

The Management Representative for Safety and Quality is the Quality Manager, who retains responsibility and authority for external communication and liaison regarding the safety and quality management system. This responsibility for communication extends to ensuring there is sufficient information relating to safety throughout the supply chain. This communication includes documented agreements, contracts, specifications, product information, 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 Quality Management System Procedures are pre-fixed QM and are as follows:

- QM 1.1 Management Commitment
- QM 1.1.1 Product Safety and Quality Policy
- QM 1.1.2 Resource Provision
- QM 1.1.3 Safety and Quality Objectives
- QM 1.1.4 Management Review
- QM 1.1.5 Management Review Communication
- QM 1.1.6 Communication
- QM 1.2 Responsibility and Authority
- QM 1.3 Job Descriptions
- QM 2 Hazard Analysis and Critical Control Points
- QM 3 Quality Management System
- QM 3.1.2 Document Control
- QM 3.1.3 Record Control
- QM 3.2 Internal Audits
- QM 3.3 Corrective Action and Preventative Action
- QM 3.4 Contractual Arrangements
- QM 3.5.1 Purchasing, Orders and Verification of Purchased Materials
- QM 3.5.2 Contract Services
- QM 3.6 Identification and Traceability
- QM 3.7 Product Recall and Withdrawal
- QM 3.8 Incident Management Procedure
- QM 3.9 Control of Non-Conforming Product
- QM 3.10 Management of Customer Complaints
- QM 4 Site and Building Standards
- QM 5 Vehicle Operating Standards
- QM 6.1 Equipment Standards
- QM 6.2 Maintenance
- QM 6.3 Calibration
- QM 6.4 Housekeeping and Hygiene
- QM 6.5 Waste Management

Document Reference Quality Management System QM 3

Revision 1 1<sup>st</sup> February 2017

Owned by: Quality Manager

Authorised By: Managing Director



The Criteria and Methods required to ensure that the operation and control of these processes are effective are documented in these procedures and records.

These procedures are supported by second tier documents specific to each area including:

- Work Instructions
- Specifications
- Inspection schedules
- Risk assessments
- Job Descriptions
- HACCP Plans
- Critical Control Point Monitoring Procedures

Measurement, monitoring and review are carried out by analysis of data in key areas including:

- Critical Control Point monitoring
- Inspections
- Complaints analysis
- Key Quality performance indicators
- Standard Exception Reporting
- Results of Inspections
- Results of Internal audits
- Results of External Audits

The company has assessed the resources required to implement, maintain, and improve the Quality Management System and these resources have been provided including:

- Skilled Personnel
- Suitable Materials
- Suitable Equipment
- Appropriate Hardware and Software
- Infrastructure
- Information
- Finances
- Audit resource
- Training resource

Action is taken in response to results in order to correct and prevent deficiencies and to improve the probability of achieving company objectives.

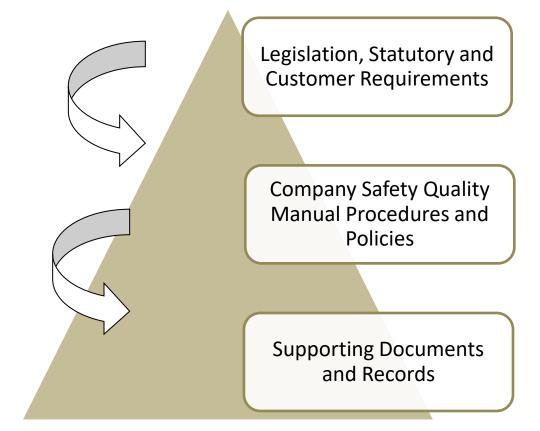
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#### **Document Hierarchy**





### **Quality System Process Diagram**



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