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The Small Business FSMS Starter Pack has been designed to also include the key requirements of ISO 9001 for Quality Management Systems and ISO 22000 for Food Safety Management Systems and so is a good starting point for any small business.

Included in the Small Business FSMS Starter Pack:

- ✓ Food Safety Management Procedures A Comprehensive set of 28 top level documents that form the basis of your food safety quality management system
- √ Food Safety Records A set of 36 easy to use record templates
- ✓ GMP Manual A set of 10 Fundamental Good Manufacturing Practice Procedures

Food Safety Management System Procedures

A Comprehensive set of 28 top level documents that form the basis of your food safety quality management system:

- QM 001 Food Safety Quality Management System
- QM 002 FSQM Manual Summary
- QM 003 Document Control
- QM 004 Customer, Statutory and Regulatory Conformance
- QM 005 Record Control
- QM 006 Management Commitment
- QM 007 Quality and Food Safety Policy
- QM 007 Quality and Food Safety Objectives
- QM 008 Responsibility, Authority and Communication
- QM 009 Management Review
- QM 010 Resources and Training
- QM 011 Infrastructure and Work Environment
- QM 012 Product Realization and Contract Review
- QM 013 Design and Development
- QM 014 Purchasing, Orders and Verification of Purchased Materials
- QM 015 Prerequisite Programmes
- QM 016 Identification and Traceability
- QM 017 Customer Property
- QM 018 Planning Product Realisation and Contract Review
- QM 019 Calibration
- QM 020 Hazard Analysis and Critical Control Points System
- QM 021 Verification and Improvement
- QM 022 Customer Satisfaction
- QM 023 Internal Audit
- QM 024 Monitoring and Measuring QMS, Analysis of Data
- QM 025 Control of Non-Conforming Product
- QM 026 Corrective Action and Preventive Action
- QM 027 Crisis Management
- QM 028 Product Recall

Food Safety Management System Procedures



QM 001 Food Safety Quality **Management System**

Introduction

The company has planned, established, documented and implemented a food safety and quality management system for the site, which is maintained in order to continually improve its effectiveness in accordance with legislation, international standards and best industry practice. The company has planned and developed the processes that contribute to meeting the requirements of these standards and producing safe products.

The scope of the Food Safety Quality Management System includes a product categories, processes and activities conducted on site. These requirements are aligned with the policies and objectives of the site.

The Food Safety Quality Manual demonstrates due diligence of the company in the effective development and implementation of the food safety management system. These documents are fully supported by the completion of the records specified in this manual for the monitoring of planned activities, maintenance and verification of control measures and by taking effective actions when non-conformity is encountered.

The company is committed to supplying safe products for consumption. As part of this commitment, all products and processes used in the manufacture of food products are subject to food safety hazard analysis based on the Codex Alimentarius guidelines to the application of a HACCP system. All food safety hazards, that may reasonably be expected to occur, are identified by this process and are then fully evaluated and controlled so that our products do not represent a direct or indirect risk to the consumer. New information regarding food safety hazards is continually reviewed by the Food Safety team to ensure that the Food Safety and Quality Management system is continually updated and complies with the latest food safety requirements.

Should the company be required to outsource any process that may affect product conformity to the defined standards of the Food Safety

Document Reference QM 001 Food Saf Revision 2 13th July 2010 Owned by: Technical Manager Authorised By: Managing Director





QM 001 Food Safety Quality **Management System**

Quality Management System then the site will assume control over this process. This is fully defined in all Sub-Contract Agreements.

Communication

The company has established and documented clear levels of communication for suppliers, contractors, customers, food authorities and staff within the lood safety quality management system. Detailed communication arrangements and food safety communication responsibilities for all levels of management are contained in the food safety and quality manual. The scope of the communication procedures applies to all members of staff, both full time and temporary.

The Management Representative for Quality and Food Safety is the Technical Manager, who retains responsibility and authority for external communication and liaison regarding the food safety management system. This responsibility for communication extends to ensuring there is sufficient information relating to food safety throughout the food chain. This communication includes documented agreements, contracts, specifications, product information, food safety leaflets, allergen advice and reports.

Procedure

These processes and their interaction are documented within this manual and its procedures.

