ISO 9001 Quality Manual

This manual complies with the requirements of the ISO 9001:2015 International Standard.

WaveTherm

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1 Introduction

Our Quality Management System Commitment

As the **President** of **WaveTherm**, I am committed to the quality management system, taking full accountability, and supporting other roles of leadership. Management uses the process approach and risk-based thinking to ensure the management system is integrated into our business processes to achieve intended results.

I am committed to provide the resources and training needed to ensure an effective quality management system that is necessary for our success and improvement. We provide a work environment that allows our employees to be successful in meeting our customers' needs.

The Quality Policy is established to be the driving force behind our quality management system, and I will continue to ensure that it remains compatible with the context and strategic direction of our organization.

David W. Mosier / President

WaveTherm

Quality Policy

WaveTherm Corporation strives to satisfy customer requirements through the development of innovative solutions, on-time delivery of products and services, and continual improvement of product quality and reliability.

2 Management System Approach

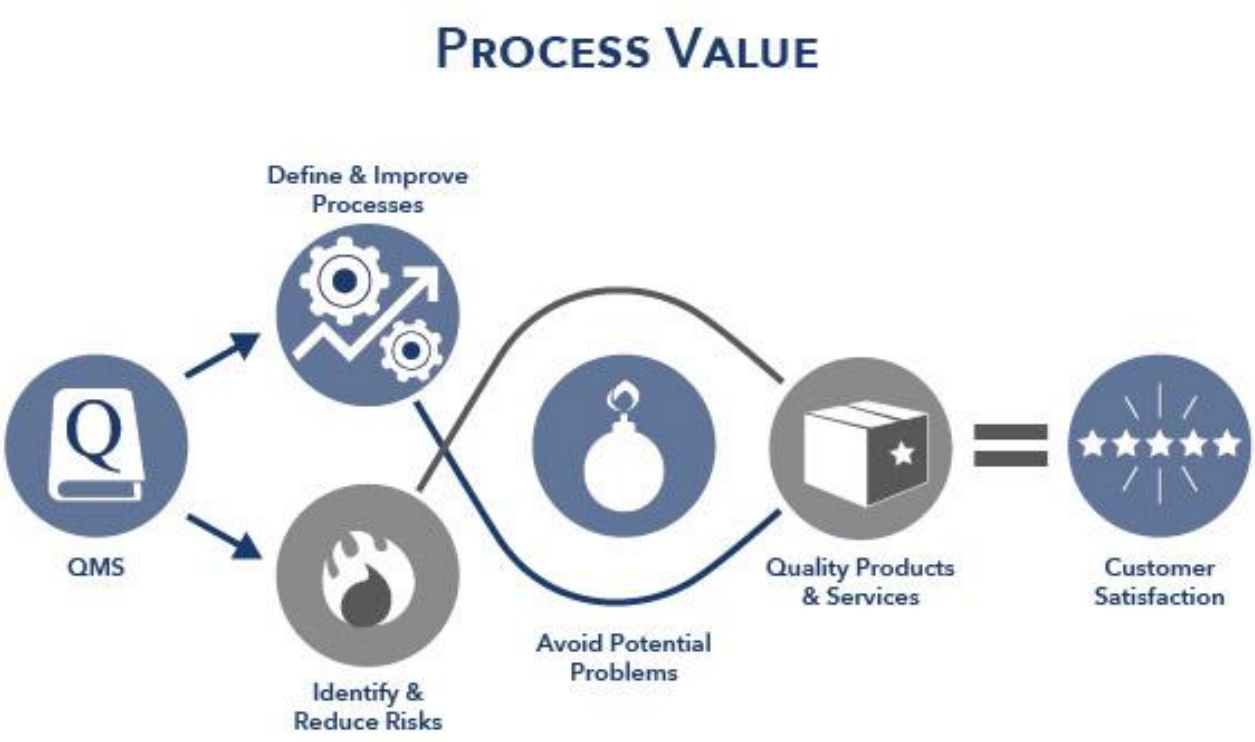
Our approach to our quality management system is based on the Plan, Do, Check, Act cycle (PDCA). The basis of our business beliefs is represented in **three** pillars:

Customer Focus

Our customers are the reason we exist. We aim to meet or exceed their needs and expectations to make them successful. We will even try to anticipate their needs and introduce solutions they have not seen before in the spirit of true partnership. Our success depends upon our customers' success.

Process Approach

To deliver on our commitment to total customer focus, we constantly work on our internal processes to maximize their effectiveness and efficiency. We recognize that it takes countless individual activities to deliver our products and services and that the process approach ties them all together. Our business is a process that transforms several inputs (customer requirements, resources, skilled employees, etc.) into an output that meets our customer's needs. Within our business are several key processes that make it all work. Our processes are dependent upon one another and individually need continual attention and improvement. We are constantly challenging ourselves to refine and change how we do things to reduce the time it takes to get something done with the least errors. When errors do occur, we use them as opportunities to learn and improve. We are never satisfied with how things are working now and strive to raise our game every day.



Risk-Based Thinking

Looking ahead to anticipate what could happen is the reason we employ risk-based thinking throughout our organization. At several points in our process, we purposely stop and ask two probing questions:

- "What could go wrong?"
- "Is there a way to improve?"

This perspective of constantly watching for risks and opportunities leads us to action which we carefully manage to ensure timely implementation and effective results. This gives us an attitude of being proactive to take advantage of every opportunity to improve.

We intend these three basic beliefs to cause our customers to stand up and take notice of the difference we provide to them on a daily basis. Our quality management system described in this Quality Manual has been carefully crafted to make these three pillars a real part of what makes us work.

3 Quality Manual Structure

This Quality Manual is presented in a PDCA manner and describes our approach to the requirements of ISO 9001:2015. The manual is divided into four sections with all applicable sub-clauses represented in each section as below:

NOTE: In the sections that follow, **Bold Blue Text** refers to related documentation where additional documentation is maintained and/or records are retained.

SECTION 1: PLAN

With an ever-changing world, we are faced with new challenges on a continuing basis. The issues, changes and trends within our industry and the broader economy present us with risks and opportunities from cultural, technological, competitive, regulatory, market, economic and social factors. Not only can these factors affect our business, but there are also other interested parties and organizations that we deal with on a day-to-day basis and these present additional requirements that we must account for.

All of these factors may affect our business negatively (risks) or positively (opportunities). The risks may be relevant to us and have the potential to affect our business or our customers in a negative way. These aspects of our business environment may also create opportunities for us to improve our organization or take advantage of expanded current or new business ventures.

Planning other aspects of our organization is also very important. Our planning process also includes people, their knowledge and training, infrastructure, environment, documented information, and communication. All planning efforts are structured, include decision-makers, and are documented when required.

Our extensive planning process puts us in the best position possible to forecast these challenges and take actions when necessary. It also establishes the needed foundations for us to provide our products and services.

