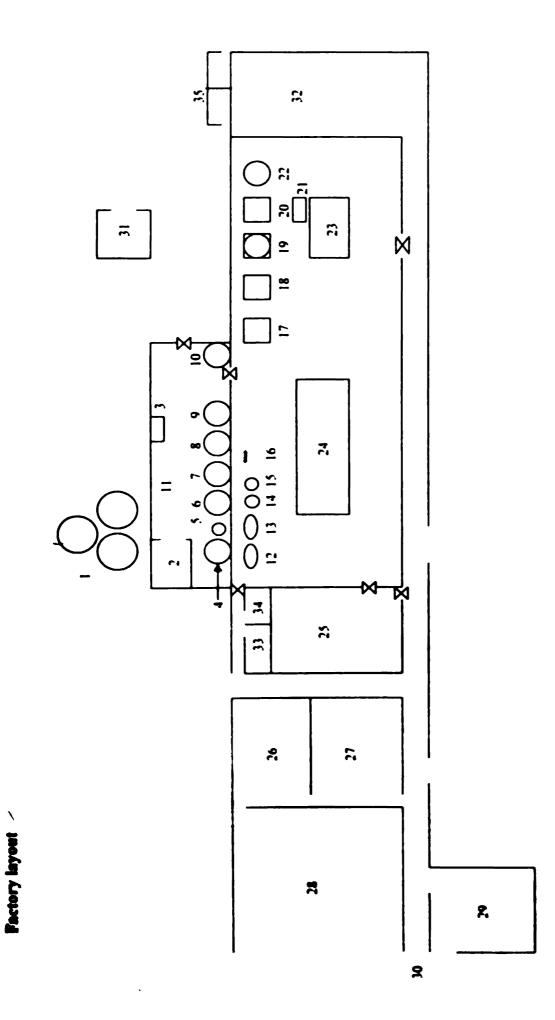


Fig. 4.5: Process flow diagram for Shaa cola, cream soda, mandarin, tamarind and lemonade in standard icons



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# Lecations indicated by numbers:

- 1. Raw water storage tank
- 2. Raw material storage area

Sugar, acidualant and preservatives (bags)

Colours. Rayours, and emulsion (cans)

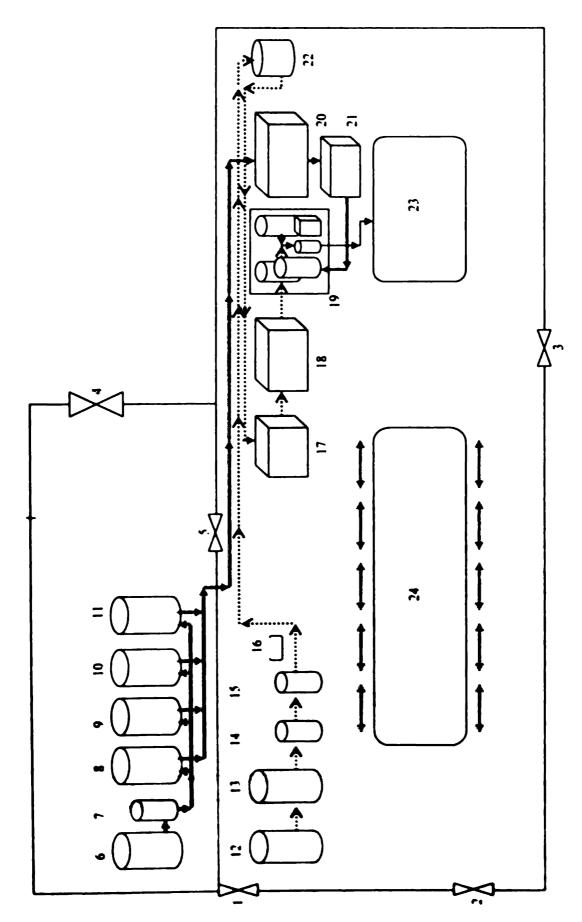
- Electric halance
- Sugar mixing tank
- Sugar filtration (0.5 micron filters)
- 6 Syrup storage tank-1
- 7 Syrup storage tank-2
- 8 Syrup storage tank-3
- Syrup sionage tank -4
- 10 Water storage tank
- 11 Syrup river
- 12 Water filtration (sand filter)
- 13 Water filtration (Active carbon filter)
- 14 Water filtration (0 5 micron filters)
- 15 Water filtration (0.2 micron filters)
- 16. UV stenlizer
- 17 Water chiller (10°C)
- 18 (Continues water chilling tank (10°C)

### 19. Carbonation plant

- 20. Syrup chiller (8°C 9°C)
- 21. Continues syrup chilling tank
- 22. Treated water storage tank
- 23. Filling and Capping machine
- 24. Labeling & packing area
- 25. Raw material storage area
- (PET bottles and lids)
- (Labels and corrugated boxes)
- 26. Laboratory
- 27. Office
- 28. Sogo drinking water bottling section
- 29. Finished product storage area
- 30. Factory entrance
- 31. Canteen
- 32. Maintenance equipment storage area
- 33. Changing room male
- 34. Changing room female
- 35. Toilets



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# Lacations indicated by numbers:

- 1. Workers entrance
- 2. Raw material entrance (PET bottles, lids, cartons)
- 3. Finished product exit
- 4. Raw material entrance (sugar and other ingredients)
- 5. Symp room entrance
- 6 Sugar mixing tank
- 7 Sugar filtration (0.5 micron filters)
- 8 Syrup worage tank-1
- 9. Syrup storage tank-2
- 10 Syrup storage task-3
- 11 Syrup storage tank-4
- 12 Water filtration (sand filter)
- 1) Water filtration (Active carbon filter)
- 14 Water filtration (0 5 micron filters)

15. Water filtration (0.2 micron filters)

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- 16. UV sterilizer
- 17. Water chiller (10°C)
- 18. Continues water chilling tank (10°C)
- 19. Carbonation plant
- 20. Syrup chiller (8°C 9°C)
- 21. Continues syrup chilling tank
- 22. Treated water storage tank
- 23. Filling and Capping machine
- 24. Labeling & packing area
- Concentrated syrup flow (brix 60° -65°)
  ....... Water flow
  Diluted syrup flow (brix 10° -15°)
  - --- Workers flow

	Significant hazards	Is climination/r cduction of	is control of the hazard	
Process step	Biological: B C'hemical: C	the hazard essential to the	needed to enable the defined	Control measure
	Physical: P	production of a safe food?	acceptable levels to be met?	
I. Raw matchal				
• Sugar	B: Pathogenic bacteria, Yeasts, Moulds	Yes	Yes	OPRP/HACCP
	C: Sulphate ash. Cleaning chemicals. Antibiotics. Pesticides	Ycs	Yes	COPRP-HACCP
	P: Water insoluble matters. Dead bodies (pest/insects).	Yes	Ycs	OPRP/HACCP
	Dust particles. Metal particles			
(0)	B: Not applicable	No	·	No need
	C: presence of toxic gasses	Yes	Yes	<b>OPRP</b> <sup>HACCP</sup>
	P: Foreign matters	Yes	Yes	OPRP/HACCP
• Flavours.	B: Air born contaminants	Yes	Ycs	OPRP/HACCP
E-mulsions.	C: Toxic substances. Oxidants, Antibiotics	Yes	Yes	<b>OPRP-HACCP</b>

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preservatives	P: Foreign matters	Yes	Yes	OPRP/HACCP
• PET bottles &	~~~			
lids	B: Contamination with vegetative pathogens	Yes	Yes	OPRPHACCP
	C: Toxic chemicals	Yes	Yes	OPRP HACCP
	P. Foreign matters (dust/plastic particles)	Ycs	Yes	OPRP HACCP
2 Raw water filtration	B. Remaining Algae spores	Yes	Yes	· OPRP HACCP
	C: Remaining organic matters, Cleaning chemicals	Yes	Yes	OPRP-HACCP
	P. Remaining suspended particles. Impurities	Yes	Ycs	OPRP HACCP
3 Water UV	B: Proliferation or survival of vegetative pathogens	Yes	Yes	OPRP/HACCP
stenization	C. Not applicable	No	•	No need
	P: Not applicable	No	•	No need
4 Water chilling	B. Survival of pathogenic bacteria	Yes	Ycs	OPRPHACCP
	C: Cleaning chemical residues	Yes	Yes	OPRP-HACCP
	P. Contamination by gasket materials	Yes	Yes	<b>OPRP-HACCP</b>
5 Sugar dissolving	B: Survival of pathogenic bacteria or yeasts	Yes	Yes	OPRP HACCP
	C: Formation of acid floc, Cleaning chemical residues	Yes	Ycs	OPRP-HACCP
	P: Drop of foreign matters in to the tank, contamination of	Yes	Yes	<b>OPRP HACCP</b>
	gasket materials			

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6. Sugar filtration	B: Remaining yeasts	Yes	Yes	OPRP/HACCP
	C: Remaining acid floc	Yes	Ycs	OPRP/HACCP
	P: Remaining un wanted particles & other turbidity components	Yes	Yes	OPRPTHACCP
7. Ingredient weighing	B: Contamination by person	Yes	Yes	OPRPHACCP
	C: Measuring of excess amount of food additives	Yes	Yes	ОРКР НАССР
	P. Drop of foreign matters	Yes	Yes	ОРКР НАССР
8 Mixing ingredients	B. Contamination by person	Ycs	Ycs	OPRP HACCP
with sugar solution	( Cleaning chemical residues	Yes	Ycs	ОРКР НАССР
	P. Drop of foreign matters	Yes	Ycs	OPRP/HACCP
9. Syrup chilling	B: Survival of pathogenic bacteria	Yes	Ycs	OPRP HACCP
	(' ('leaning chemical residues	Yes	Ycs	<b>OPRP-HACCP</b>
	P Contamination by gasket materials	Yes	Ycs	OPRP-HACCP
10 Carbonation &	B. Not applicable	No No	•	No need
adjusting brix	C: Excess carbonation, excess amount of syrup	Yes	Ycs	OPRP/HACCP
	P. Contamination by gasket materials	Yes	Ycs	OPRP/HACCP
II. filling & capping	B: Presence of pathogen organisms in filling pipes/bottles/lids	Yes	Yes	OPRP/HACCP
	Contamination by pathogen organisms due to improper bottle			
	scaling			-
	C: Cleaning chemical residues	Yes	Yes	OPRP/HACCP

	P: Contamination by metal particles in the filling machine Contamination by foreign matters in bottles/lids	Yes	Ycs	OPRP/HACCP
12.Storage and dispatch	12.Storage and dispatch B: Post contamination of pathogen organisms	Ycs	Yes	OPRP-HACCP
of hottles	C: Not applicable	No	•	No need
	P. Post contamination of foreign matters	Yes	Yes	d.).)VH dldO

### 4.8 Food safety hazard evaluation by categorizing control measures

Table 4.3: Food safety hazard evaluation by ca	ategorizing control measures
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Process step	Type of hazard	Risk level	Categorization of control measure
1. Raw material receiving			
<ul> <li>Sugar</li> </ul>	Biological	Major	OPRP/RRI & PRP/SA
	Chemical	Major	OPRP/RRI & PRP/SA
	Physical	Major	OPRP/RRI & PRP/SA
• CO <sub>2</sub>	Biological	-	-
	Chemical	Major	OPRP/RRI & PRP/SA
	Physical	Major	OPRP/RRI & PRP/SA
• Flavours,	Biological	Major	OPRP/RRI & PRP/SA
Emulsions, Acids,	Chemical	Major	OPRP/RRI & PRP/SA
colours &	Physical	Major	OPRP/RRI & PRP/SA
preservatives			
• PET bottles & lids	Biological	Major	OPRP/RRI & PRP/SA
	Chemical	Major	OPRP/RRI & PRP/SA
	Physical	Major	OPRP/RRI & PRP/SA
2. Raw water filtration	Biological	Major	OPRP/FPM
~	Chemical	Major	OPRP/FPM
	Physical	Major	OPRP/FPM
3. Water UV sterilization	Biological	Critical	HACCP plan
	Chemical	•	-
	Physical	•	-
4. Water chilling	Biological	Major	OPRP/CPM
C C	Chemical !	Important	PRPCS
	Physical	Important	PRP/CS
5. Sugar dissolving	Biological	Major	OPRP/SRM
	Chemical	Major	OPRP/SRM
	Physical	Important	PRP/CS
6. Sugar filtration	Biological	Major	OPRP/FPM
	Chemical	Major	OPRP/FPM
	Physical	Major	OPRP FPM

7. Ingredient weighing	Biological	Important	PRP PH
	Chemical	Major	OPRP/SRM & PRP C
	Physical	Important	PRP/PH
8. Mixing ingredients with	Biological	Important	PRP/PH
sugar solution	Chemical	Important	PRP/CS
	Physical	Important	PRP/PH
9. Syrup chilling	Biological	Major	OPRP/CPM
	Chemical	Important	PRP/CS
	Physical	Important	PRP/CS
10. Carbonation &	Biological	-	-
adjusting brix	Chemical	Major	OPRP/CPM
	Physical	Important	PRP/CS
11. filling & capping	Biological	Critical	HACCP plan
	Chemical	Important	PRP/CS
	Physical	Major	OPRP/SRM
12.storage and dispatch of	Biological	Major	OPRP/FPI
bottles	Chemical	-	-
	Physical	Major	OPRP/FPI

### 4.9 Development of Prerequisite programs (PRPs)

By studying the entire factory, production flow, internal and external environment, prerequisite programs (PRPs) were developed.