The top level procedures of the Food Safety Quality Management System Procedures are pre-fixed QM and are as follows:

QM 001 - Food Safety Quality Management System
QM 003 - PSQM Manual Summary
QM 003 - Document Control
QM 004 - Customer, Statutory and Regulatory Conformance
QM 005 - Record Control
QM 006 - Management Commitment
QM 007 - Quality and Food Safety Policy
QM 007 - Quality and Food Safety Objectives
QM 008 - Responsibility, Authority and Communication
QM 009 - Management Review
QM 010 - Resources and Training
Document Reference QM 003 Food Safety Management System





QM 019 Calibration

Introduction

The company has established, documented and implemented a Calibration system for monitoring and measuring equipment on site, which is maintained in order to ensure conformity to product requirements in accordance with international standards and best industry practice. The processes that contribute to meeting the requirements of these standards have been determined.

The scope of the Calibration System includes all equipment used to measure, monitor and manufacture product on site and activities conducted on site.

These requirements are aligned with the policies and objectives of the site and include those of the ISO 9001:2008 standard.

The company maintains this procedure for the calibration of monitoring and measuring equipment on site

An inventory of all monitoring and measuring equipment critical to product quality and safety or whose results can affect the conformity of product requirements is maintained by the Engineering Manager. Each piece of equipment is labelled with a unique identification code which is also used to identify it on all relevant documentation including calibration coefficients.

All of the Measuring and monitoring Equipment is subject to regular All of the Measuring and monitoring Equipment is subject to regular servicing and preventative maintenance as per the Preventative Maintenance Schedule for Critical Equipment. The Equipment is also covered by maintenance contracts with the supplier. Records of all work including maintenance, servicing and calibration of all equipment are maintained and retained on site for a minimum of 3 years.

All measuring and monitoring equipment on site is used and maintained in accordance with the instructions laid down in the manufacturer's handbooks/manuals. Operating and maintenance instructions are displayed or held next to the equipment. Monitoring and measuring

Document Reference QM 019 Calibration Revision 2 14th July 2010 Owned by: Technical Manager Authorised By: Managing Director



oument Reference QM 001 Food Safety Management System rision 2 13th July 2010 ned by: Technical Manager

Food Safety Management System Procedures



QM 020 Hazard Analysis and Critical **Control Points**

The company is committed to supplying safe products for consumption. As paid this commitment, all products and processes used in the manufacture of foo products are subject to hazard analysis based on the Codex Alimentariu HACCP principles.

The Food Safety Quality Manual demonstrates due diligence of the company in the effective planning, development and implementation of the food safety management system. These documents are fully supported by the completion of a HACCP plan and the records specified in this manual for the monitoring of planned activities, maintenance and verification of control measures and by taking effective actions when non-conformity is encountered. All food safety hazards, that may reasonably be expected to occur, are identified by this process and are then fully evaluated and controlled so that our products do not represent a direct or indirect risk to the consumer.

The Food Safety Management System is fully supported by established verification procedures and validation of the control measures/combinat control measures that are implemented through the operational Good Manufacturing Practices or the HACCP plan.

Management Commitment

We are a leading food company committed to produce safe and legal products in line with legislation and to continuously improve our standards of hygiene, quality and safety in relation to both our product range and the environment in which we manufacture these products.

HACCP is a system, which identifies specific hazards and implements measures for their control. All the HACCP's contained in this manual have been developed not write common. All the MACLE's contained in this manual have been devel taking legislation requirements into consideration and using the seven basic principles detailed below: -

Prepare a flow diagram of the steps in the process. Conduct a hazard analysis by identifying potential hazards. Assess likelihood of occurrence of these hazards and identify control options

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Authorised By: Managing Director





QM 020 Hazard Analysis and Critical **Control Points**

- Details of packaging Preparation and/or handling before use or processing Food Safety Acceptance criteria Intended use

All specifications are maintained, updated and approved by the Food Safety Team leader who identifies legal food safety requirements related to the items purchased. Raw material specifications are reviewed and updated if necessary when there is new design or redesign of the food safety management system.

The food safety team document the end product characteristics, including legal food safety requirements, for the purpose of conducting the Hazard Analysis. The product description includes:

- Product name
 Origin of ingredients
 What will the purchaser will do with it
 Details of the packaging
 How the product is processed or manufactured
 Composition of the product
 Chemical characteristics relevant for food safety such as pH or Aw
 Biological characteristics relevant for food safety treatment such as
 heating, freezing, brining or smoking
 Physical characteristics relevant for food safety
 Shelf life
 Prescribed storage temperature
 Prescribed storage conditions
 Intended use and reasonably expected handling
 Packaging

- Intended use and reasonably expected nationing Packaging Target consumers Possible unintended mishandling or misuse of the product Where the product is stored How the product is sold Labelling including instructions for handling, preparation and usage Prescribed delivery conditions

End product descriptions are reviewed and updated if necessary when there is new design or redesign of the food safety management system.