4 Context of the Organization

4.1 Understanding the organization and its context

***Requirement**: Determine the external and internal issues that are relevant to the purpose and strategic direction and that affect the ability to achieve the intended result(s) of the quality management system.*

Our Approach: Issues (4.1) stemming from trends and changes in our industry may affect our business purpose and strategic direction. Those that present risks and/or opportunities are initially addressed by top management, recorded on the **Improvement Plan** and then monitored through the **Improvement Plan** and Management Review meetings.

4.2 Understanding the needs and expectations of interested parties

***Requirement:** Determine the interested parties, and their requirements that are relevant to the quality management system.*

Our Approach: On our **QMS Plan**, we identify requirements from relevant interested parties (4.2) that impact our ability to meet our customer and applicable statutory and regulatory requirements. Requirements are monitored on an ongoing basis through management activities and Management Review meetings.

4.3 Determining the scope of the quality management system

***Requirement:** Determine the boundaries and applicability of the quality management system to establish the scope, considering:*

- *external and internal issues;*
- *requirements of relevant interested parties;*
- *products and services.*

The scope is available and maintained as documented information stating the:

- *products and services covered by the quality management system;*
- *justification for any instance where a requirement of ISO 9001 cannot be applied.*

Our Approach: The contextual issues and interested party requirements are considered to determine the scope (4.3) of our quality management system:

Considering these external and internal issues and requirements, we have established the scope of our quality management system as:

Scope

The scope of the Quality Management System includes the major product and service categories associated with the primary functions of manufacturing wedgelocks, heatframes chassis and thermal management studies and solutions at their Raleigh North Carolina location and distributing the products to major aerospace and defense contractors

4.4 Quality management system and its processes

Requirement: *Establish, implement, maintain and continually improve the quality management system, including the processes needed and their interactions.*

For the processes needed, determine:

- *the inputs required, and the outputs expected;*
- *their sequence and interaction;*
- *the criteria, methods, including monitoring, measurements and related performance indicators needed to ensure their effective operation, and control;*
- *the resources needed and their availability;*
- *the assignment of the responsibilities and authorities;*
- *the risks and opportunities, and plan and implement the appropriate actions to address them;*
- *the evaluation and, if needed, the changes to processes to ensure that they achieve intended results;*
- *and improvement.*

Our Approach: The processes (4.4) needed to achieve intended outcomes, results and to continually improve our quality management system are identified on the **QMS Plan**, are maintained on **Process Plans**, and reviewed during Management Review meetings.

5 Leadership

5.1 Leadership and commitment

5.1.1 General

Requirement: *Demonstrate leadership and commitment with respect to the quality management system by:*

- *taking accountability for its effectiveness;*
- *establishing a quality policy and objectives that are compatible with the context and strategic direction;*
- *integrating the QMS requirements into business processes;*
- *promoting the use of the process approach and risk-based thinking;*
- *ensuring that the resources needed are available;*
- *communicating its importance and conforming to its requirements;*
- *achieving intended results;*
- *engaging, directing and supporting people to contribute to the QMS;*
- *promoting improvement;*
- *supporting other relevant management roles to demonstrate their leadership as it applies to their areas of responsibility.*

Our Approach: Our top management holds the ultimate responsibility for the quality management system. Our top management is dedicated and committed (5.1) to ensuring that our quality management system is effective, understood and improved.

Top management includes the following members:

- **David Mosier - President**
- **Rodney Bame – VP Engineering**
- **Josh Jenkins – VP Sales**

5.1.2 Customer focus

Requirement: *Demonstrate leadership and commitment with respect to customer focus by ensuring that:*

- *applicable requirements are determined, understood and consistently met;*
- *risks and opportunities that can affect conformity of products; services and enhancement of customer satisfaction are determined and addressed;*
- *focus on enhancing customer satisfaction is maintained.*

Our Approach: Top management demonstrates leadership and commitment to ensure that all applicable requirements are met, risks and opportunities are addressed, and the focus on customer satisfaction is maintained (5.1.2) through our **QMS Plan**, **Improvement Plan**, **Process Plans** and **Quality Policy**.

5.2 Policy

5.2.1 Establishing the quality policy

Requirement: *Establish, implement and maintain a quality policy that:*

- *is appropriate to the purpose and context and supports the strategic direction;*
- *provides a framework for setting quality objectives;*
- *includes a commitment to satisfy applicable requirements;*
- *includes a commitment to continual improvement of the QMS.*

Our Approach: The top-level requirement that directs our entire quality management system is our **Quality Policy.** It is reviewed at least annually during Management Review.

5.2.2 Communicating the quality policy

Requirement: *The quality policy:*

- *is available and maintained;*
- *is communicated, understood and applied;*
- *is available to relevant interested parties.*

Our Approach: The quality policy (5.2) is maintained and available in this **Quality Manual** and on the **QMS Plan**. It is communicated to appropriate staff. It is made available to interested parties upon request.

5.3 Organizational roles, responsibilities and authorities

Requirement: *Ensure that the responsibilities and authorities for relevant roles are assigned, communicated and understood.*

Assign quality management system responsibilities and authority for:

- *ensuring that it conforms to the requirements of ISO 9001:2015;*
- *ensuring that processes are delivering their intended outputs;*
- *reporting on its performance, and opportunities for improvement, to top management;*
- *ensuring the promotion of customer focus;*
- *ensuring that its integrity is maintained when changes are planned and implemented.*

Our Approach: Responsibilities and authorities (5.3) for our process owners are assigned, communicated and understood in our **QMS Plan, Process Plans,** and through management of our employees. The **Quality Manager** has been appointed as the Management Representative of our QMS.

6 Planning

6.1 Actions to address risks and opportunities

Requirement: *When planning for the QMS, consider the issues (4.1) and the requirements (4.2) and determine the risks and opportunities that need to be addressed to:*

- *assure that the QMS can achieve its intended result(s);*
- *enhance desirable effects;*
- *prevent, or reduce, undesired effects;*
- *achieve improvement.*

Plan:

- *actions to address these risks and opportunities;*
- *how to:*
  - *integrate and implement the actions into our processes;*
  - *evaluate their effectiveness.*

Our Approach: We respond appropriately to the risks and opportunities (6.1) identified in the **Process Plans** and **Improvement Plan,** and in the Management Review meeting.

The actions are integrated into our quality management system process and are evaluated for effectiveness during reviews.

6.2 Quality objectives and planning to achieve them

Requirement: *Establish objectives at relevant functions, levels and processes that:*

- *are consistent with the quality policy.*
- *are measurable.*
- *take into account applicable requirements.*
- *are relevant to conformity of products and services and the enhancement of customer satisfaction.*
- *are monitored.*
- *are communicated.*
- *are updated as appropriate.*

Retain documented information on the quality objectives.

Our Approach: We establish objectives (6.2) at relevant functions, levels, and processes on our **Measurement Plan**. The results of these objectives and plans to achieve them are reviewed and retained on the **Management Review Minutes**.