Table 4.4: Prerequisite	programs (PRPs)
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Prerequisite program		Reference
Calibration (C)	PRP 1	Appendix A
Cleaning and Sanitizing (CS)	PRP 2	Appendix B
Personal Hygiene (PH)	PRP 3	Appendix C
Pest Control (PC)	PRP 4	Appendix D
Supplier Assessment (SA)	PRP 5	Appendix E
Waste & Sewage Disposal (WSD)	PRP 6	Appendix F
Water, Air and Energy supply (WAE)	PRP 7	Appendix G

### 4.10 Establishment of the Operational Prerequisite programs (OPRPs)

By considering identified control points (raw material receiving, ingredient weighing, filtration, chilling, carbonation, finished product inspection), Operational Prerequisite programs (OPRPs) were established.

Process step		OPRP program	Reference
Raw material Receiving	CP-1	Raw material Receiving and Inspection (RRI) OPRP 1	Appendix H
Ingredient weighing	CP-2	Syrup Room Management (SRM) OPRP 2	Appendix 1
Sugar dissolving	CP-3	Syrup Room Management (SRM) OPRP 2	Appendix I
Sugar filtration	CP-4	Filtration Process Management (FPM) OPRP 3	Appendix J
Water filtration	CP-5	Filtration Process Management (FPM) OPRP 3	Appendix J
Carbonation	СР-6	Carbonation Process Management (CPM) OPRP 4	Appendix K
Storage and dispatch of	bottles P-7	Finished Product Inspection (FP1) OPRP 5	Appendix L

 Table 4.5: Operational Prerequisite programs (OPRPs)

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<b>•</b> \	it of the HACCP plan
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Table 4.6: HACCP plan

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Process step	Identified Hazard	Control limit	Monitoring	Corrective action	HACCP Records	Verification Procedure	Responsibility
Raw water sterilization CCP-1	Biological	Maintain the. • Acrobic plate count - 1.0•10 <sup>4</sup> per ml • Fotal coliform count (mpN) • 10per 100ml • E-coli count none in 100ml	<u>What</u> • Acrobic plate count • Total coliform coliform count thow Microchem laboratory Who Lab assistant Frequency Once a week	c Inform to the maintenance department and maintain wave length at 254 nm or replace the UV bulb after finishing recommende d working hours (9000hrs).	<ul> <li>Working hours of I V bulb - log sheet</li> <li>Changing of UV bulb - log sheet</li> <li>Maintenanc</li> <li>records</li> </ul>	olab assistant collects the records and reported to the factory engineer	Factory engineer
			4				

3.	
Responsibility	Factory cngineer
Verification Procedure	o machine operator inform to the factory cngineer
HACCP Records	o machine maintenanc e records
<b>Corrective</b> action	<ul> <li>Inform to the machine operator and maintain the capping strength</li> </ul>
	stion thour
Monitoring	What Capping strength How Measuring capping strength Who Production team leader Frequency Once a hour
Control limit Monito	Maintain the What capping strength Cappin at least 10 bar strength How Cappin strengt Produc team lo
<u></u>	Maintain the capping strength at least 10 bar

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### 4.12 Validation of control measure combinations

Prior to implementation of control measures included in OPRPs and HACCP plan, validation was done to determine that control measures were scientifically and technically sound. Information needed to validate was collected from in-house observations and measurements, scientific studies, expert advisers and evaluations.

For OPRPs validation; brix, CO<sub>2</sub> gas volume, and pH value of each Shaa products were measured and capability of control measures were evaluated by using process capability analysis. HACCP plan validation was done by information gathered from third part laboratory test reports, in-house observations and measurements and scientific studies.

### Validation of control measure combinations using process capability analysis



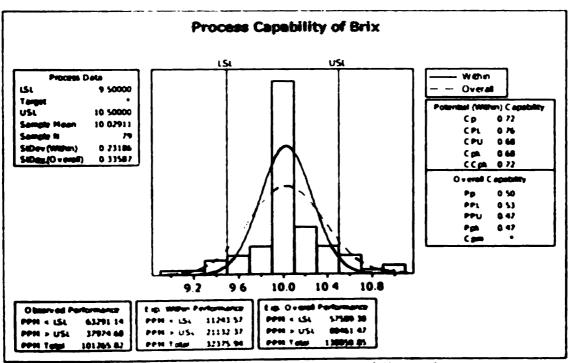


Fig. 4.8: Process capability analysis for brix value of Shaa cola

### Interpretation:

**Cp** = 0.72

1.0 >= Cp

Process capability is inadequate; improvements should be made

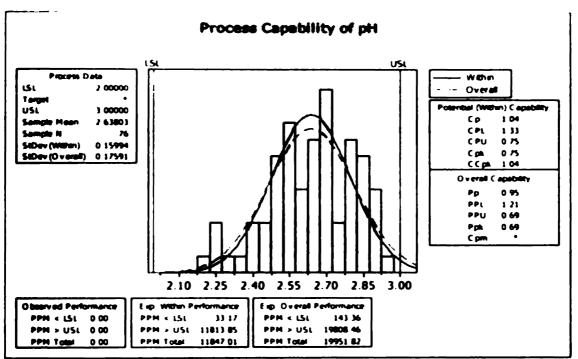


Fig. 4.9: Process capability analysis for pH value of Shaa cola

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Cp = 1.04

1.33 >= Cp > 1.0

Process is acceptable but process capability does not completely meet

specifications; process control should be continued.

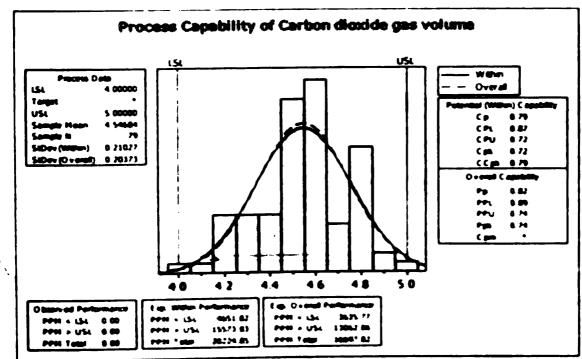


Fig. 4.10: Process capability analysis for CO; gas volume of Shaa cola

Cp 0.79

1.0 ° Cp

Process capability is inadequate; improvements should be made.

According to the process capability analysis for pH value, process is acceptable. Cp values for brix and  $CO_2$  gas volume also near to 1.0. Therefore process becomes acceptable. But process capability does not completely meet specifications; process control should be continued.

### Shaa mandarin

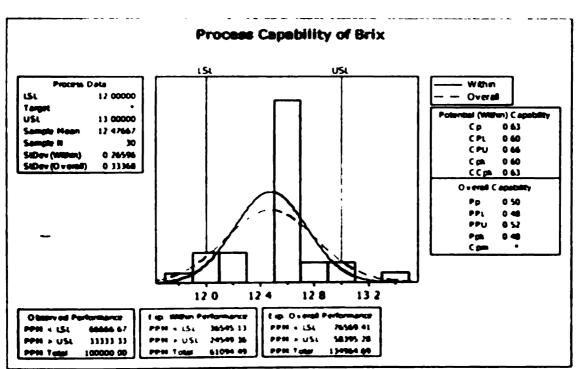


Fig. 4.11: Process capability analysis for brix value of Shaa mandarin

Interpretation:

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**Cp** = 0.63

1.0 > · Cp

Process capability is inadequate; improvements should be made.

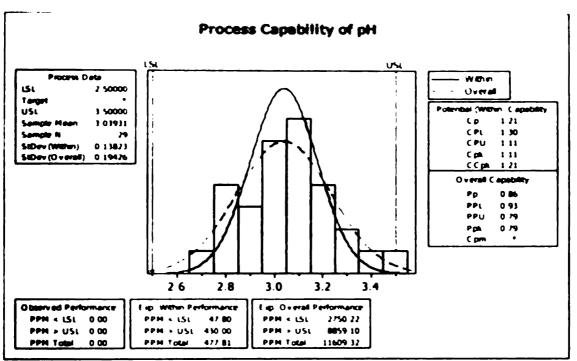


Fig. 4.12: Process capability analysis for pH value of Shaa mandarin

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Cp = 1.21

1.33 > Cp > 1.0

Process is acceptable but process capability does not completely meet

specifications; process control should be continued.

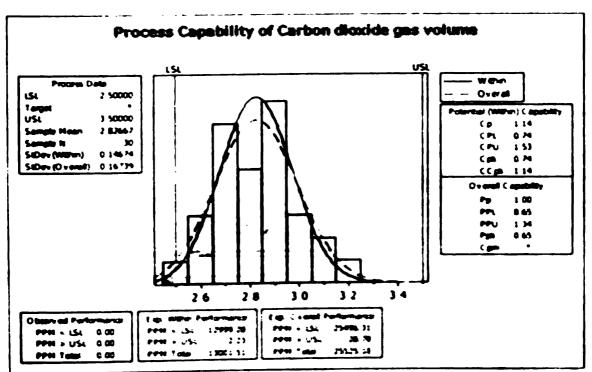


Fig. 4.13: Process capability analysis for CO; gas volume of Shaa mandarin

Cp 1.14 1.33 >= Cp + 1.0

Process is acceptable but process capability does not completely meet

specifications; process control should be continued.

According to the process capability analysis for pH value and  $CO_2$  gas volume, process is acceptable. Cp values for brix also near to 1.0. Therefore process becomes acceptable. But process capability does not completely meet specifications; process control should be continued.

### Shaa soda

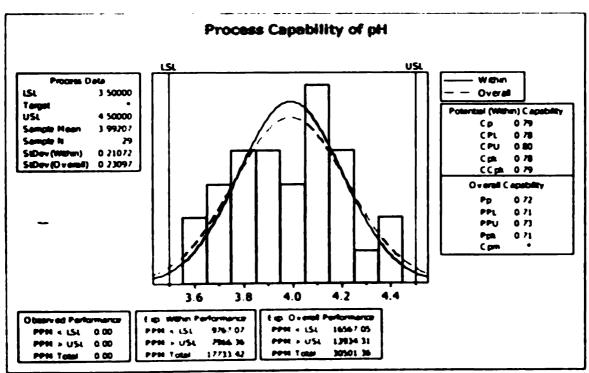


Fig. 4.14: Process capability analysis for pH value of Shaa soda

Interpretation:

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**Cp = 0.79** 

1.0 > = Cp

Process capability is inadequate, improvements should be made

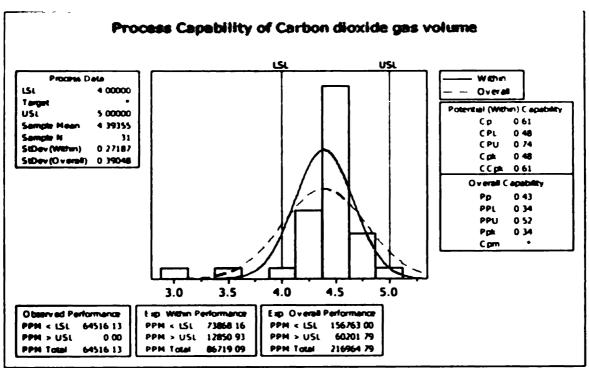


Fig. 4.15: Process capability analysis for CO2 gas volume of Shaa soda

**Cp** = 0.61

1.0 >= Cp

Process capability is inadequate; improvements should be made.

Shaa soda process capability is inadequate for both pH value and  $CO_2$  gas volume. Therefore improvements should be made for soda production process.