July 2010 ned by: Technical Manager norised By: Managing Directo





QM 023 Internal Audits

Introduction

The company has established, documented and implemented an internal audit system, which is maintained in order to verify the Food Safety Quality Management System is effectively implemented and updated and complies with planned arrangements legislation, international standards and best industry practice.

The scope of the Internal Audit System includes all products manufactured on site and activities conducted on site including the food safety quality management system.

These requirements are aligned with the policies and objectives of the site and include those of international standard ISO 9001:2008.

The Senior Management has a total commitment to the food safety The Settion in transgement has a total commitment to the food safety quality management system and provides adequate resource in the form of trained and qualified personnel to carry out a comprehensive Internal Audit Schedule. Internal audits are performed to confirm that company management systems are working effectively and to promote continuous improvement. Our philosophy is simply audit, review and improve.

The Internal Audit Schedule is planned annually and is designed to comprehensively cover all areas of the Food Safety Quality Management system including procedures, policies and activities.

The Technical Manager draws up the Internal Audit Schedule based on the following criteria:

- Risk associated with the procedure or activity Results of Previous audits
- Number of Corrective Actions raised or outstanding Customer Complaint Analysis Number of Preventative Actions raised or outstanding Results of the Management Review



QM 028 Product Recall Procedure

The company has established, documented and implemented a Product Recall Procedure for the site which is maintained in order to ensure products found to have major defects are withdrawn from the market in an efficient manner to minimise the risk to the consumer.

The scope of the Product Recall Procedure includes all products manufactured on site and activities conducted on site.

This procedure details the action that should be taken if for any reason a defective product reaches a customer. The action taken would depend upon the nature of the defect. A customer is defined as anyone who receives any product that is sold by the company.

Should non-conforming product be delivered to a customer causing a potential product recall then this is reported immediately to Technical Manager. The Technical Manager assesses the situation and may chose to contact the customer for a concession or if the non-conformity relates to a food safety hazard outside of acceptable limits instigate the Initial Procedure of a Product Recall.

The handling of customer complaints is categorized into non-critical and critical. Non-Critical Quality complaints from customers are directed to the Customer Services Manager who co-ordinates the customer response with the Quality Manager.

Critical or Serious complaints such as a claim of alleged injury or poisoning are notified to the Technical Manager who will instigate a immediate investigation which may involve crisis and product recall

Critical complaint is defined as an unsafe product with an aspect of the product that will result in injury or illness to the customer. This includes metal or glass in the product, contamination with dangerous chemicals the presence of food poisoning bacteria or their toxins.

t Reference QM 023 Internal Audits Revision 2



Document Reference QM 028 Product Recall Procedure Revision 2 14th July 2010 Owned By: Technical Manager Authorised By: Managing Director



Food Safety Management System Records

A comprehensive range of 36 easy to use record templates including:

QMR 001	Management Review Minutes
QMR 002	Training Record
QMR 003	Product Release Record
QMR 004	Design and Development Records
QMR 005	Supplier Assessment Record
QMR 006	Validation Record
QMR 007	Identification and Traceability Record
QMR 008	Register of Customer Property
QMR 009	Calibration Record
QMR 010	Internal Audit Record
QMR 011	Records of Non-conforming Product
QMR 012	Corrective Action Request Form
QMR 013	Preventative Action Request Form
QMR 014	Supplier Self Assessment and Approval Form
QMR 015	Equipment Commissioning Record
QMR 016	Return to Work Form
QMR 017	Hygiene Policy Staff Training Record
QMR 018	Complaint Investigation Form
QMR 019	Prerequisite Audit Checklist
QMR 020	Knife Control Record
QMR 021	Knife Breakage Report
QMR 022	Goods in Inspection Record
QMR 023	Equipment Cleaning Procedure
QMR 024	Glass and Brittle Plastic Breakage Record
QMR 025	Metal Detection Record
QMR 026	First Aid Dressing Issue Record
QMR 027	Cleaning Schedule
QMR 028	Cleaning Record
QMR 029	Engineering Hygiene Clearance Record
QMR 030	Glass and Brittle Plastic Register
QMR 031	GMP Audit Checklist
QMR 032	Vehicle Hygiene Inspection Record
QMR 033	Outgoing Vehicle Inspection Record
QMR 034	Pre Employment Medical Questionnaire
	Visitor Questionnaire
QMR 036	Product Recall Record