6.3 Planning of changes

Requirement: *Where needed, carry out changes to the Quality Management System in a planned manner considering:*

- *the purpose of the change and any of its potential consequences.*
- *the integrity of the quality management system.*
- *the availability of resources.*
- *the allocation or reallocation of responsibilities and authorities.*

Our Approach: Changes (6.3) that are needed are planned and carried out carefully considering the consequences, the integrity of our QMS, resources and associated responsibilities. The changes are managed and are recorded in the **Management Review Minutes** as appropriate for the change.

7 Support

7.1 Resources

7.1.1 General

Requirement: *Determine and provide resources needed for maintenance and continual improvement of the QMS considering:*

- *capabilities, constraints and existing resources;*
- *needs from external providers.*

Our Approach: During our Management Reviews, our top management discusses all internal and externally provided resources needed (7.1.1) for maintenance and continual improvement of our quality management system and ensures that they are provided.

7.1.2 People

Requirement: *Determine and provide the people necessary to effectively implement the QMS and for the operation and control of processes.*

Our Approach: During our Management Reviews and through daily management activities, our top management determines the people necessary (7.1.2) for the effective implementation of our QMS and for the operation and control of our processes and ensures that the resources are provided.

7.1.3 Infrastructure

Requirement: *Provide and maintain the infrastructure for the operation of processes and conformity of products and services.*

Our Approach: To ensure that our infrastructure resources (7.1.3) remain adequate, they are reviewed and discussed during Management Reviews.

7.1.4 Environment for the operation of processes

Requirement: *Provide and maintain the environment necessary for the operation of processes and to achieve conformity of products and services.*

Our Approach: Our top management ensures that our work environment (7.1.4) is sufficient to achieve conformity of our products and services as discussed during Management Reviews.

7.1.5 Monitoring and measuring resources

7.1.5.1 General

Requirement: *Provide the resources needed to ensure results when monitoring and measuring is used to verify conformity of products and services.*

Our Approach: We determine and provide the resources needed to monitor and measure (7.1.5.1) our products and services to ensure that they continue to meet requirements and specifications.

We ensure these resources are:

- suitable for the specific type of monitoring and measurement activities;
- maintained to ensure fitness for purpose.

This is documented in the **Measurement Plan** and reviewed during **Process Plan** audits**.**

7.1.5.2 Measurement Traceability

Requirement: *When measurement traceability is a requirement, or considered essential to provide confidence in the validity of the measurement results, measuring equipment is:*

- *calibrated or verified or both, at specified intervals, against measurement standards that are traceable to international or national standards. When no standards exist, the basis for calibrations needs to be documented.*
- *Identified in order to determine status;*

- *Safeguarded from adjustments, damage or deterioration.*

When measuring equipment is found to be unfit, the organization must determine if the validity of previous measurement results has been adversely affected and take appropriate action.

Our Approach: Measurement traceability is essential to providing confidence in the validity of the measurement results. Details of this process are maintained within the **Calibration Plan** and **Calibration Form,** and a list of monitoring and measurement equipment is maintained on the **Calibration Log.**

If measuring equipment is found to be unfit, management will take appropriate action, including initiating a **Corrective Action Request** if needed.

7.1.6 Organizational Knowledge

Requirement: *Determine the knowledge necessary for the operation of processes and to achieve conformity of products and services. Maintain this knowledge and, make it available to the extent necessary.*

When addressing changing needs and trends, consider current knowledge and determine how to acquire or access the necessary additional knowledge and required updates.

Our Approach: All current knowledge (7.1.6) sources, requirements, changes, needs and trends are determined by top management, maintained, and discussed during Management Reviews.

7.2 Competence

Requirement: *Determine the necessary competence of people doing work under organizational control that affects the performance and effectiveness of the QMS and:*

- *ensure they are competent on the basis of education, training, or experience;*
- *where applicable, take actions to acquire the necessary competence, and evaluate the effectiveness of the actions taken;*
- *retain appropriate documented information as evidence of competence.*

Our Approach: We determine the required competencies (7.2) for our employees, whose work may impact the effectiveness and performance of our QMS. We hire employees with specific knowledge, skills and education that best fit our needs and provide training to fulfill any missing competencies.

Evidence of this process is retained in employee files and maintained by the **Operations Manager**.

As of the initial release of this document, all current employees are considered to be competent.

7.3 Awareness

Requirement: *Make people doing work under organizational control aware of:*

- *the quality policy;*
- *relevant quality objectives;*
- *their contribution to the QMS and the benefits of an improved system;*
- *the implications of not conforming to the QMS requirements.*

Our Approach: People doing work under our control are made aware (7.3) of our quality policy, objectives, how our quality management system works and the implications of not working within our quality management system as defined on the **Communication and Awareness Plan**. This plan is reviewed during Management Review**.**

7.4 Communication

Requirement: *Determine all elements of internal and external communications relevant to the quality management system.*

Our Approach: Communication (7.4) is very important to our operation's success. Our communication methods are maintained on the **Communication and Awareness Plan**, which is reviewed periodically during Management Review.

7.5 Documented Information

7.5.1 General

Requirement: *The organization's quality management system will include: documented information required by the ISO standard and documented information deemed necessary by the organization for the effectiveness of the quality management system.*

Our Approach: We have determined which documented information (7.5) is necessary for the effectiveness of our quality management system and control as per the requirements listed below.

7.5.2 Creating and updating

Requirement: *The organization will ensure appropriate:*

- *identification and description (such as title, date, author or reference number);*
- *format (such as language, software version, graphics) and media (ex: paper or electronic);*
- *review and approval for suitability and adequacy.*

Our Approach: This documented information is identified, formatted, reviewed and approved, and controlled according to applicable requirements primarily through the use of our CORE Compliance Platform.

7.5.3 Control of documented information

7.5.3.1

Requirement: Control documented information so that:

- *it is available for use where and when it is needed;*
- *it is adequately protected (ex: from loss of confidentiality, improper use, or loss of integrity).*

Our Approach: The CORE Compliance Platform is a cloud-based system, so it is available as needed. To ensure data protection, the CORE Compliance Platform has the following security features:

Server and network security:

- encryption, where appropriate hardened operating systems and firewalls;
- regular security audits;
- rigorous systems reviews;
- state-of-the-art security related upgrades;
- a full time, dedicated security staff for software system maintenance, research and troubleshooting;
- secure facilities locations;
- tightly controlled and supervised server access.

CORE Compliance Platform security:

- unique usernames and passwords;
- encrypted transactions (including username and password submittals;
- company-specific company database realms (associated by username) with a limited database authority level based on customer defined roles;
- specific nature of user session and user authority control and encryption methods cannot be released.

For documents and records not maintained in CORE**,** we ensure data protection through periodic backup and password protection.

7.5.3.2

Requirement: *For the control of documented information, address the following:*

- *distribution, access, retrieval and use;*
- *storage and preservation, including preservation of legibility;*
- *control of changes (ex: version control;)*
- *retention and disposition.*

Documented information of external origin determined by the organization to be necessary for the QMS will be identified and controlled.

Documented information retained as evidence of conformity will be protected from unintended alterations.