### Shaa cream soda

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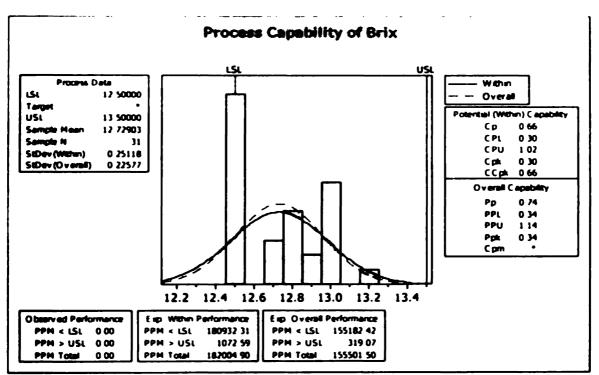


Fig. 4.16: Process capability analysis for brix of Shaa cream soda

Interpretation:

Cp = 0.66

Process capability is inadequate; improvements should be made.

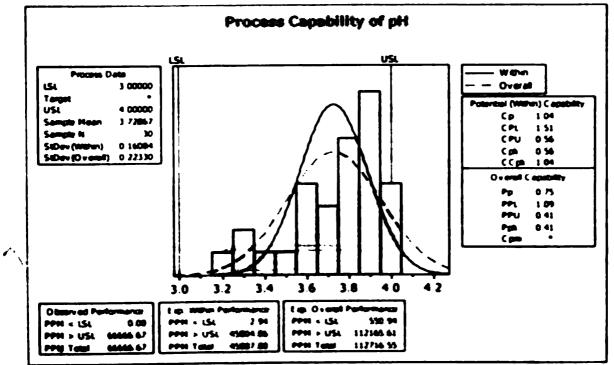


Fig. 4.17: Process capability analysis for pH value of Shaa cream soda

Cp ~ 1.04

1.33 >- Cp > 1.0

Process is acceptable but process capability does not completely meet specifications; process control should be continued.

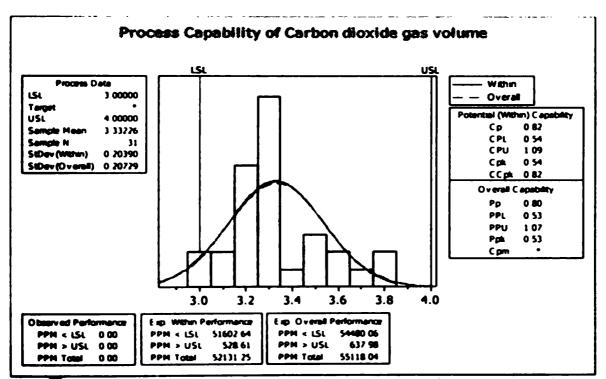


Fig. 4.18: Process capability analysis for CO2 gas volume of Shaa cream soda

Interpretation:

|-

**Cp** = 0.82

1.0 > - Cp

Process capability is inadequate; improvements should be made.

According to the process capability analysis for pH value, process is acceptable. Cp values for brix and CO<sub>2</sub> gas volume also near to 1.0. Therefore process becomes acceptable. But process capability does not completely meet specifications; process control should be continued.

### Shaa lemonade

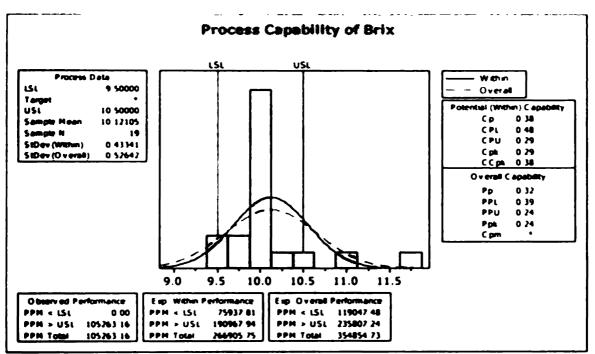


Fig. 4.19: Process capability analysis for brix of Shaa lemonade

#### Interpretation:

Process capability is inadequate; improvements should be made.

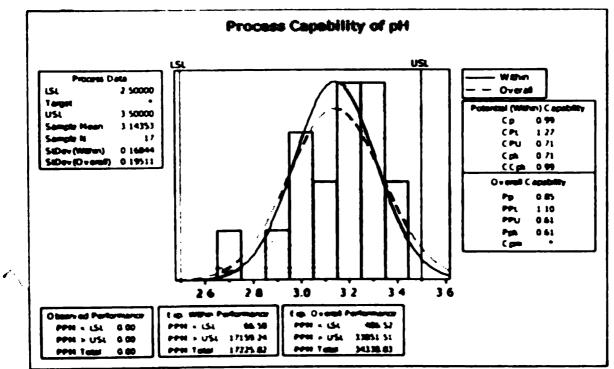


Fig. 4.20: Process capability analysis for pH value of Shaa lemonade

- Cp 0.99
- 1.0 Cp

Process capability is inadequate; improvements should be made.

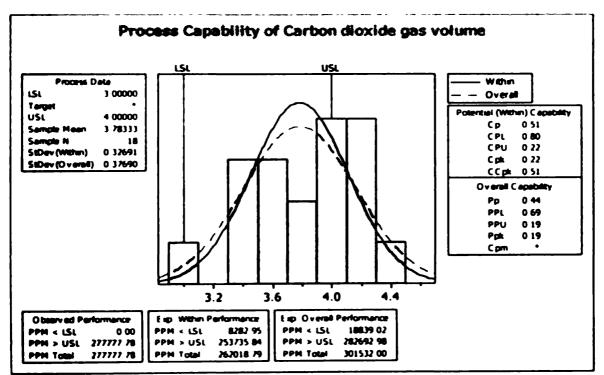


Fig. 4.21: Process capability analysis for CO2 gas volume of Shaa lemonade

### Interpretation:

1.0 >= Cp

Process capability is inadequate; improvements should be made.

Shaa lemonade process capability is inadequate for brix, pH value and CO<sub>2</sub> gas volume. Therefore improvements should be made for lemonade production process.

### Microbiological test results for raw water

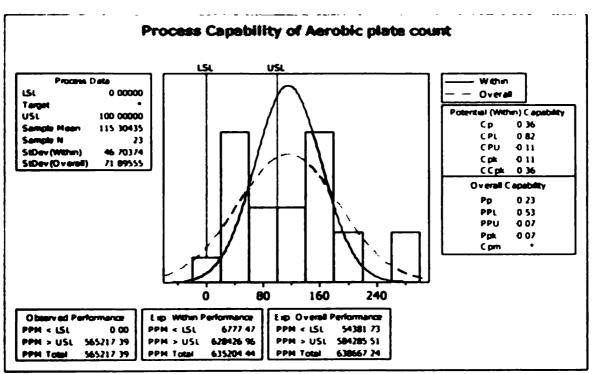


Fig. 4.22: Process capability analysis for aerobic plate count test results of raw water

Interpretation:

Process capability is inadequate; improvements should be made

Total coliform count and E.coli count in raw water was not detected (Appendix M). But process capability of aerobic plate count test is inadequate. Therefore improvements should be done by improving environment hygiene as well as personal hygiene.

### 4.13Verification and improvement of food safety management system

Food safety plan for the carbonated beverage process and internal audit plan were developed to complete the verification procedure of food safety management system.

Process step	Monitoring parameters	Frequency	Responsibility	Standard	Records	Remarks
l. Raw matcrial						T I
receiving						
Raw water	Chemical quality	()nce a month	QAM	SLS 614 part		
	Microbiological quality	()nce a week	WVD	SLS 614 pan2	OPRP/RRI/01/Form	
	phi of raw water	()nce a day	Lab assistant	SLS 614 parl	<u> </u>	_
	Few chemical parameters	Once a week	Lab assistant	SLS 614 parl		
	(pH.turbidity.conductivity)				_	
• Sugar	OPRP/RRI/03/List	~				
· ().	OPRP/RRI/02/1.ist		QAM.			
<ul> <li>Flavours,</li> </ul>			Lab assistant,	Supplier		
Emulsions,	<b>OPRP/RRI/03/List</b>		Supervisor.	assessment	OPRP/RRI/03/Form	
Acids, colours	28	く Once	Receiving		<u> </u>	
& preservatives	OPRP/RRI/04/List	received	officer			
PET bottles &     lids	OPRP/RRI/05/List					
? Raw water filtration Pressure of the gauge	Pressure of the gauge	Once a day	Supervisor		•	

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Table 4.7: Food safety plan for the carbonated beverages process

Food safety plan for the carbonated beverages process

	Turbidity of water	Once a day	Lab assistant	SLS 516 part 1	OPRP/RRI/02/Form	
		~		SLS 614 part 2		
3. Water UV	Function of UV bulb	Hourly	Supervisor	ł	Working hours of	
sterilization					UV bulb - log sheet	
	Changing of the UV bulb	Replace the	Factory	·	Changing of UV	
		UV bulb after	engineer		bulb - log sheet	
		finishing			<b></b>	
		recommended				
		working				
		hours(9000hrs).				
4 Water chilling	Chilling temperature	Hourly	Supervisor	•	OPRP/CPM/01/Form	;
5. Sugar dissolving	Physical observation (total	Every batch	Supervisor	•	•	- 
	dissolved sugar)					
6. Sugar filtration	Pressure of the gauge	Once a day	Supervisor	e	•	
7. Ingredient weighing	Weight of the ingredients	Every batch	Supervisor	I	OPRP/SRM/01/Form	
8. Mixing ingredients	Brix. Temperature and pH	Every batch	Supervisor	ı	OPRP/SRM/02/Form	
with sugar solution						
9. Symp chilling	Chilling temperature	Hourly	Supervisor	8	OPRP/CPM/02/Form	
10 Carbonation &	Gas volume, brix, pH	Hourly	Lab assistant	¢	OPRP/CPM/03/Form	
		, I				

afjusting brix						
11.Filling & capping	Capping strength	Hourly (	Supervisor	•	Machine maintenance records	
12. Finished product						
<ul> <li>Immediately</li> </ul>	Physical quality	Every batch	Lab assistant	OPRP/FPI/02/1.ist	OPRP/FPI/01 Form	
after the	Chemical quality	Once a month	QAM	SLS 183: 1997	OPRP/FPI/01/Form	
production	Microbiological quality	Once a week	QAM	SLS 183: 1997	OPRP/FPI/01/Form	
At market place	At market place Chemical quality	Once in three	QAM	SLS 183: 1997	OPRP/FPI/02/Form	
		month				
	Microbiological quality	Once in three	QAM	SLS 183: 1997	OPRP/FPI/02/Form	, .
After shelf life		month				
	Chemical quality	Once in three	QAM	SLS 183: 1997	OPRP/FPI/03/Form	
		month				-
	Microbiological quality	Once in three	QAM	SLS 183: 1997	OPRP/FPI/03/Form	
		month				

### Internal audit plan

Food safety team leader should plan the audit and he is responsible for appoint the auditors and auditees. Auditors have responsible for give the answers for auditee's questions. Auditees should be appointed from other department and he should be prepared questions before starting the audit. Internal audit should be arranged once in three months.

Once finished the internal audit, food safety team leader arrange the opening meeting with the auditees to find out non conformities that are having with each departments. Once audit is completed, get auditee to specify the corrective action to be taken for each non- conformity giving the required time period will be corrected. Finally audit findings and corrective action reports should be return to the management representative.

Department	ISO 22000:2005	Auditee	Auditors		T	ime	
	Relevant Clauses			March	June	Sep.	Dec.
1.Production							
-							
NCR						1	<b>†</b>
2. Quality Assurance Dpt.							
NCR							
3. Receiving and purchasing Dpt							
NCR			•				
4. Stores							
NCR			<b>↓</b>	<b>↓</b>			

 Table 4.8: Internal audit schedule

### **CHAPTER 05**

### **Conclusions and Recommendations**

### 5.1 Conclusions

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- PRPs were developed by considering each and every process steps and production areas.
- By identifying biological, chemical and physical hazards in the carbonated beverage process, following Control Points (CPs) and Critical Control Points (CCPs) were determined.
  - Control Points (CPs)

Raw material Receiving Ingredient weighing Sugar dissolving Sugar filtration Water filtration Carbonation Storage and dispatch of bottles

- Critical Control Points (CCPs)
  - Raw water sterilization
  - Bottle capping
- Operational prerequisite Programs (OPRPs) and Hazard Analysis Critical Control Point (HACCP) plan were established according to the identified CPs and CCPs.
- Under the process capability analysis of CO<sub>2</sub> gas volume, brix and pH; cola, mandarin and cream soda production processes were identified as barely acceptable processes and soda, lemonade production processes should be improved according to their specifications.
- Total coliform count and E.coli count in raw water was not detected. According to aerobic plate count results, process capability of beverage process is inadequate. So improvements should be made.
- Food safety plan and internal audit plan was developed to verification of the food safety management system.