QMR 002 Training Record

Name:	Employee Number:
Company Start Date:	Position:
Prior External Qualification(s), Skills & Experience :	

Period Training Required	Details of Internal Training or External Training Course	Dates of Training	Signed (Trainee)	Assessed as Competent Signed (Trainer)
Weeks 1 - 4	Induction			
	QMD 002 Quality Policy Briefing			
	QMD 003 Quality Objectives			
	Health and Safety Procedure			
	Records monitoring and control			
	Environment and Waste Management			
	Packing Procedure			
Weeks 5 - 13	Operating Procedure			
	Coding Procedure			
	Labelling Procedure			

Document Reference QMR 002 Training Record Revision 2 26 February 2010 Owned By: Training Manager Authorised By: Quality Manager





QMR 010 Food Safety Management System Audit Form

FOOD SAFETY MANAGEMENT SYSTEM AUDIT FORM				
DATE OF AUDIT		TIME OF AUDI	г	
PROCEDURE DOCUMENT O	OR AREA AUDITED			
MANUAL	DOCUMENT NUMBER	TITLE		ISSUE NUMBER
NON-CONFORMANCES FOU	ND (To be completed by	y auditor)		
ACTION TO BE TAKEN (To I	be agreed between audit	or and auditee with tim	escales)	
LOG CORRECTIVE ACTION	REQUEST NUMBERS	RAISED IN BOX BE	LOW:	
NAME (Auditor)	SIGNATURE	(Auditor)	DATE	
NAME (Auditee)	SIGNATURE		DATE	
ACTIONS COMPLETE AND	CORRECTIVE ACTIO	NS SIGNED OFF AUD		
NAME	SIGNATURE		DATE	



QMR 010 Food Safety Management System Audit Form

AUDI	TOR SYSTEM AUDIT RE	PORT
Area Conformances to requirements		
Opportunities for improvement		
Strengths and weaknesses		
Confirmation if the food safety management system is adequate in the area audited		
Recommendations for future audit planning		
Items to follow up on the next audit		
NAME (Auditor)	SIGNATURE (Auditor)	DATE

Revision Number	Summary of Changes made from previous revision	Requested By:	Authorised By
2	Update to meet the requirements of BRC Global Standard for Food Safety Issue 5	Quality Manager	Site Director

Document Reference QMR 010 Food Safety Management System Audit Form Revision 2 26 February 2010 Owned by: Quality Manager



Document Reference QMR 010 Food Safety Management System Audit Form Revision 26 February 2010 Owned by: Quality Manager



GMP Manual

A set of 10 Fundamental Good Manufacturing Practice Procedures:

- GMP 1 Hygiene and Housekeeping Management
- GMP 2 Management of Pest Control
- GMP 3 Control of Visitors and Sub-Contractors
- GMP 4 Management of Cleaning
- GMP 5 Despatch and Distribution
- GMP 6 Maintenance
- GMP 7 Hygiene Policy
- GMP 8 Glass Policy
- GMP 9 Raw Material Foreign Body Control Policy
- GMP 10 Nut Handling Procedure



The company has established, documented and implemented a Hygiene and Housekeeping Management System for the site, which is maintained as part of the GMP prerequisite programmes in order to meet the requirements of the Food Safety Quality Management System and ensure the safe production of products.

The scope of the Hygiene Management System includes all products manufactured on site and activities conducted on site.

These procedures are specifically designed to preserve the product and

- personal hygiene requirements
- dress code
- work wear supply and laundering
 glass, plastic, ceramic, blade control
 illness/foreign travel/medication reporting
- visitor and temporary staff screening
 pest control incident reporting
 clean as you go and cleaning procedures

Personal hygiene standards required are read and acknowledged by new inductees and a bulletin is posted in all production areas.

All personnel are required to comply with the company standards for personal hygiene, the hygiene policy and the hygiene code of practice.

Dress code standards are clearly displayed. The requirement to wear the correct colour coded work wear in production areas is briefed to all staff on induction. Compliance to dress code is monitored by the supervisory staff in each area. All protective clothing is designed to prevent product contamination. The captive footwear provided must worn in high risk production areas.

A daily change of work wear is provided to all staff. All work wear is professionally laundered at a Food Grade Laundry.

Document Reference GMP 1 Hygiene and Housekeeping Management Revision 2 5 July 2010 Owned by: Technical Manager Authorised By: Managing Director



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