Our Approach: Documents are stored and controlled, including change control, via CORE**.** External documents are identified and controlled as appropriate. Records are listed on the **Record Control Plan**, including their retention and disposition processes and retained in ways to prevent unintended changes.

SECTION 2: DO

Providing our customers with products and services that meet their requirements and expectations is why we are in business. This takes planning, reviewing, as well as execution of these processes to ensure that all requirements are identified and met.

In this section of the handbook, we will be describing our methods for conforming to the operational planning, requirements determination and review, design and development, purchasing, product and service provision, post-delivery activities, and what we do when something doesn't go quite as we expected.

8 Operation

8.1 Operational planning and control

Requirement: *Plan, implement and control the processes needed to meet requirements for products and services and to implement the actions determined in 6.1, by:*

- *determining requirements for the product and services;*
- *establishing criteria for the processes and for the acceptance of products and services;*
- *determining the resources needed to achieve conformity to product and service requirements;*
- *implementing control of the processes in accordance with the criteria;*
- *maintaining and retaining documented information to the extent necessary to have confidence that the processes have been carried out as planned and to demonstrate conformity of products and services to requirements.*

The output of this planning is suitable for the organization's operations.

Our Approach: The processes, including outsourced processes that affect our products and services are controlled (8.1). The details and evidence of our processes are maintained within the **QMS Plan** and **Process Plans**. All planned changes are controlled, un-planned changes are reviewed, and actions are taken to mitigate any adverse effects.

8.2 Requirements for products and services

8.2.1 Customer communication

Requirement: *Communication with customers includes:*

- *information relating to products and services;*
- *inquiries, contracts or order handling, including changes;*
- *obtaining customer feedback relating to products and services, including customer complaints;*

- *the handling or controlling of customer property, if applicable;*
- *specific requirements for contingency actions, when relevant.*

Our Approach: Open, and efficient communication with our customers (8.2.1) is very important for communicating information relevant to products and services, contract information, customer complaints, changes, property, requirements and contingency actions. We communicate primarily through email and phone conversations.

8.2.2 Determining the requirements for products and services

Requirement: *When determining the requirements for the products and services to be offered to customers, ensure that:*

- *the requirements for the products and services are defined, including; applicable statutory and regulatory requirements, and those considered necessary;*
- *the organization has the ability to meet the claims for the products and services offered.*

Our Approach: We determine the requirements (8.2.2) for our current or new products and services to ensure that all applicable customer, organizational, regulatory and statutory requirements are identified. This ensures that we can meet our claims, our customers' needs, and any other requirements.

8.2.3 Review of the requirements for products and services

Requirement: *Ensure that the ability to meet the requirements for products and services to be offered to customers is present. Conduct a review before committing to supply products and services to a customer, to include:*

- *customer requirements, including requirements for delivery and post-delivery activities;*
- *requirements not stated by the customer, but necessary for the customers' specified or intended use, when known;*
- *requirements specified by us;*
- *statutory and regulatory requirements applicable to the products and services;*
- *contract or order requirements differing from those previously expressed.*

Ensure that contract or order requirements differing from those previously defined are resolved.

Our Approach: After the requirements are determined, the **Vice President of Engineering** reviews all requirements (8.2.3) to ensure that we have the ability to meet the product and service requirements prior to offering the product or service. If our customer does not provide us with any requirements, we will confirm requirements prior to acceptance. Requirements generated from the product or service, the organization, statutory, regulatory and requirements that are differing from previous ones are reviewed. The results of the review are retained in customer specific folders.

8.2.4 Changes to requirements for products and services

Requirement: *Ensure relevant documented information is amended and that relevant persons are aware of changes.*

Our Approach: When the requirements for products and services are changed (8.2.4), the **Engineering staff and Operations Manager** ensures that relevant documented information is amended and that appropriate personnel are made aware of the changed requirements.

8.3 Design and development of products and services

8.3.1 General

Requirement: *Establish, implement and maintain a design and development process that is appropriate to ensure the subsequent provision of our products and services.*

Our Approach: We maintain a design and development process that is appropriate to the products and services that we offer.

Our design and development process is performed in controlled stages for:

- inputs;
- controls;
- outputs; and,
- changes.

8.3.2 Design and development planning

Requirement: *Determine the stages and controls for design and development, considering:*

- *the nature, duration and complexity of the activities;*
- *the required process stages, including applicable reviews;*
- *the required verification and validation activities;*
- *the responsibilities and authorities involved;*
- *the internal and external resource needs;*
- *the need to control interfaces between individuals and parties involved;*
- *the need for involvement of customer and user;*
- *the requirements for subsequent provision of products and services;*
- *the level of control expected for the processes by customers and other relevant interested parties;*
- *the necessary documented information to confirm that design and development requirements have been met.*

Our Approach: We have determined the stages and controls for our design and development process (8.3.2). The **Design Plan and Schedule** controls all stages of the design and development process. The documented information is retained in the **Design Plan and Schedule.**

8.3.3 Design and development inputs

Requirement: *Determine the requirements essential for the specific types of products and services being designed and developed, considering:*

- *functional and performance requirements;*
- *information derived from previous similar activities;*
- *applicable statutory and regulatory requirements;*
- *standards or codes of practice that we have committed to implement;*
- *the potential consequences of failure due to the nature of the products and services;*

Inputs are adequate for design and development purposes, complete, and unambiguous. Conflicts among inputs are resolved.

Documented information on design and development inputs is retained.

Our Approach: We determine our requirements taking into consideration all input elements. Our design and development inputs (8.3.3) are included within the **Design Plan and Schedule**.

8.3.4 Design and development controls

Requirement: *Apply design and development controls that ensure:*

- *the results to be achieved are defined;*
- *reviews are conducted to evaluate the ability of the results of design and development to meet requirements;*
- *verification activities are conducted to ensure that the outputs meet the input requirements;*
- *validation activities are conducted to ensure the products and services meet the requirements for the application or intended use;*
- *any actions are taken on problems determined during the reviews, or verification and validation activities;*
- *documented information of these activities is retained.*

Our Approach: We apply controls (8.3.4) to our design and development process to ensure that we:

- achieve defined results;

- review and evaluate our Design and Development requirements;
- ensure that outputs meet input requirements;
- meet requirements for application or intended use;
- resolve problems that may arise during design and development;
- retain documented information.

Our design and development controls include the **Design Plan and Schedule**.

8.3.5 Design and development outputs

Requirement: *Ensure that design and development outputs:*

- *meet input requirements;*
- *are adequate for the processes for the provision of products and services;*
- *include or reference monitoring and measuring requirements, as appropriate, and acceptance criteria;*
- *specify the characteristics of the products and services that are essential for their intended purpose and their safe and proper provision.*

Retain documented information resulting from the design and development outputs.

Our Approach: Our design and development outputs (8.3.5) are adequate, and include essential information needed to ensure that all input requirements are met. The design and development outputs are retained on the **Design Plan and Schedule**.