### 5.2 Recommendations

- PRPs, OPRPs and HACCP plan records should be maintained continuously.
- Introduction of machine for capping strength measurement is important.
- Comprehensive food safety management system verification should be carried out by maintaining food safety plan and internal audit plan.
- Gap analysis needed to be done by identified requirements with ISO 22000:2005 food safety management system requirements for the successful implementation of ISO 22000 to the carbonated beverage process.

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## Shaa Cola Beverage (Pvt) Limited

### ISO 22000:2005

### Pre Requisite Programme Calibrations (C)

-	Issue Date:
	<b>Revision No:</b>
	Controlled Copy No:

Copy Holder	Copy No.
Quality Assurance Manager	1
Lab Assistant	2
Supervisor	3

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SHAA	ISO 22000:2005 Pre Requisite Programme (PRP) Calibrations	Section No: PRP 01 Date of Issue: Revision No: 00 Page No: 01 of 03
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APPROVED BY	NGNATURE	

### 1. CALIBRATION OF REFRACTOMETER, THERMOMETER, PH METER, PRESSURE GAUGE, ANALYTICAL BALANCE AND ELECTRIC BALANCE

### 1.1 Objectives

Shae Cola Beverage (Pvt) Limited shall establish, implement and maintain procedure for calibration of Refractometer, Thermometer, pH meter, Pressure meter, Analytical balance and Electric balance to assist in reducing the likelihood of introducing food safety hazards to the product throughout the work environment

### 1.2Scope

Calibration procedure is applies to the Refractometer, Thermometer, pH meter, Pressure meter, Analytical balance and Electric balance used in the syrup room, production area and laboratory

### 1.3 Responsibility

The overall responsibility to establish, implement and maintain the calibration procedure is vested with the Quality Assurance Manager while the functional responsibilities are vested with lab assistant and supervisors

### **1.4 Calibration** Policy

### 1.4.1 Third party calibrations

Refractometer, Thermometer, pH meter, Pressure gauge, Analytical balance and Electronic balance are calibrated once a year from an independent calibration laboratory.

### 1.4.2 List of calibrated instruments

PRP/C/01/List

Instrument	Location	Calibration	Calibration certificate no. & date	Responsible
1. Refractometer-1 Refractometer-2	Production	10 <sup>8</sup> - 32 <sup>8</sup> 50° - 65°	• • • · · · ·	•
2 Thermometer	Production area	0 °C - 100 °C		
3. pH meter	Laboratory	25-70		
4. Pressure gauge	Production	0 - 60 ps	· · · · · · · · · · · · · · · · · · ·	
5.Analytical balance	Laboratory	0 01 g - 20g	DW/1439/2008 2008 07 22	
6. Electronic balance 1	Syrup room	1 kg - 14 5 kg		
7. Electronic belance 2	Syrup room	1 kg - 300 kg	•	

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### 1.4.3 In house calibration

 $\mathcal{L}_{\mathcal{A}}$ 

pH meter and Refractometer is calibrated once a month by lab assistant

PRP/C/01/Form

### In house calibration reports - pH meter

Date of		Standards	,	Calibrated	Remarks
calibrated	pH 2.00	pH 4.00	pH 7.00	by	
		• · • •		· · · · · · · · · · · · · · · · · · ·	
		•			·
		•	• •	· · •	
		<b>↓</b> -	• -	•	•
		• · - ·	•••••	•	·

PRP/C/02/Form

### In house calibration reports - Refractometer1

Date of calibrated	Brix 10 <sup>8</sup>	Standards Brix 20 <sup>8</sup>	Brix 30 <sup>8</sup>	Calibrated by	Remarks
	· _ ·	• •	<b>.</b> -		
		• •		· · · · · · · · · · · · · · · · · · ·	•
			4		
	)	• • •	<b>é</b>		• ·
	<b></b>	• *	٩		· _ · · · · · · ·
		• •	•		

PRP/C'/RJ/Form

## In house calibration reports - Refractometer2

Date of calibrate	d Brix 50 <sup>8</sup>	Standards Brix 60 <sup>8</sup>	Brix 65	Celibrated by	Remerks
		• - • • -	<b></b>	• = =	
-		• -	♦ •	•	• ·- ·- ·
		•-	•	•	∳
		•	•	• ·	۰ <b>۱</b>

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### **1.5 Related** documents

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- i. Calibration certificates
- ii. List of calibrated instruments PRP C 01 List
- iii. In house calibration reports (pH meter and Refractometer) PRP/C/01/Form, PRP/C/02/Form, PRP C 03 Form

Appendix B

# Shaa Cola Beverage (Pvt) Limited

### ISO 22000:2005

### Pre Requisite Programme Cleaning and Sanitizing (CS)

Issue Date: Revision No: Controlled Copy No:

Copy Holder	Copy No.
Quality Assurance Manager	1
Lab Assistant	2
Supervisor	3

SHAA	ISO 22000:2005 Pre Requisite Programme (PRP) Cleaning and Sanitizing	Section No: PRP 02 Date of Issue: Revision No: 00 Page No: 01 of 03
ISSUED BY	SIGNATURE	
APPROVED BY	NGNATURI	

### 2. CLEANING AND SANITIZING

### 2.1 Objectives

Shaa Cola Beverage (Pvt) Limited shall establish, implement and maintain program for cleaning and sanitizing to assist in reducing the likelihood of introducing food safety hazards to the product throughout the work environment.

### 2.2 Scope

Cleaning and sanitizing program applies to the raw material storage area, syrup room, beverage production site and finished product storage area

#### 2.3 Responsibility

The overall responsibility to establish implement and maintain the cleaning and sanitizing is vested with the Quality Assurance Manager while the functional responsibilities are vested with supervisors, lab assistant and cleaner.

### 2.4 Cleaning and sanitizing procedure

PRP/CS/01/List

Cleaning & sanitizing place/ item	How	Chemical used	Frequency	Who	records
1 syrup room & production area	<ol> <li>Remove food soil</li> <li>Wash &amp; rub using potable water</li> <li>Use detergent for further cleaning</li> <li>Wash with potable water until complete removal of detergents</li> </ol>	Detergent - Tepol	After the production	Helper	
2. water storage tanks, syrup tanks, pipelines & filling	1 Wash & rub using 10ppm chionne water 2 Wash with potable water until complete removal of chionne	Chlorine water	once a week	Helper	
Machine	* Syrup tanks, pipelines & filling machine wash with potable water until complete removal of flavours and colours	 	After the each production	, Helper	
3 filters	1 Pass 10ppm chlonne water through filters 2 Wash with potable water unbl complete removal of chlonne	Chionne weler	Once a month	Helper	

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4 equipment used for beverage production	<ol> <li>Remove food soil</li> <li>Wash &amp; rub using potable water</li> <li>Use detergent for further cleaning</li> <li>Rinse with potable water until complete removal of detergents</li> </ol>	Detergent - Tepol	After the production	Food handler	
5. PET bottles	Wash using potable water coming through pressure guns of the machine	Potable water	During production	Machine operator	
6 plastic creates	<ol> <li>Dip in to 20ppm chlonne water bath</li> <li>Rub using a brush</li> <li>Use detergent for further cleaning</li> <li>Wash with potable water until complete removal of chlorine &amp; detergent</li> </ol>	Chlorine water Detergent - Tepol	Once a week	Helper	
7 tile floor	1 Wet the tile with detergent mixed water 2 Rub the floor with a brush 3 Mop out until all detergent goes out	Detergent – tile cleaner	Once a day	Cleaner	
8 walls & caling	1 For tile walls follow the above procedure 2 For masonry wells and ceiling clean the cob webs and dust in a suitable way	Detergent - tile cleaner	Once a week	Cleaner	
9 giass	1 Wet the glass with detergent mixed water 2 Rub the glass with a punch 3 mop out with other punch until all detergent goes out	Detergent – glass cleaner	Once a week	Cleaner	

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### 2.5 Format for cleaning and sanitizing log – Syrup room & production area PRP/CS/01/Form

Cleaning and sanitizing log - Syrup room & production area

ľ

	Areas							
Dete	Water storage tank	filters	Syrup room	Syrup storage tank	Production area	Filling machine	Packing area	Sıg
				Ţ				
	<b></b>	÷		+		↓		
	<u> </u>		♦	<b>+</b>	4		+	
	<u>+</u>	- <b>+</b>	<b>∳</b> =	1	1			
							L	

### 2.6 Related documents

- i. Copy of detergent and sanitizer certificates
- ii. Cleaning and sanitizing procedure PRP/CS/01/List
- iii. Cleaning and sanitizing log Syrup room & production area PRP/CS/01/Form

Appendix C

## Shaa Cola Beverage (Pvt) Limited

### ISO 22000:2005

### Pre Requisite Programme Personal Hygiene (PH)

Issue Date: Revision No: Controlled Copy No:

Copy Holder	Copy No.
Quality Assurance Manager	1
Lab Assistant	2
Supervisor	3

1.

SHAA	ISO 22000:2005 Pre Requisite Programme (PRP) Personal Hygiene	Section No: PRP 03 Date of Issue: Revision No: 00 Page No: 01 of 03
ISSUED BY	SIGNATURE	
APPROVED BY:	SIGNATURE	

### 3. PERSONAL HYGIENE

### 3.1 Objectives

Shaa Cola Beverage (Pvt) Limited shall establish, implement and maintain program for personal hygiene to assist in reducing the likelihood of introducing food safety hazards to the product throughout the work environment regarding the personal who has direct contact with the product.

### 3.2 Scope

Personal hygiene program applies to raw material handlers in syrup room and beverage handlers in production site

### 3.3 Responsibility

The overall responsibility to establish implement and maintain the personal hygiene is vested with the Quality Assurance Manager while the functional responsibilities are vested with supervisors and lab assistant

### **3.4 Personal Cleanliness**

### 3.4.1 Bathing

- Workers must bathe daily
- Use of good soap is important to wash away sweat and dirt
- A good deodorant should be used after a bathe and undergarments should be changed every day

### 3.4.2 Hair

Hair can be a breeding ground for bacteria found on the skin. Employees are required to wear head coverings such as hair bands, nets or caps. It helps to;

Keep hair out of food

Prevents contamination by staphylococci

Prevent long hair from getting entangled in machinery

- Hair should be neatly tied if long.
- Hair length for men should be up to mid-ears
- . Hair should be shampooed regularly
- Moustaches and beards should be clean and trimmed
  - Men without moustaches and beards should be clean shaven

### 3.4.3 Eyes

- · Eyes must be kept clean and washed frequently
- · Rubbing of eyes should be avoided
- . An employee suffering from sore eyes should no be allowed to work

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### 3.4.4 Teeth and Mouth

- Teeth should be brushed twice a day
- Mouth should be rinsed well and gargling is a must after meal

### 3.4.5 Hands

Hands are possibly the most unsafe equipment in the chain of infection in the entire food production operation. Therefore;

- hands should be free from cuts and abrasions. It should be covered with waterproof dressing
- have nails manicured, preferably trimmed short and free from nail polish.
- refrain from wearing jewellery.

Hands should be washed.

- before beginning work and after a break
- before handling raw materials and foods
- after eating or smoking a cigarette
- after using the toilet
- after touching infected or unsanitary areas of the body or combing hair.
- after using a handkerchief, sneezing or coughing in to the hands.
- - after handling money
- + after scullery or any cleaning operation
- after handling waste food or refuse
- when ever hands are dirty

Proper hand washing procedure

- 1. Turn the water on and moisten the hands under the water
- 2. Apply liquid water using dispenser lathering well beyond the wrists and up the arms to the elbows
- 3. Pay special attention to the areas between the fingers and around the nails
- 4. Rub one hand against the other in a rotating motion using friction
- 5 Rinse thoroughly under the running water, allowing the water to flow from the elbows down to the fingertips
- 6. Turn water faucet off with a single service towel or your elbow or foot
- 7. Dry hands thoroughly with a hot air dryer or single service towel
- 8. Repeat this procedure as often as necessary

### 3.4.6 Uniforms, Caps and Masks

Each food handler is given 2 sets of uniforms, caps and masks. Before starting each shift they should wear washed and sanitized uniforms with caps and masks.

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### 3.4.7 Medical examination

All the food handlers who has direct contact with food are subjected to medical examination once a year to make sure that they don't have communicable diseases.

### 3.5 Monitoring of personal hygiene

PRP/PH/01/Form

### Personal hygiene check list

Month:....

Signature of inspector	h(30 columns)	Days per	Name	Employee Name
		• • • •	······································	
		• •		
-		•		

Checking for

í

- N Nails not Cleaned
- W Wounds and Earldoms
- H -No proper combing and protectig to hair
- 8 Relocated due to sickness
- J -Weanng hanging Jewellenes
- R Relocated one taken or production

\* If the food handler shows above signs the relevant cage should be marked with letter or letters, which has been given above

### **3.6 Related documents**

- I. Medical certificates of food handlers
- # Personal hygiene check ist PRP-PH/01 Form

Appendix D

## Shaa Cola Beverage (Pvt) Limited

### ISO 22000:2005

### Pre Requisite Programme Pest Control (PC)

Issue Date: Revision No: Controlled Copy No:

Copy Holder	Copy No.
Quality Assurance Manager	1
Lab Assistant	2
Supervisor	3
Security officer	4

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SHAA	ISO 22000:2005 Pre Requisite Programme (PRP) Pest Control	Section No: PRP 4 Date of Issue: Revision No: 00 Page No: 01 of 04
ISSUED BY	NGNATURE	-
APPROVED BY	SIGNATURE	

### 4. PEST CONTROL

### 4.1 Objectives

Shaa Čola Beverage (Pvt) Limited shall establish, implement and maintain pest control procedure to assist in reducing the likelihood of introducing food safety hazards to the product throughout the work environment

### 4.2 Scope

1-

Pest control program applies to the entire building and the surrounding area of the factory.

### 4.3 Responsibility

The overall responsibility to establish implement and maintain the pest control procedure is vested with the Quality Assurance Manager while the functional responsibilities are vested with cleaner, supervisors and lab assistant.

The pest control program of the company has been out soused to a licensed pest control and janitorial service which is Sadow service

### 4.4 Chemicals used for pest controlling

Pest	Name of the chemical	Dosage	Mixing level
Cockroaches	Propoxur	100 ml	Add water up to 4 L and mix
Flies	Coopex	100 g	Add water up to10 L and mix
Mosquitoes	Baytex Ec & Cyslin Ec	50 mi	Add water up to 8 L and mix
Ants	Coopex	50 g	Add water up to 5 L and mix
Rat / Mice Rat Glue		50 g	Put the chemical in a Bart Tray and leave it in the area where the pest live
	Rat Glue	25 g	Paste the glue in a board and leave it in the area where the rats/mice live

SHAA	ISO 22000:2005 Pre Requisite Programme (PRP) Pest Control	Section No: PRP 4 Date of Issue: Revision No: 00 Page No: 02 of 04
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APPROVED BY	NGNATURE	

## 4.5 Manufacturers specifications of material/chemicals used for pest

Pest	Chemical	Formulation	Group	Antidote/Treatment	
1. Rats /	Fentrol	0 005% RB	Difeneacum	Vitamin K	
Mice			Anti Coagulant		
2.		20% Ec	Organo	Atropine Sulphate	
Cockroaches	Propoxur	Emulsified	Carbonate		
		Concentrate		(Don't induce vomiting)	
3.	RBT	50%	Ferstheige		
Mosquitoes		concentrate	Fernthrion	Symptomatic treatment	
4. Flies			WP		
	•	Wettable	Permethrin S	Symptomatic treatmen	
	Coopex	Powder			
		25%			
5.Flies		50%	• • • • • • • • • • • • • • • • • • • •		
	RBT	Emulsified	Fernthrion	Symptomatic treatme	
		Concentrate	1		
6.Flies ~				No specific	
		Cislin	25% Ec	Delta Methrin	Antidote

### 4.6 Method of controlling

### 4.6.1 Control of Rodents (Rats and Mice)

A plan for setting up baiting stations throughout the premises using latest rodenticides will be prepared by a trained technician of Sadow service Checking and replenishing of baits as well as setting up additional baiting stations will be identified by regular visits of pest controlling organization

### 4.6.2 Control of Cockroaches

Effective control is achieved by spraying a residual insecticide to all locations likely to be infused in the premises

### 4.6.3 Control of Ants

Spraying with a residual insecticide will be carried out to all locations likely to be infested internally and externally

### 4.7 Pest control operations

Pest control operations are carried out once a month by a trained technician of the pest control organization. All the departments are informed about the date and time of the pest control operations. All the staff members are made aware of the preceutionary actions to be taken prior to the pest control operations.

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APPROVED BY	SIGNATURE	

#### 4.8 identification of pests and their locations

The different types of pests and the potential locations for pest harborage are identified with the support of all the staff members. The form is made available in every department. This form is filled by staff members who notice pests or their locations and hand over to lab assistant. These records are used by pest control organization when they plan the programs.

PRP/PC/01/Form

### Pest identification form

Name of the pest or pest sign (droppings, foot marks etc.)	ـــــــــــــــــــــــــــــــــــــ	
Location	· · · · · · · · · · · · · · · · · · ·	
Date and time		
Action taken		
Name and signature of the stat member who notice the pest sign	f	
Signature of Lab Assistant	•	
Date reported to the pest control organization	₩ - •	
Corrective or preventive action	•	
Signature of pest controller	•	

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APPROVED BY	NGNATURI	

PRP/PC/02/Form

### Periodic check list of pest control program

Date:....

Pest control service

Treatment area	Treatment method	Checked by
1.Production	•	
- syrup room	1	
- labeling & packing		
2. Stores	İ	
- raw material storage	1	
- finish product storage		
3. Laboratory	• · · · · · · · · · · · · · · · · · · ·	
4. Security office	•	
5. Office	•	
4. Canteen	•	
5. Changing rooms	•	
6. Toilets	•····	

### 4.9 Related documents

- i. Copy of pest controller's certificate of registration
- ii Pest identification form PRP PC 01 Form
- m Periodic check list of pest control program PRP PC/02/Form

Appendix E

# Shaa Cola Beverage (Pvt) Limited

### ISO 22000:2005

### Pre Requisite Programme Suppliers Assessments (SA)

Issue Date: Revision No: Controlled Copy No:

Copy Holder	
Purchasing Manager	
Assistant	
iving Officer	

Copy No. 1 2 3

SHAA	ISO 22000:2005 Pre Requisite Programme (PRP) Suppliers Assessments	Section No: PRP 5 Date of Issue: Revision No: 00 Page No: 01 of 04
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APPROVED BY	NGNATURI	

## 5. SUPPLIERS ASSESSMENTS

## 5.1 Objectives

Shaa Cola Beverage (Pvt) Limited shall establish, implement and maintain procedure for Suppliers Assessment to assist in reducing the likelihood of introducing food safety hazards to the product throughout the work environment.

## 5.2 Scope

Suppliers Assessment procedure applies to all raw material suppliers.

### 5.3 Responsibility

The overall responsibility to establish implement and maintain the Suppliers Assessment procedure is vested with the purchasing manager while the functional responsibilities are vested with lab assistant and receiving officer

### 5.4 Method of Suppliers Assessment

The suppliers assessment questionnaire is circulated among suppliers by the purchasing manager twice a year for regular suppliers or at the time that new suppliers are recruited Filled questionnaires are evaluated by a supplier assessment committee nominated by the purchasing manager

The suppliers who are not certified for HACCP. ISO 9001 or at least for SLS product standards are audited by the nominated member of the company food safety audit team.

SHAA SSUED BY	ISO 22000:2005 Pre Requisite Programme (PRP) Suppliers Assessments	Section No: PRP 5 Date of Issue: Revision No: 00 Page No: 02 of 04
ISSUED BY	NGNATURE	
APPROVED BY	NGNATURE	

# 5.5 Suppliers Assessment Questionnaire

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PRP/SA/01/Form

Name of the company		· · · · · · · · · · · · · · · · · · ·
Contact person		
Address:	• •	· ··· · · · · · · · · · · · · · · · ·
Tel / Fax/ e-mail		
Name of the product supplied		··· · ·
Person responsible for food safety		···· · · · · · · · · · · · · · · · · ·

Yes	No	Not applicable
	L	
	1	
· ••••	• · · · · · · · · · · · · · · · · · · ·	
	Yes	Yes No

- -

\_ \_ . .

# Completed by Designation Date

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SHAA	ISO 22000:2005 Pre Requisite Programme (PRP) Suppliers Assessments	Section No: PRP 5 Date of Issue: Revision No: 00 Page No: 03 of 04
ISSUED BY.	NGNAIT RI	
APPROVED BY:	SIGNATURE	

# **5.6 Supplier Evaluation**

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The above questionnaires are evaluated by suppliers evaluation committee.

PRP/SA/02/Form

**Supplier Evaluation Form** 

Supplier name: Assessment date:

Questions	<b>Evaluation marks</b>			
		5	0	
1. Does the company operate a food safety management system? If yes attach a copy of food safety policy				
2. Has the company being certified for HACCP/ ISO9001/ SLS product certificate? If yes attach copy of certificates				
3. Do you evaluate the customers? What are the evaluation criteria?				
4. Does the company have products recall procedure? If yes attach a copy of it				
5. Is chemical/microbiological test carried out internal/external laboratory? If yes attach copy of reports.				
6. Did you carry out staff training in food safety during last 3 months? If yes attach copy of training records				
7. Do you carry out internal audit on food safety?				
8. Do you have specifications for all raw materials that you use for production of supplement product?	 			
9. Has the company being inspected by a legal enforcement agency in the last 12 months? If yes attach copies of relevant documents				
10. Do you have distribution procedure? If yes attach a copy of it		ĭ ♠	▲	

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Assessed by Varified by (purchasing manager)

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### **5.7 Suppliers Evaluation Criteria**

Suppliers who gain more than 65 marks are registered as raw material supplier for carbonate beverage production. However the supplier who gains less than 65 has to be audited by the company audit team.

### 5.8 Registered supplier list

The results of suppliers assessment is used for preparation of registered supplier list

PRP/SA/03/Form

#### **Registered supplier list**

Raw material	Registered supplier
	•
	·
••••••••••••••••••••••••••••••••••••••	

### 5.9 Related documents

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- i. Suppliers assessment filled questionnaire PRP/SA/01/Form with relevant documents
- ii. Supplier evaluation form PRP SA 02 Form
- iii. Registered supplier list PRP SA 03 Form

Appendix F

# Shaa Cola Beverage (Pvt) Limited

# ISO 22000:2005

# Pre Requisite Programme Waste and Sewage Disposal (WSD)

Issue Date: Revision No: Controlled Copy No:

Copy Holder	Copy No.	
Quality Assurance Manager		1
Lab Assistant		2
Supervisor		3

•

SHAA	ISO 22000:2005 Pre Requisite Programme (PRP) Waste and Sewage Disposal	Section No: PRP 6 Date of Issue: Revision No: 00 Page No: 01 of 02
ISSUED BY	NGNATURE	
APPROVED BY	SIGNATURE	

## 6. WASTE AND SEWAGE DISPOSAL

### 6.1 Objectives

Shaa Cola Beverage (Pvt) Limited shall establish, implement and maintain waste and sewage disposal procedure to assist in reducing the likelihood of introducing food safety hazards to the product throughout the work environment

### 6.28cope

Waste and sewage disposal program applies to the entire factory

#### 6.3 Responsibility

The overall responsibility to establish implement and maintain the waste and sewage disposal procedure is vested with the Quality Assurance Manager while the functional responsibilities are vested with cleaner and supervisors.

#### 6.4Waste water disposal

Waste water is directly connected to the waste water treatment plant and treated water is passed to the lake in the separated area away from the factory

## 6.5 Solid waste disposal

The entire solid wastes are collected separately. Non degradable solid wastes (damage PET bottles, lids, cans, bags, etc.) are recycled by sundry sale and degradable solid wastes are removed in to a separated place away from the factory premises which is under the supervision of Quality Assurance department. Solid wastes are removed from the production area twice a day

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APPROVED BY	NGNATURI	

PRP/WSD/01/Form

# Waste and sewage disposal check schedule

Month & week

Area	Mon.	Tues.	Wed.	Thu.	Fri.	Sat
1.Production	1	1	1		T	T
- syrup room		1		i		
- labeling & packing	·	} ◆	•	ĺ		+
2. Stores		•			1	
<ul> <li>raw material storage</li> </ul>	1			1 1		
<ul> <li>finish product storage</li> </ul>	•	•	↓ ↓	i	÷	+
3. Laboratory	1	1	•		<b>_</b>	
4. Security office		1		İ		
5. Office	•	•	•	•		
4. Canteen	•		•	•		
5. Changing rooms	T	-	•	•	· · · ·	
6. Toilets	• _ ·	•	- -		Ī _	L

- Fully removed
- Half 🛆
- Not removed ☆

# 6.6 Related documents

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i Waste and sewage disposal check schedule PRP WSD/01 Form

Appendix G

# Shaa Cola Beverage (Pvt) Limited

# ISO 22000:2005

# Pre Requisite Programme Water, Air and Energy Supply (WAE)

Issue Date: Revision No: Controlled Copy No:

Copy Holder Operational Manager Technical Executive

1.