8.3.6 Design and development changes

Requirement: *Review, control and identify changes made during, or subsequent to, the design and development of the products and services, to ensure that there is no adverse impact on conformity to requirements.*

Retain documented information on:

- *design and development changes;*
- *the results of reviews;*
- *the authorization of the changes;*
- *the actions taken to prevent adverse impacts.*

Our Approach: Any changes (8.3.6) that are made during the design and development stages of the products and services are controlled and identified. We retain documented information of the changes, reviews, authorizations and adverse impact actions through the **Design Plan and Schedule**.

8.4 Control of externally provided processes, products and services (Purchasing)

8.4.1 General

Requirement: *Ensure that externally provided processes, products, and services conform to requirements.*

Apply controls to externally provided processes, products and services when:

- *products and services are provided for incorporation into the organization's products and services;*
- *products and services are provided directly to the customer(s) on behalf of the organization;*
- *a process, or part of a process, is provided as a result of our decision.*

Establish and apply criteria for the evaluation, selection, monitoring of performance and re-evaluation of external providers based on their ability to provide processes or products and services in accordance with specified requirements.

Retain documented information of the results of the evaluations, monitoring of the performance and re-evaluations.

Our Approach: We ensure that all of our suppliers of processes, products and services (8.4) conform to all applicable requirements. We apply sufficient controls to any provider of products or services that:

- are directly incorporated into our products or services;
- are provided directly to the customer on our behalf; or,
- provide a process, or part of a process requested by us.

Our criteria for selection, evaluation, performance and re-evaluation practices, is described in the table below:

Criteria	Selection	Evaluation/Re-evaluation
Capability and Experience	X	X
Ability to Meet Technical Specifications	X	X
Price and availability	X	X
On time delivery		X
Customer Specified Supplier	X	X
Product/ Service Quality	X	X
Mgt. System/Certification	X	X
Project Completion		X

The Operations Manager is responsible for controlling the purchasing process and for maintaining appropriate records. Approved suppliers are listed in our Accounting software. External providers are evaluated using the **Supplier Evaluation Form** and reviewed during our Management Reviews.

As of the initial release of this document, all current suppliers in good standing are considered to be approved.

8.4.2 Type and extent of control

Requirement: *Ensure that externally provided processes, products and services do not adversely affect the ability to consistently deliver conforming products and services to customers by:*

- *ensuring that externally provided processes remain within the control of the QMS;*
- *defining both the controls that are intended to be applied to an external provider and those intended to be applied to the resulting output;*
- *taking into consideration the potential impact of the externally provided processes, products and services on the ability to consistently meet customer and applicable statutory and regulatory requirements; and the effectiveness of the controls by the external provider;*
- *determining the verification, or other activities, necessary to ensure that the externally provided processes, products and services meet requirements.*

Our Approach: The controls (8.4.2) that we apply to our external providers are decided on an individual basis. We ensure all suppliers remain in control in our quality management system and apply other controls as necessary by product, service or situation.

8.4.3 Information for external providers

Requirement: *Ensure adequate requirements prior to communicating to the external provider, and communicate the requirements for:*

- *the processes, products and services to be provided;*
- *approval or release of products and services, methods, processes or equipment;*
- *competence of personnel, including necessary qualification;*
- *their interactions with the QMS;*
- *the control and monitoring of the external provider's performance to be applied;*
- *verification or validation activities that the organization, or customers, intend to perform at the external provider's premises.*

Ensure the adequacy of specified requirements prior to communicating to the external provider.

Our Approach: Prior to communicating with suppliers, we ensure that all applicable requirements are clearly identified. These may include requirements relating to products, services, supplier processes, certifications or personnel, and any verification or validation that the supplier provides at their premises.

The purchasing information (8.4.3) is communicated to suppliers via purchase orders and/or contracts.

8.5 Production and service provision

8.5.1 Control of production and service provision

Requirement: *Implement controlled conditions, including, as applicable:*

- *the availability of documented information that defines the characteristics of the products and services, and the results to be achieved;*
- *monitoring and measurement activities at appropriate stages to verify that criteria for control of processes and process outputs, and acceptance criteria for products and services, have been met;*
- *the use, and control of suitable infrastructure and process environment;*
- *the availability and use of suitable monitoring and measuring resources;*
- *the competence and, where applicable, required qualification of people;*
- *the validation, and periodic revalidation, of the ability to achieve planned results of any process for production and service provision where the resulting output cannot be verified by subsequent monitoring or measurement;*
- *the implementation of actions to prevent human error;*
- *the implementation of products and services release, delivery and post-delivery activities.*

Our Approach: We control all phases of our product or service realization (8.5.1). These controls may include; documented characteristics, monitoring and measurement, validations or reviews of products and/or processes, and release and post-delivery activities.

The Operations Manager is responsible for controlling all phases of product and service provision and for maintaining appropriate records.

8.5.2 Identification and traceability

Requirement: *Use suitable means to identify outputs when it is necessary to ensure the conformity of products and services. Identify the status of outputs with respect to monitoring and measurement requirements throughout production and service provision. Control the unique identification of the outputs when traceability is a requirement and retain the documented information necessary to enable traceability.*

Our Approach: Products or services are identified (8.5.2) by means of procedures defined in the Identification and Traceability procedure Where traceability (8.5.2) is a requirement, we use methods suitable to identify outputs to ensure conformity of our products or services. The method(s) used for traceability is determined by the **Identification and Traceability Procedure** and is accomplished through the use of the Traceability batch number log.

8.5.3 Property belonging to customers or external providers

Requirement: *Exercise care with property belonging to the customer or external providers while it is under organizational control or being used. Also, identify, verify, protect and safeguard the customer's or external provider's property provided for use or incorporation into our products and services.*

Property of the customer or external provider which is lost, damaged or found to be unsuitable, is reported to the customer or external provider and documented information of what occurred is retained.

Our Approach: There may be times that we use property belonging to customers or external providers (8.5.3). When this occurs, we identify, verify and protect the provider's property.

The **Operations Manager** is responsible for controlling and recording customer property.

In the rare occurrence of customer or provider's property becoming lost, damaged or unusable, the Operations Manager will contact the provider. The record of communication is retained in customer files.

8.5.4 Preservation

Requirement: *Ensure preservation of process outputs during production and service provision, to the extent necessary to maintain conformity to requirements.*

Our Approach: We use methods necessary to ensure that our product or service maintains conformance to the requirements.

8.5.5 Post-delivery activities

Requirement: *Meet requirements for post-delivery activities associated with products and services, considering:*

- *customer requirements;*
- *the nature, use and intended lifetime of the products and services;*
- *customer feedback;*
- *statutory and regulatory requirements;*
- *the potential, undesired consequences associated with its products and services.*

Our Approach: Post-delivery activities (8.5.5) include customer feedback and a delivery quality performance log and are developed considering applicable requirements, product/service use, customer feedback and potential risks.

8.5.6 Control of changes

Requirement: *Review and control changes for production or service to the extent necessary to ensure conformity with specified requirements.*

Retain documented information describing the results of the review of changes, the personnel authorizing the change, and any necessary actions arising from the review.