Copy No. 1 2

SHAA	ISO 22000:2005 Pre Requisite Programme (PRP) Water, Air and Energy Supply	Section No: PRP 7 Date of Issue: Revision No: 00 Page No: 01 of 01
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APPROVED BY	NGNATURI	

# 7. WATER, AIR AND ENERGY SUPPLY

### 7.1 Objectives

Shaa Cola Beverage (Pvt) Limited shall establish, implement and maintain water, air and energy supply procedure throughout the factory to assist in reducing the likelihood of introducing food safety hazards to the products by means of water, air and energy and also ensure that availability and adequacy

### 7.2Scope

Water, air and energy supply procedure applies to the entire building and the surrounding area of the factory

#### 7.3 Responsibility

The overall responsibility to establish implement and maintain the water, air and energy supply is vested with the Operational Manager while the functional responsibilities are vested with Technical Executive and electricians

# 7.4 Method of water, air and energy supply

1. Water	Method of supply Water from tube well to storage tank and distribute throughout the factory	Records Cleaning and maintenance records of water distribution system and water test reports
2. Air	Natural air flow, Air ducts and Air condition machines	Preventive maintenance programs and records of air conditions and air ducts
3. Energy	Electricity from electricity board / generator	Preventive maintenance programs of the electric system

### 7.5 Related Documents

- Cleaning and maintenance records of water distribution system and water test
   reports
- ii. Preventive maintenance program documents and records of air conditions and air ducts
- iii. Preventive maintenance program documents of the electric system

Appendix H

# Shaa Cola Beverage (Pvt) Limited

# ISO 22000:2005

# Operational Pre Requisite Programme Raw material Receiving and Inspection (RRI)

Issue Date: Revision No: Controlled Copy No:

Copy Holder	Copy No.
Quality Assurance Manager	1
Lab assistant	2
Supervisors	3
Receiving officer	4

1

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	Raw material Receiving & Inspections	Page No: 01 of 07
ISSUED BY	SIGNATURE	1
APPROVED BY	NGNATURE	

# **1. RAW MATERIAL RECEIVING AND INSPECTIONS**

## 1.10bjectives

Shaa Cola Beverage (Pvt) Limited shall establish, implement and maintain raw material receiving and inspection procedure to assist in reducing the likelihood of introducing food safety hazards to the product throughout raw materials and transportation

# 1.2 Scope

Raw material receiving and inspection procedure applies to the

- + Water
- Carbon dioxide (cylinders)
- Sugar, acidualant and preservatives (bags)
- Colours, flavours, and emulsion (cans)
- PET bottles and lids
- Labels and corrugated boxes

# 1.3 Responsibility

The overall responsibility to establish implement and maintain the raw material receiving and inspection procedure is vested with the Quality Assurance Manager while the functional responsibilities are vested with lab assistant, receiving officer and supervisors

# 1.4Rew material inspection procedure

All the raw materials (apart from the water) are purchased form the approved list of suppliers and packaging materials are purchased from fully owned subsidianes of JC group of companies. Inspection criteria are described as mentioned in the following table. Raw materials are inspected by lab assistant and once the inspection is finished the relevant record sheet should be filled.

Other relevant raw materials are inspected by receiving officer and satisfactory, unsatisfactory conditions are mentioned in the invoice it self

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Raw material	Mode of delivery	Supplier	inspection criteria	Records
1.Water	Pipe lines	Tube well in the factory	OPRP/RRI/01/List	OPRP/RRI/01/Form OPRP/RRI/02/Form
2.Carbon dioxide (cylinders)	Lorry	Ceylon oxygen limited	OPRP'RR1/02/List	OPRP/RRI/03/Form
3.Sugar (bags)	Lorry	C W mackie PLC (sugar trading division	OPRP'RRI/03/List	OPRP/RRI/03/Form
4.Acidualant and preservatives (bags)	Lorry	CHEMaster International	OPRP RRI/03/List	OPRP/RRI/03/Form
6.Colours, flavours and emulsions (cans)	Lorry	ABmauri lanka (pvt) limited	OPRP RRI/04/List	OPRP/RRI/03/Form
7.PET bottles	Lorry	609 packaging (pvt) limited	OPRP RR1/05/List	OPRP/RRI/03/Form
8.Labels and corrugated boexs	Lorry	Fast printers, Midaya packaging (pvt) limited	OPRP/RR1/06/List	OPRP/RRI/03/Form

# 1.5 Format for water inspection records

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OPRP/RRI/01/Form

# Water inspection records

Date	Time	pH	Chemical	treatment) Microbiological Analysis	Comments	Signature
			• • • • •	·		
7.		•	•	-	·····	
····		-	• •	1		i
√S	atusfactory			Unsatisfactory		
Microbio √S	atisfactory	nalysis:	•	Unsatisfactory		
			1	PERMANENT REFERE	weeks I throng	•

SHAA	ISO 22000:2005 Operational Pre Requisite Programme (OPRP)	Section No: OPRP 1 Date of Issue: Revision No: 00 Page No: 03 of 07	
	Raw material Receiving & Inspections		
ISSUED BY	NGNATURE	-	
APPROVED BY	NUNATURE		
Data Time	(After treatment)	PRP/RRI/02/Form	

Date	Time	рН	Analysis	Analysis	Comments	Signature
		ļ	•	•		
·	<b>-</b>	ł	•	•		
			•	• • • • • •		
	I Analysia		i ▲ -	• •		L
✓S	atisfactory		X	Unsatisfactory		

× Unsatisfactory

✓ Satisfactory
 Microbiological Analysis:
 ✓ Satisfactory

· · .

1.6 Format for inspection records

OPRP/RRI/03/Form

-	- Inspection records						
Delivery date	Time	Product	Product condition	Supplier	Expire date	Comments	Receipt signature
	• •		•	•	• •		
		·····	• • •	•	•		
Product		•	•	•	• -		•
	tisfacto		×	Unsatisfac	ctory		

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APPROVED BY	NGNATURI	

### **1.7Water inspection criteria**

Raw water inspected results were recorded in OPRP RR101 Form Treated water (after filtration and sterilization) inspected results were recorded in OPRP/RRI/02/Form

Chemical Analys	is		OPRP/RRI/01/List
Characteristics	Max. desirable level	Max. permissible level	Units
1. Colour	5 units	30 units	PCU
2. Odour	Unobjectionable	Unobjectionable	
3. Taste	Unobjectionable	Unobjectionable	
4. Turbidity	2 units	8 units	NTU
5. pH range	50-65	45-70	
6. Conductivity	750	3500	µs/Cm
7. Chloride	200	1200	mg/i
8. Alkalinity	200	400	mg/l

Microbiological Analysis	
Test	Acceptance limit
1. Aerobic plate count per ml (48 hrs at 37 °C) (test is carned out according to SLS516:part1 1991)	< 1 0 × 10 <sup>4</sup>
2. Total coliforms count(MPN)per 100ml (test is carried out according to SLS614 part2 1983)	<10 per 100ml
3. E-coli count (MPN)per 100ml (test is carried out according to SLS614 part2 1983)	None in 100ml

# 1.8 Carbon dioxide (cylinders) inspection criteria

Observe the below mention parameters and recorded in OPRP RRI03 Form 2

OPRP RRI021.ist

- + Leakages
- Carbon dioxide composition(mention in the label)
- Expire date(mention in the label)
- Weight of the cylinder(mention in the label)
- Any damages

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	Raw material Receiving & Inspections	Page No: 05 of 07
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APPROVED BY	SIGNATURI	

#### **1.9 Sugar, acidualant and preservatives (bags) inspection criteria Observe the below** mention parameters and recorded in OPRP/RRI/03/Form

• Correct weight(mentioned in the label)

OPRP/RRI/03/List

- Contaminations with dust, sand etc.
- Expiry date(mention in the label)
- Any damages
- Moisture content(mention in the label)

### Characteristics of sugar

Product name	White – fine granulated refine sugar
Colour	Max 45 ICUMSA
Moisture	Max 0 04%
Polarization	Min 99 9%
<b>Reducing sugars</b>	Max 0 04%
Sulphate ash	Max 0 02%
Mean aperture	Max 0 7 mm
Packaging	In 50 kilos net polylined polypropylene bags with machine stritched and fit for rough handling

### **Characteristics** of acidualant

•

Product name Appearance	Ascorbic acid (E 30 White crystalline po	-
Particle size		min 95%
	through 100 mesh	min 5%
Melting point	about 190°C	
Loss of drying	=<0 1%	
pH (5% w/v)	21-26	
Deaduct come	Citric acid (E.330)	

Product name Citric acid (E 330) Appearance White crystalline powder

Product name	Sodium citrate (E 331)
Appearance	White crystalline powder

# Characteristics of preservatives

Product name	Sodium benzoate (E 211)
Appearance	White powder
Product name	Sodium metabisulphate (E 223)
Appearance	White powder

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APPROVED BY	SIGNATURE	

# **1.10 Colours, flavours, and emulsion (cans) inspection criteria Observe the below mention parameters and recorded in OPRP/RRI/03/Form**

•	ate(mentioned in the label)		PRP/RRI/04/List
<ul> <li>Leakage</li> </ul>	8		
Swelling			
Rust			
Dent			
	al odour/ colour/ texture		
<ul> <li>Appeara</li> </ul>	nce		
Characteristic	s of Colours		
Colour type	sunset yellow (E 110)	Colour type	tartrazine (E 102)
Colour type	caramel colour		
Characteristic	s of flavours		
Flavours type	dalandan flavour (nature i	dentical)	
Appearance	clear pale yellow green lic		
	•••••••••••••••••••••••••••••••••••••••		
Flavours type	natural lemon flavour		
Appearance	clear pale yellow liquid		
Compounds	lemon extract (R)-p-ment	ha-1,8-diene	
Flavours type	cola part B flavour		
Appearance	dark brown colour liquid		
Compounds	ortho phosphoric acid, ca	Teine	
Flavours type	tamarınd flavour		
Appearance	dark brown colour liquid		
Compounds	2-furaldehyde		
Flavours type	cream soda flavour		
Appearance	clear light yellow liquid		
Compounds	polypropylene glycol		
Characteristic	cs of emulsions		
Emulsion type	· cola emulsion		
Appearance	dark brown colour liquid		
Compounds	sulphur dioxide		
Emulsion type	neutral clouding flavour		
Appearance	white emulsion		
Compounds	orange, sweet, ext		

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# 1.11 PET bottles and lids inspection criteria

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Observe the below mention parameters and recorded in OPRP/RRI/03/Form

OPRP/RRI/05/List

0		Lloinht	Colour	Circum	stances
Category	Weight	t Height Colour Body area	Neck area		
350 ml	22 3 g	18 5 cm	white & green	18.5 cm	8.0 cm
500 ml	28 3 g	23 0 cm	white & green	21 0 cm	8.0 cm
1500 ml	48 0 g	30 5 cm	white & green	28.5 cm	8.0 cm

Lids		_		
Colour	Weight		Diameter	Circumstance
Green, red, orange. violet, ash and brown	2 8 g	·	3 2 cm	9.5 cm
VIOLET, BEIT BILD OLOWIT		•		A

# 1.12 Labels and corrugated boxes inspection criteria

Observe the below mention parameters and recorded in OPRP/RRI/03/Form

OPRP/RR1/06/List

Labels			
Category	Length	Width	Diameter
350 ml (s*)	190 cm	4 0 cm	9.5 cm
350 ml (p*)	195 cm	4 0 cm	975 cm
500 ml (p*)	216 cm	4 5 cm	10 8 cm
1500 ml (p*)	30 3 cm	7 2 cm	15 15 cm
e*_ebriok label	p* - print label		• • • • • • • •

# Compared boxes (3 Ph)

Category	Length	•	Width	Height
350 ml	36 2 cm	- • -	23 7 cm	193 cm
/300 ml	38 9 cm	i	27 0 cm	237 cm
1500 ml	368 cm	•	27 2 cm	32 0 cm

# 1.13 Related documents

- i Registered supplier list PRP SA 03 Form
- i Water inspection records OPRP RRI01 Form and OPRP/RRI/02/Form
- m Other raw material inspection records ()PRP RRI/03 Form

Appendix

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# Shaa Cola Beverage (Pvt) Limited

# ISO 22000:2005

# Operational Pre Requisite Programme Syrup Room Management (SRM)

Issue Date: Revision No: Controlled Copy No:

1 m	
Copy Holder	Copy No.
Quality Assurance Manager	1
· · ·	<b>`</b>
Lab assistant	•
Supervisors	3

SHAA	ISO 22000:2005 Operational Pre Requisite Programme (OPRP)	Section No: OPRP 2 Date of Issue: Revision No: 00
	Syrup Room Management	Page No: 01 of 03
ISSUED BY	SIGNALL RE	1
APPROVED BY	SIGNATURE	

## 2. SYRUP ROOM MANAGEMENT

### 2.1 Objectives

Shaa Cola Beverage (Pvt) Limited shall establish, implement and maintain syrup room management procedure to assist in reducing the likelihood of introducing food safety hazards to the product throughout the syrup room including syrup preparation area and storage tanks

### 2.2 Scope

Syrup room management procedure applies to the entire syrup room including syrup preparation area and storage tanks

### 2.3 Responsibility

The overall responsibility to establish implement and maintain the syrup room management procedure is vested with the Quality Assurance Manager while the functional responsibilities are vested with lab assistant and supervisors.