Our Approach: Any changes (8.5.6) that occur during our product or service provision are controlled by change orders, and recorded on the change order log.

8.6 Release of products and services

Requirement: *Implement planned arrangements at appropriate stages to verify that product/service requirements have been met.*

Release of products/services to the customer does not proceed until the planned arrangements have been satisfactorily completed, unless otherwise approved by a relevant authority and, as applicable, by the customer. Retain documented information on the release of products/services includes:

- *evidence of conformity with acceptance criteria;*
- *traceability to the person(s) authorizing the release.*

Our Approach: The release of our product/service (8.6) is indicated by means of the Design and Development Process and recorded in the Design Review Summary and does not occur until all planned arrangements have been completed and is only released by authorized persons.

8.7 Control of nonconforming process outputs (products and services)

Requirement: *Ensure process outputs that do not conform to requirements are identified and controlled to prevent unintended use or delivery.*

Take action based on the nature of the nonconformity and its effect on the conformity of products/services. This applies also to nonconforming products/services detected after delivery of the products or during or after the provision of the service.

As applicable, deal with nonconforming outputs in one or more of the following ways:

- *correction;*
- *segregation, containment, return or suspension of products and services;*
- *informing the customer;*
- *obtaining authorization for acceptance under concession.*

Where nonconforming outputs are corrected, conformity to the requirements is verified.

Retain documented information that:

- *describes the nonconformity;*
- *describes the actions taken;*

- *describes any concessions obtained;*
- *identifies the authority deciding the action in respect of the nonconformity.*

Our Approach: Any output that does not conform (8.7) to requirements is identified and controlled to prevent unintended use or delivery.

We take appropriate action to deal with the nonconformity. Resolutions are described on the **Nonconformance Form**, **Customer Complaint Form** and/or **Corrective Action Form**.

SECTION 3: CHECK

We make great efforts to be data-driven decision makers. This can be accomplished only by ensuring that we maintain accurate data and that the data is properly interpreted.

We take the time to analyze data from various areas that supplies us with data on:

- customer satisfaction;
- process effectiveness;
- product/service conformity;
- effectiveness of our QMS;
- external providers;
- our planning efforts;
- the associated risks and opportunities.

Our thorough "checking" process allows us to have confidence in our quality management system and identify improvement areas.

9 Performance Evaluation

9.1 Monitoring, measurement, analysis and evaluation

9.1.1 General

Requirement: *Determine:*

- *what needs to be monitored and measured;*
- *the methods for monitoring, measurement, analysis and evaluation to ensure valid results;*
- *when the monitoring and measuring will be performed;*
- *when the results from monitoring and measurement will be analyzed and evaluated.*

Evaluate the performance and effectiveness of the quality management system through the Management Review process and retain documented information as evidence of the results.

Our Approach: Our method of monitoring, measurement, analysis and evaluation is maintained within our **Measurement Plan**. The review of this plan is retained in our Management Review minutes.

9.1.2 Customer satisfaction

Requirement: *Monitor customer perceptions of the degree to which their needs and expectations have been fulfilled.*

Our Approach: We obtain and, monitor customer perception by means of a Customer Satisfaction Survey. The customer satisfaction data is discussed during Management Reviews.

9.1.3 Analysis and evaluation

Requirement: *Analyze and evaluate appropriate data and information arising from monitoring, measurement and other sources to evaluate:*

- *conformity of products and services;*
- *the degree of customer satisfaction;*
- *the performance and effectiveness of the QMS;*
- *planning implementation;*
- *the effectiveness of actions taken to address risks and opportunities;*
- *the performance of external provider(s);*
- *needs or opportunities for improvements to the QMS.*

Our Approach: Our sources and evaluations (9.1.3) are described within our **Measurement Plan** and retained within our **Management Review Meeting Minutes**.

9.2 Internal audit

Requirement: *Conduct internal audits at planned intervals to provide information on whether the QMS conforms to requirements, is implemented and maintained.*

The organization shall:

- *plan, establish, implement and maintain an audit program(s) including the frequency, methods, responsibilities, planning requirements and reporting, which takes into consideration, the importance of the processes concerned, changes affecting the organization, and the results of previous audits;*
- *define the audit criteria and scope for each audit;*
- *select auditors and conduct audits to ensure objectivity and impartiality of the audit process;*
- *ensure that the results of the audits are reported to relevant management;*
- *take appropriate correction and corrective actions without undue delay;*
- *retain documented information as evidence of the implementation of the audit program and audit results.*

Our Approach: Our internal audit program is implemented, maintained and is used to ensure that our QMS is maintained and effective. Our internal audits are planned according to importance on our **Internal Audit Plan and**

Schedule. The scope of an individual audit is defined on the **Process Plan** for that specific key process. Our auditors are objective and impartial and report the results to management. Auditors are qualified based on completion of an auditor training course or previous experience. Records of this training are maintained by the Management Representative**.** Corrective Actions resulting from internal audits are taken without undue delay.

The Management Representative is responsible to oversee the internal auditing system and for retaining appropriate documented information. Internal audit results and status are discussed during Management Review.

9.3 Management review

Requirement: *Top management conducts planned reviews of the QMS to ensure its suitability, adequacy, effectiveness and alignment with the strategic direction considering:*

- *the status of actions from previous management reviews;*
- *changes in external and internal issues that are relevant to the QMS;*
- *information on the performance and effectiveness of the quality management system, including trends in:*
    - *customer satisfaction and feedback from relevant interested parties;*
    - *the extent to which quality objectives have been met;*
    - *process performance and conformity of products and services;*
    - *nonconformities and corrective actions;*
    - *monitoring and measurement results;*
    - *audit results;*
    - *the performance of external providers;*
    - *the adequacy of resources;*
    - *the effectiveness of actions taken to address risks and opportunities;*
    - *opportunities for improvement.*

The outputs of management review are to include decisions and actions related to:

- *opportunities for improvement;*
- *any need for changes to the quality management system;*
- *resource needs.*

Retain documented information as evidence of the results of management reviews.

Our Approach: Our management reviews are planned and occur at planned intervals using the **Management Review Plan.** These reviews are attended by:

- **David Mosier - President**
- **Rodney Bame – VP Engineering**
- **Josh Jenkins – VP Sales**
- **Brandey Sweat – Operations Manager**

- **Chris Busch – Quality Manager**

The Management Reviews are planned using a schedule and meeting agenda consisting of all required inputs. The meetings are retained on the **Management Review Meeting Minutes**.

Outputs from our Management Reviews include the actions and decisions relating to any opportunities for improvement, needed changes to the QMS and resource needs. Outputs are also retained on the **Management Review Meeting Minutes**.

SECTION 4: ACT

This final step within our Plan, Do, Check and Act quality management system serves two purposes. First, it is the step which is used to make the decision of taking or not taking action based on the analysis and evaluations that occur during the "check" step. Whether we decide to take action or not, the decision will always be metric-driven, and risk-based.

The second purpose of the "Act" step is that it serves as the pivoting step that guides our QMS back to the Plan phase to begin the PDCA cycle and support continual improvement.

This last section of our manual covers our approach to improvements and corrective actions.

10 Improvement

10.1 General

Requirement: *Determine and select opportunities for improvement and implement actions to meet customer requirements and enhance customer satisfaction, including (as appropriate):*

- *improving products and services to meet requirements as well as to address future needs and expectations;*
- *correcting, preventing or reducing undesired effects;*
- *improving the performance and effectiveness of the quality management system.*

Our Approach: We select opportunities to:

- improve our products and services;
- correct, prevent or reduce undesired effects;
- improve our QMS.

We retain the documented information regarding improvements on our **Improvement Plan**, **Corrective Action Forms** and **Management Review Meeting Minutes**.

10.2 Nonconformity and corrective action

Requirement: *When a nonconformity occurs, including those arising from complaints, we:*

- *react to the nonconformity, and as applicable:*
  - *take action to control and correct it;*
  - *deal with the consequences;*
- *evaluate the need for action to eliminate the cause(s) of the nonconformity, in order that it does not recur or occur elsewhere:*

- *review and analyze the nonconformity;*
- *determine the causes of the nonconformity;*
- *determine if similar nonconformities exist, or could potentially occur;*

- *implement any action needed;*
- *review the effectiveness of any corrective action taken;*
- *update risks and opportunities determined during planning, if necessary;*
- *make changes to the quality management system, if necessary.*

Corrective actions are appropriate to the effects of the nonconformities encountered.

Retain documented information as evidence of:

- *the nature of the nonconformities and any subsequent actions taken;*
- *the results of any corrective action.*

Our Approach: Nonconformities are taken seriously and are reacted to as applicable. We take any actions necessary to ensure that the nonconformity does not recur or occur elsewhere. Nonconformities are documented on our **Nonconformance Form** and/or **Corrective Action Form** and discussed during Management Review.

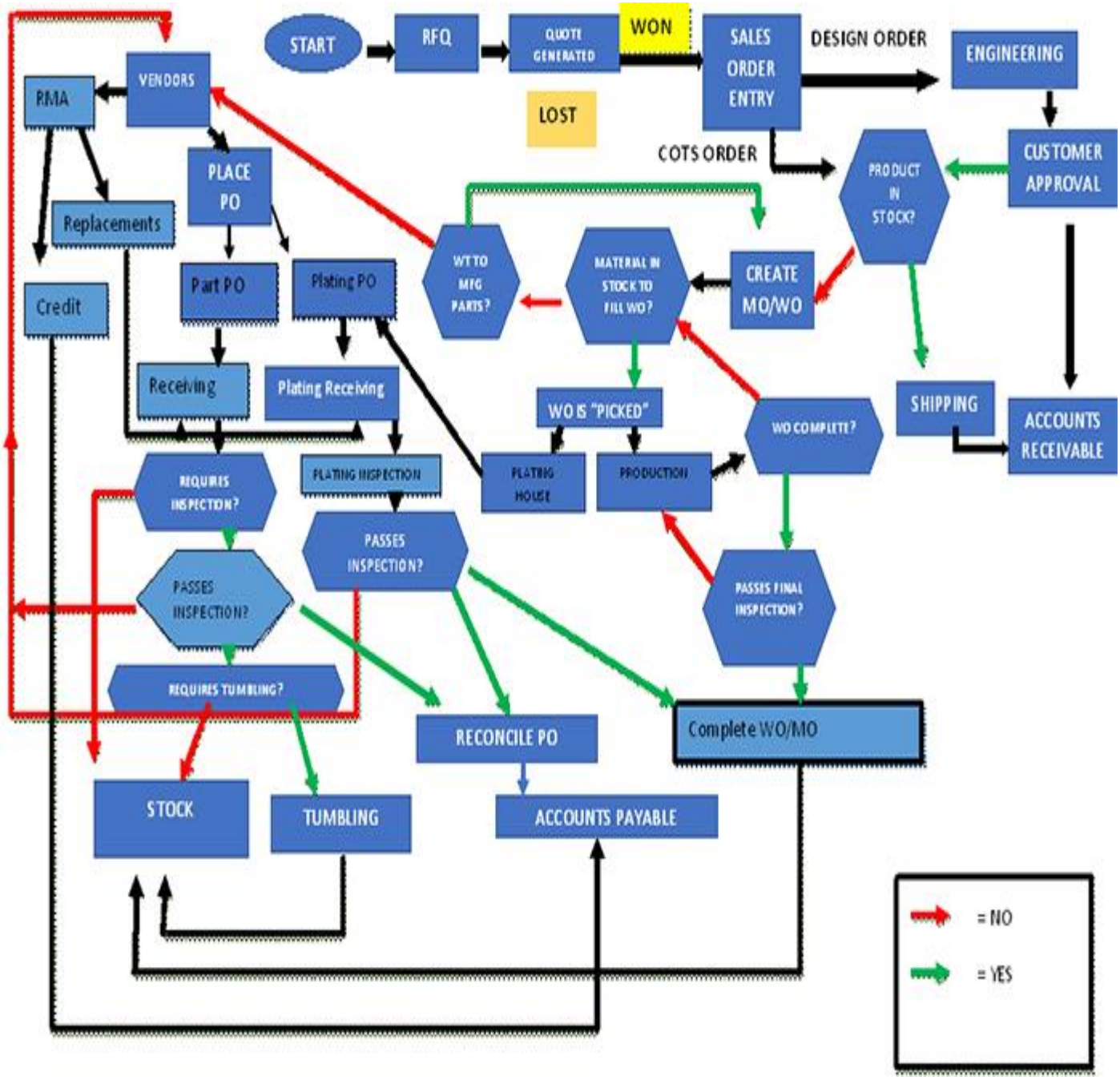
10.3 Continual improvement

Requirement: *Continually improve the suitability, adequacy, and effectiveness of the QMS*.

Our Approach: We consider the results of analysis and evaluation, and the outputs from Management Review, to confirm if there are needs or opportunities to be addressed as part of continual improvement.