#### 2.4Syrup room operational practices

Workers should follow the below mention operation practices to work efficiently

- . Flow the 'FIFO' (First In First Out) method when using raw material for syrup preparation
- + Half of the sugar/preservatives/acidualant bags should be seal well and keep in a dry place
- Colours, flavours and emulsion cans should be closed tightly
- Separate syrup storage tanks should be maintained for each category
- · Sugar dissolving should be done throughout the required time period and do not store over time in to the tank

# **2.5**Cleaning and Sanitizing

After the preparation of syrup, all the equipments and entire syrup room should be cleaned well

Once finish the production as well as before start the production, all the pipe lines and machines should be washed properly

Procedure reference No PRP 02

## 2.6 Ingredient weighing

All the ingredients used for preparation of each syrup should be weighed accurately by the trained person or experienced person and record it under OPRP/SRM/01/Form

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	Syrup Room Management	Page No: 02 of 03
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APPROVED BY	NONATURE	

# 2.7 Calibration of Electric balance, pH meter, Refractometer and Thermometer

Procedure reference No PRP 01

## 2.8 Sugar syrup filtration

Prepared sugar solution should be filtered using 0.5 µm filter and filters should be change according to the PRP 02

#### 2.9 Syrup operational conditions in the storage tanks

- I Brix range of syrup storage tank should be 60° 65°
- ii Temperature range of syrup storage tank should be 9°C 11°C
- iii. pH range of syrup storage tank should be 2 5 4 0

# 2.10 Format for amount of ingredients used per each syrup

OPRP/SRM/01/Form

Amount of ingredients list	
(per each syrup)	

Date	Time	Type of beverage	Ingredients	Amount	Signature (prepared person)
	•		•	÷	
	•	-	•	•	· · · · · · · · · ·
	•		• -	↓	
			•		
			•	÷	
			• • • •	•	······································
	• •	-	•	•	•

SHAA	ISO 22000:2005 Operational Pre Requisite Programme (OPRP)	Section No: OPRP 2 Date of Issue: Revision No: 00	
	Syrup Room Management	Page No: 03 of 03	
ISSUED BY	SIGNATURE	1	
APPROVED BY	SIGNATURE		

# 2.11 Format for syrup operational conditions monitoring log

Monitor the syrup operational conditions in the storage tank and record it under OPRP/SRM/02/Form by supervisor.

OPRP/SRM/02/Form

# Syrup operational conditions monitoring log

Date	Time	Syrup storage tank no.	Type of beverage	Brix	Temp.	рН	signature
	····· ··	• •	·	∳	···		
	· · · · ·	◆ ↓ ↓	• • · · -	• • •			
			• · · · · ·	∳ ∳	↓		

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# 2.12 Related documents

- I Amount of ingredients list OPRP SRM 01 Form
- I Syrup operational conditions monitoring log OPRP/SRM/02/Form

Appendix J

# Shaa Cola Beverage (Pvt) Limited

# ISO 22000:2005

# Operational Pre Requisite Programme Filtration Process Management (FPM)

Issue Date: Revision No: Controlled Copy No:

Copy Holder	
Quality Assurance	Manager
Lab assistant	
Supervisors	

SHAA	ISO 22000:2005 Operational Pre Requisite Programme (OPRP)	Section No: OPRP 3 Date of Issue: Revision No: 00
	Filtration Process Management	Page No: 01 of 02
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# 4. FILTRATION PROCESS MANAGEMENT

#### 4.1 Objectives

Shaa Cola Beverage (Pvt) Limited shall establish, implement and maintain filtration process management procedure to assist in reducing the likelihood of introducing food safety hazards to the product throughout the filtration

### 4.2 Scope

Filtration process management procedure applies to the raw water filtration and sugar syrup filtration system

#### 4.3 Responsibility

The overall responsibility to establish implement and maintain the filtration process management procedure is vested with the Quality Assurance Manager while the functional responsibilities are vested with lab assistant and supervisors.

#### 4.4 Raw water filtration process

Sand filter, Activated carbon filter, 0.5 micron filters and 0.2 micron filters are used for raw water filtration process

# 4.5 Sugar syrup filtration process

0.5 micron filters are used for sugar syrup filtration process

#### 4.6 Cleaning of filters

Before start the production, pressure gauge should be checked by the supervisor and if it is exceeded more than 1 bar filters should be cleaned

Sand filter and active carbon filter cleaning is carned out by washing the media using a reverse flow of water and backwashing the floc out of the filter to drain

## 4.7 Changing of filters

Filters should be changed if.

- + Visible particles in filtered water
- + Turbidity of filtered water is exceeded more than maximum permissible level
  - · Off tastes in filtered water
    - · Patches, tom areas or any damages in micron fifters

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# 4.8 Format for Changing of filters

OPRP/FPM/01/Form

# Changing of filters

Filters	month	(12 columns)	
Water filtration			
Multiple Media Filter			
Active carbon Filter			
0.2 Micron Filter			
0.5 Micron Filter			
Syrup filtration	· · · · · · · · ·		
0.5 Micron Filter	·····		

# 4.9 Related documents

- I. Changing of filters log sheet OPRP/FPM/01/Form
- ii. Water inspection records (After treatment) OPRP/RRI/02/Form

Appendix K

# Shaa Cola Beverage (Pvt) Limited

# ISO 22000:2005

# Operational Pre Requisite Programme Carbonation Process Management (CPM)

Issue Date: Revision No: Controlled Copy No:

Copy Holder	Copy No.
Quality Assurance Manager	1
Lab assistant	2
Supervisors	3

SHAA ISO 22000:2005 Operational Pre Requisite Programme (OPRP)		Section No: OPRP 4 Date of Issue: Revision No: 00
	Carbonation Process Management	Page No: 01 of 02
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APPROVED BY	SIGNATURE	

## **3. CARBONATION PROCESS MANAGEMENT**

## 3.1 Objectives

Shaa Cola Beverage (Pvt) Limited shall establish, implement and maintain carbonation process management procedure to assist in reducing the likelihood of introducing food safety hazards to the product throughout the carbonation.

### 3.2 Scope

Carbonation process management procedure applies to the carbonation section.

### 3.3 Responsibility

The overall responsibility to establish implement and maintain the carbonation process management procedure is vested with the Quality Assurance Manager while the functional responsibilities are vested with lab assistant and supervisors.

#### 3.4 Carbonation process handling practices

Trained person/ experienced person should be appointed for handling of carbonation process.

Water chiller temperature (10 °C) and syrup chiller temperature (8 – 9 °C) should be recorded in OPRP/CPM/01/Form and OPRP/CPM/02/Form hourly by supervisor.

Gas volume, Brix and pH should be recorded in OPRP/CPM/03/Form hourly by lab assistant and if any deviations occur from the inspection criteria under the OPRP/CPM/01/List, lab assistant should be informed to the supervisor/ appointed person immediately

Carbon dioxide cylinders should be changed by the technician under the supervision of the supervisor

# 3.5 Inspection criteria for each carbonated beverage

OPRP/CPM/01/List

Beverage type	Gas volume range	Brix range	pH range
mandarin	2.5 - 3.5	120-130	2.5 - 3 5
Cola	40-50	95-105	20-30
Lemonade	3.0 - 4.0	95-105	25-35
Tamarind	30.35	125-130	25-35
Cream soda	30-40	125-135	3.0 - 4.0
Soda	40-45	•	4.0 - 4 5

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	Carbonation Process Management	Page No: 02 of 02
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# 3.6 Format for inspection records

OPRP/CPM/01/Form

Inspection records (Water chiller)			
Date Time Temperature			

Inspection records (Syrup chiller)			OPRP/CPM/02/Form
Date	Time	Beverage type	Temperature

OPRP/CPM/03/Form

# Inspection records (During production)

Dete	Time	Beverage type	Gas volume	Brix	pH	Signature
						-
		· · · · · · · · · · · · · · · · · · ·				<u></u>

# 3.7 Calibration of Refractometer, Thermometer, pH meter, Pressure meter

Procedure reference No PRP 01

# 3.8 Related documents

i. Inspection criteria for each carbonaled beverage OPRP/CPM/01/List ii. Inspection records OPRP/CPM/01/Form, OPRP/CPM/02/Form and OPRP/CPM/03/Form

Appendix I.

# Shaa Cola Beverage (Pvt) Limited

# ISO 22000:2005

# **Operational Pre Requisite Programme Finished Product Inspection** (FPI)

**Issue Date: Revision No: Controlled Copy No:** 

Copy Holder	Copy No.
Quality Assurance Manager	1
Cab assistant	2
Supervisor	٦

SHAA	ISO 22000:2005 Operational Pre Requisite Programme (OPRP)	Section No: OPRP 5 Date of Issue: Revision No: 00
	Finished Product Inspection	Page No: 01 of 05
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APPROVED BY	SIGNATURE	

### **1. FINISHED PRODUCT INSPECTION**

#### 1.1Objectives

Shaa Cola Beverage (Pvt) Limited shall establish, implement and maintain finished product inspection procedure to assist in reducing the likelihood of introducing food safety hazards to the product throughout production, storage, transportation and market place.

#### 1.2Scope

Finished product inspection procedure applies to the

- Finished products (immediately after the production)
- Finished products (at the market place)

#### 1.3 Responsibility

The overall responsibility to establish implement and maintain the finished product inspection procedure is vested with the Quality Assurance Manager while the functional responsibilities are vested with lab assistant and supervisor

# **1.4Finished** product inspection procedure

# 1.4.1 Finished products (immediately after the production)

The number of bottles to be selected from a lot accordance with OPRP/FPI/01/List.

	OPRP/FPI/01/List
Scale of sampling	L
Number of bottles to be selected	Size of the sub sample
8	4
10	5
12	6
15	6
20	10
	Number of bottles to be

SHAA	ISO 22000:2005 Operational Pre Requisite Programme (OPRP)	Section No: OPRP 5 Date of Issue: Revision No: 00
	Finished Product Inspection	Page No: 02 of 05
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#### 1.4.1.1 Physical parameters

Samples (selected finished products accordance with OPRP/FPI/01/List) were inspected daily according to OPRP/FPI/02/List.

#### Parameters List

OPRP/FPI/02/List

- A. Label (accordance with SLS 467)
  - Name of the product Registered trade mark or brand name, if any Batch or code number Name and address of the manufacturer and/or distributor Volume of the contents in liters or milliliters and Date of expiry (optional)
- B. Leakages (Closer Leakage Test accordance with S L S 714 1985) Selected bottles were kept within 2 hours in up side down and checked for leakages
- C Damages (Drop Test accordance with S L S 714 1985) Selected bottles were dropped down.
  - a vertically in normal way (bottom touched with floor)
  - b vertically in up side down way (closure touched with floor)

#### 1.4.1.2 Chemical parameters

Samples (selected finished products accordance with OPRP/FPI/01-(List) were tested for chemical parameters once in three months from third party laboratory (SLS 183 1997)

Samples (selected finished products accordance with OPRP/FPI/01 1.ist) were tested for chemical parameters once in six months from Sn Lanka Standard Institution (SLS 183 1997)

OPRP/FPI/03/List

Preservatives		
Characteristic	Requirement	Method of test
1. Benzoic acid mg/kg. max *	160	SLS 581
2 Sulfur dioxide mg/kg, max **	70	SLS 581

• When a product contains more than one preservative the quantity of each preservative expressed as a percentage of the maximum permitted limit of the preservative shall be calculated. The sum of these percentages shall not exceed 100.

\*\* Products packed in metal containers shall not contain sulfur dioxide

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OPRP/FPI/04/List

Heavy metals	OFRE/FEI/04/LISt	
Heavy metals	Limit	Method of test
1 Arsenic, mg/kg, max	0.2	SLS 312
2. Copper, mg/kg, max	2	SLS 301
3. Lead, mg/kg, max	02	SLS 311

#### 1.4.1.3 Microbiological parameters

Samples (selected finished products accordance with OPRP/FPI/01/List) were tested for microbiological parameters once a month from third party laboratory (SLS 183 1997)

Samples (selected finished products accordance with OPRP.FPI.01 List) were tested for microbiological parameters once in three months from Sri Lanka Standard Institution (SLS 183.1997)

OPRP/FP1/05/List

Limit	Method of test
100	SLS 516 : pert 1
Absent	SLS 516 : pert 3
Absent	SLS 516 pert 2
	100 Absent

# 1.4.2 Finished products (at the market place)

Samples at the market place (selected finished products accordance with OPRP FPI:01/List) were tested for chemical and microbiological parameters once in six month from third party laboratory (SLS 183 1997)

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#### 1.5 Format for Finished product(immediately after the production) inspection records

OPRP/FPI/01/Form

#### Finished product (immediately after the production) inspection records

Date	Produ ct	Physical parameter				Chemical Microbiol- Remarks 1 parameters ogical				Signature
	type	A	B		C b	Preser- vatives	Heavy metals	parameter -s		
·					; ◆	· •	; •	· · · · · · · · · · · · · · · · · · ·		
			 	•	} ╃─── ♦───	, •	• • • • • • • • • • • • • • • • • • • •	• • •		
				∳	<b>↓</b>	•	•	• •		
Phys	cal Para	me				• •	<u>.</u>	<u>+</u>		

Satisfactory

× Unsatisfactory

#### Chemical Parameters: ✓ Satisfactory Microbiological Parameters: ✓ Satisfactory

- × Unsatisfactory
- × Unsatisfactory

# 1.6 Format for Finished product (at market place) inspection records

OPRP/FPI/02/Form

# Finished product (at market place) inspection records

Dete	Product type	1	Chemical Microbiological Remain parameters parameters	Remarks	Signature	
.,,,,,,,,,,,,,,,,,,,,,,,,,,,,,,,,,,,,,,	Preser- vatives	Heavy metals				
		· · · · · · · · · · · · · · · · · · ·	- •			
-	cal Parameti Satisfactory Nological Pa		= U	Insatisfactory	<u> </u>	<b>_</b>
	Satisfactory		= U	<b>insetsfactory</b>		

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SHAA	ISO 22000:2005 Operational Pre Requisite Programme (OPRP)	Section No: OPRP 5 Date of Issue: Revision No: 00
	Finished Product Inspection	Page No: 05 of 05
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#### 1.7 Identification of shelf life

Shelf life of the carbonated drink is six months. Samples were kept for a period of six months and tested for chemical and microbiological parameters to verify the shelf life.

OPRP/FPI/03/Form

Date	Product type	Cher		Microbiological parameters	Remarks	Signature
		Preser- vatives	Heavy metals			

**Chemical Parameters:** 

- ✓ Satisfactory
- × Unsatisfactory

#### **Microbiological Parameters:**

✓ Satisfactory

× Unsatisfactory

#### **1.8 Related documents**

i. Scale of sampling OPRP/FPI/01/List

ii. Physical, Chemical and Microbiological parameters OPRP/FPI/02/List, OPRP/FPI/03/List, OPRP/FPI/04/List, OPRP/FPI/05/List

iii. Finished product (immediately after the production) inspection records OPRP/FPI/01/Form

iv. Finished product (at market place) inspection records OPRP/FPI/02/Form

v. Shelf life verification records OPRP/FPI/03/Form

Appendix M

Date	Time	Brix	CO,	pН
1/10/2008	1 30	98	49	2 70
	3 30	10 0	46	2 63
2/10/2008	8 50	10 0	4 5	2 59
	10 40	98	4 5	2 96
	2 30	10 0	47	2 27
	4 00	10 0	4 1	2 49
8/10/2008	9 50	99	49	2 82
	10 55	10 0	46	2 88
	12 30	10 0	44	2 89
	1 30	10 0	4 6	2 88
	2 30	10 0	48	2 86
	3 40	10 0	4 5	2 75
10/10/2008	10 00	10 0	4 5	2 88
	11 40	95	4 6	2 4 1
	2 15	10 0	43	2 66
	3 45	10 0	46	2 53
16/10/2008	2 00	10 5	46	2 53
	3 00	10 3	46	2 4 1
	4 30	10 2	44	2 51
17/10/2008	7 55	11 0	48	2 49
	9 05	11 0	46	2 43
	11 30	10 0	4 8	2 85
	12 05	10 0	4 5	2 68
	1 35	10 2	44	2 84
	3 30	10 2	48	2 66
	4 30	10 0	42	•
27/10/2008	12 40	94	47	2 81
	2 50	10 5	4 5	2 78
	4 20	10 0	4 3	2 70
3/11/2008	2 50	10 2	48	2 90
	4 45	10 2	46	•
4/11/2008	8 20	92	46	2 32
	9 40	10 0	4 5	2 82
5/11/2008	8 30	98	42	2 80
	9 45	90	44	2 68
	11 20	94	48	2 86
	12 50	10 2	46	2 74
	1 50	10.4	48	2 79
	3 50	10 2	45	2 67
	4 50	10 0	46	2 56
6/11/2006	11 50	98	46	2 85
	2 30	98	48	2 72

# Brix, CO, gas volume and pH values for shaa cola

7/11/2008	8 30	10 0	47	2 49
	9 30	10 0	4 5	2 55
	10 40	10.4	44	2 85
	12 05	10 4	4 5	•
	1 40	10 0	44	2 45
	4 30	10 0	4 2	2 70
21/11/2008	9 15	94	4 5	2 25
	12 25	96	43	2 71
	2 25	10 0	4 2	2 56
	3 45	10 0	43	2 68
	4 40	10 0	42	2 46
22/11/2008	11 15	10 0	46	2 52
	2 30	10 0	4 2	2 87
	4 30	10 0	4 5	2 68
25/11/2008	8 50	10 0	5 0	2 38
	10 00	10 0	4 5	2 22
4/12/2008	8 30	10 0	47	2 59
	9 30	95	48	2 52
	11 00	10 0	4 0	2 79
	4 10	10 0	48	2 59
11/12/2008	1 50	10 4	4 6	2 68
	2 50	10 2	4 5	2 36
	4 15	98	4 5	2 60
13/12/2008	8 50	10 5	48	2 53
	1 15	95	43	2 72
	2 15	10.4	4 3	2 74
	3 30	10 2	4 6	2 57
17/12/2008	1 10	10 0	48	2 78
	2 40	10 0	4 5	2 65
	3 30	10 0	46	2 52
	4 50	10 2	4 5	2 65
20/12/2008	11 00	10 0	48	2 55
	12 10	10 0	46	2 65
	2 20	10 0	46	2 53
	4 05	10 0	45	2 25
23/12/2008	8 00	10 5	47	2 61
	11 00	10 8	46	2 64

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Date	Time	Brix	co,	pH
3/10/2008	9 35	12 5	29	3 20
	11 00	12 5	31	3 27
	12 30	12 5	29	3 17
	1 15	12 5	28	3 12
	2 50	13 0	3 0	3 00
	3 40	12 5	28	2 98
4/10/2008	8 4 5	12 2	29	3 27
	10 30	12 5	29	3 10
	2 30	12 6	27	3 19
	3 40	12 5	26	2 79
15/10/2008	9 50	12 5	3 0	3 11
	11 15	12 5	32	3 14
	2 15	11 9	30	2 84
	4 00	11.8	29	2 78
	4 55	12 5	29	3 00
3/11/2008	11 00	12 8	27	3 47
	12 20	12 0	26	3 40
13/11/2008	2 30	12 1	27	3 05
	3 30	13 0	27	2 85
	4 45	12 5	27	2 76
19/11/2008	8 50	12 5	27	2 66
	4 55	12 5	28	•
5/12/2008	10 45	12 2	28	2 88
	11 45	12 5	29	2 88
h	2 10	12 5	26	3 20
	3 20	12 8	29	3 00
22/12/2008	1 30	134	28	2 97
	2 40	12 5	25	2 95
	3 50	12 5	31	3 06
	4 50	12 0	27	3 05

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# Brix, CO, gas volume and pH values for Shaa mandarin

Date	Time	Brix	CO,	рH
1/10/2008	11 00		4 5	4 10
6/10/2008	11 30		4 3	4 40
16/10/2008	8 20		44	3 78
	10 00		4 4	3 79
	11 30		4 5	4 24
	1 00		4 5	4 22
18/10/2008	9 10		4 6	3 72
	12 30		4 6	4 22
30/10/2008	3 40		47	4 12
3/11/2008	1 45		4 5	4 21
6/11/2008	10 45		44	3 93
10/11/2008	12 50		4 6	4 02
	2 50		4 5	4 07
	3 55		4 5	3 93
	4 45		4 1	3 86
13/11/2008	9 05		42	3 71
24/11/2008	9 20		42	361
	1 40		4 5	3 76
8/12/2008	11 15		48	3 97
	12 15		47	4 13
9/12/2008	11 40		48	4 4 1
	1 30		29	•
	2 30		49	3 62
	3 40		34	4 32
10/12/2008	8 4 5		42	3 80
	9 50		43	4 00
16/12/2008	1 45		42	4 13
	3 50		4 5	•
17/12/2008	11 40		45	4 10
19/12/2008	11 20		4 5	3 86
	12 55		4 5	3 74

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# Brix, CO, gas volume and pH values for Shaa sode

PERMANENT REFERENCE Seberagemuwe l'inversity Library

Date	Time	Brix	CO3	рН
22/10/2008	11 00	12 5	38	3 32
	1 00	12 7	33	3 66
	2 20	13 0	30	3 89
	3 30	12 5	33	3 87
	4 45	12 5	31	•
31/10/2008	9 00	13 0	38	3 89
	10 15	12 7	35	3 75
	11 15	12 5	3 5	3 82
	1 00	12 8	32	3 77
1/11/2008	9 30	13 0	36	3 55
	11 50	12 9	33	4 01
	1 50	13.0	32	4 01
	3 00	12 5	31	3 93
	4 45	12 5	3 2	3 53
14/11/2008	9 55	12 8	3 5	3 58
	11 55	12 5	33	3 85
	12 55	13 2	33	3 86
	2 20	12 5	36	3 89
20/11/2008	9 45	12.8	33	3 94
	12 10	12 9	32	3 95
	2 30	12 5	32	3 36
	3 40	12 7	37	3 26
8/12/2008	1 00	12 5	32	3 62
	1 45	12 5	33	3 68
	2 50	12.8	32	3 75
	4 00	12 5	33	3 80
	4 45	13 0	33	3 96
9/12/2008	8 15	13 0	34	3 74
	9 15	12 5	3	3 57
	10 20	12 8	33	3 83
19/12/2008	4 45	13 0	33	3 22

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**1** 

Brix, CO2 gas volume and pH values for Shaa cream soda

Date	Time	Brix	CO2	pН
6/10/2008	2 30	9.8	3.8	3.30
	3.30	10.0	4.4	3.32
	4.40	10.0	3.5	2.71
7/10/2008	9.30	10.1	4.1	2.98
	11 00	10.1	4.2	3.17
	1.00	10.0	4.2	3.43
	2.40	10.0	3.9	3.24
	3.50	10.0	3.7	•
20/10/2008	1 50	9.5	3.6	•
	4.20	10.0	3.4	2.90
21/10/2008	8.55	9.5	3.4	3.00
	10.30	10.5	4.1	2.98
30/10/2008	9.10	11.8	3.0	3.26
	10 10	10.0	3.6	3.18
6/12/2008	8 30	10.2	4.0	3.08
	9.40	9.8	4.0	3.43
	11 20	10.0	3.9	3.25
8/12/2008	8 50	11.0	•	3.16
18/12/2008	4 20	10 0	3.3	3.05

1

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Brix, CO<sub>2</sub> gas volume and pH values for Shaa lamonade

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# Microbiological test results for raw water

Date	Aerobic plate count	Coliform	E coli
7/7/2008	94	none	none
	38	none	none
	32	none	none
	58	none	none
	33	none	none
3/9/2008	140	none	none
	160	none	none
	180	none	none
	140	none	none
	260	none	none
12/10/2008	270	none	none
31/10/2008	92	none	none
	160	none	none
	180	none	none
	58	none	none
	120	none	none
23/12/2008	120	none	none
	160	none	none
	150	none	none
	64	none	none
	110	none	none
26/12/2008	20	none	none
	13	none	none

1

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