Appendix A

Process Flow Diagram



Appendix B

Documented Information List

Doc. #	Description	Responsibility	Revise date
QM-001	QMS Manual – Document Information	President	11/17/2021
P-400	Organizational context	President	11/17/2021
P-500	Leadership	President	11/17/2021
P-600	Planning for the QMS	Quality Manager	11/17/2021
P-710	Resource management	Operations manager	11/17/2021
P-715	Control of monitoring and measuring equipment	Quality manager	11/17/2021
P-720	Competence and awareness	Operations manager	11/17/2021
P-740	Communication	Quality Manager	11/17/2021
P-750	Control of documented information	Engineering	11/17/2021
WI-750-001	Document numbering system	Engineering	11/17/2021
P-810	Operational planning and control	Operations Manager	11/17/2021
P-820	Customer related process	Sales, Marketing manager	11/17/2021
P-830	Design and development	Engineering	11/17/2021
P-840	Control of external providers	Operations Manager	11/17/2021
SOP 8.5.5.x	SOPs	Operations Manager	11/17/2021
P-852	Identification and traceability	Operations Manager	11/17/2021
P-870	Control of nonconforming outputs.	Quality manager	11/17/2021
P-910	Monitoring, measurement, analysis and evaluation	Quality Manager	11/17/2021
P-912	Customer Satisfaction	Sales, marketing manager	11/17/2021
P-920	Internal audits	President	11/17/2021
P-930	Management review	President	11/17/2021
P-1010	Improvement	Management Group	11/17/2021
P-1020	Nonconformity and corrective action	Operations Manager	11/17/2021
F-440-001	QMS-Process identification worksheet	Quality Manager	11/17/2021
F-440-002	Organization context worksheet	Quality Manager	11/17/2021
FD-440-001	Process interaction	Quality Manager	11/17/2021
A-520-001	Quality policy	Quality Manager	11/17/2021
A-530-001	Organization chart	Quality Manager	11/17/2021
F-600-001	Strategic Direction & Quality Plan Template	Quality Manager	11/17/2021

F-600-002	SWOT Form Template	Quality Manager	11/17/2021
A-610-001	SWOT-20171010 Risk Analysis Variable Force	Quality Manager	11/17/2021
A-610-002	SWOT-20170425 Risk Analysis_30D	Quality Manager	11/17/2021
A-610-003	SWOT-20170711 Risk Analysis Moving	Quality Manager	11/17/2021
A-610-004	SWOT-20190524_End Mill	Quality Manager	11/17/2021
A-620-001	Strategic Direction & Quality Plan 2019	Quality Manager	11/17/2021
A-620-002	Strategic Direction & Quality Plan 2020	Quality Manager	11/17/2021
FD-620-001	PDCA Objectives planning	Quality Manager	11/17/2021
F-620-001	Quality objectives planning record	Quality Manager	11/17/2021
FD-620-003	Material Rejection Flow Chart	Operations / Engineering	11/17/2021
F-620-005	FEMA Template	Operations / Engineering	11/17/2021
F-620-007	Non Conforming Analysis Form	Operations / Engineering	11/17/2021
A-620-007	Non Conforming Log	Operations / Engineering	11/17/2021
A-715-001	Equipment calibration list	Quality Manager	11/17/2021
A-715-002	Caliper Calibration Log	Quality Manager	11/17/2021
A-715-003	Crimper Wear	Operations / Engineering	11/17/2021
F-720-001	Training action plan	Operations / Engineering	11/17/2021
F-720-002	Job Description	Operations / Engineering	11/17/2021
A-720-001	Training Records	Operations / Engineering	11/17/2021
A-720-002	Master Training Record	Operations / Engineering	11/17/2021
F-740-001	Customer Satisfaction report	Sales	11/17/2021
F-750-001	List of documented information	Operations / Engineering	11/17/2021
F-750-006	Document Revision Checklist	Operations / Engineering	11/17/2021
F-750-007	Software inventory spreadsheet	Quality Manager	11/17/2021
F-750-004	Quality records table	Quality Manager	11/17/2021
WI-750-001	Document numbering system	Operations / Engineering	11/17/2021
WI-750-002	Sales Order Entry Procedure	Sales	11/17/2021
FD-810-001	Process flow diagram	Operations / Engineering	11/17/2021
F-810-002	FEMA Form	Operations / Engineering	11/17/2021
A-810-001	FEMA Process Solid Wedge	Operations / Engineering	11/17/2021
A810-002	FEMA Design Ejector	Operations / Eng	11/17/2021

A-810-003	FEMA Process Ejector	Operations / Engineering	11/17/2021
A-820-001	Customer Complaint Log	Sales	11/17/2021
A-820-002	Customer Satisfaction Log	Sales	11/17/2021
A-820-003	Quoting Procedure	Sales	11/17/2021
F-820-001	Client assessment report	Sales	11/17/2021
F-820-002	Production Order	Operations	11/17/2021
F-830-001	Design & Input Review	Engineering	11/17/2021
F-830-002	Design review record	Engineering	11/17/2021
F-830-003	Design & Development Cover Sheet	Engineering	11/17/2021
F-830-004	Design Change Form	Engineering	11/17/2021
A-830-001	Strategic Direction & Quality Plan 2019	Management Team	11/17/2021
F-840-001	Supplier PO Ts&Cs	Operations	11/17/2021
F-840-002	New Supplier Approval Form	Operations	11/17/2021
F-840-003	Material Rejection Form	Operations	11/17/2021
Fishbowl	Acceptable Source	Operations	11/17/2021
FD-840-001	Material Rejection	Operations	11/17/2021
F-840-005	Inventory Adjustment Form	Operations	11/17/2021
A-850-001	SOP & Work Instruction Document List	Operations	11/17/2021
F-851-001	Job Order Traveler	Operations	11/17/2021
F-851-002	Stock Order Traveler	Operations	11/17/2021
F-851-003	Work Instruction Template	Operations	11/17/2021
A-851-001	Job Order Traveler Example	Operations	11/17/2021
A-851-002	Job Order Traveler Example	Operations	11/17/2021
A-851-003	Backstrap Image	Operations	11/17/2021
A-851-004	Rubber Band Image	Operations	11/17/2021
A-851-005	Setup Image	Operations	11/17/2021
A-851-006	Specification List & Signoff	Operations	11/17/2021
A-851-007	Loctite Location Image	Operations	11/17/2021
A-851-008	Plating Legend	Operations	11/17/2021
WI 8.5.1.1	Work Inst - Drive Assy.	Operations	11/17/2021
WI 8.5.1.2	Work Inst - Crimping LH #6.Drive Scr	Operations	11/17/2021
WI 8.5.1.4	Work Inst - Crimping LH #8.Drive Scr	Operations	11/17/2021
WI 8.5.1.6	Work Inst - Body Assy, Gen 4	Operations	11/17/2021
WI 8.5.1.7	Work Inst - Body Assy, Gen 5	Operations	11/17/2021
WI 8.5.1.8	Work Inst - Helicoil Insertion, tanged	Operations	11/17/2021
WI 8.5.1.12	Work Inst - Thermal Interface	Operations	11/17/2021

WI 8.5.1.13	Work Inst - Thermal Machine Cutting	Operations	11/17/2021
WI 8.5.1.14	Work Inst - Crimping LH #10 Screw	Operations	11/17/2021
WI 8.5.1.16	Work Inst - Body Assy Gen 1B	Operations	11/17/2021
WI 8.5.1.18	Work Inst - Captive Screw Insertion HKE	Operations	11/17/2021
WI 8.5.1.20	Work Inst - Inspecting Keyance Parts	Operations	11/17/2021
WI 8.5.1.21	Work Inst - Crimping Instructions	Operations	11/17/2021
WI 8.5.1.2	Ejector Mounting	Operations	11/17/2021
SOP 8.5.1.1	SOP - Solid Wedge Gen 4	Operations	11/17/2021
SOP 8.5.1.2	SOP - Solid Wedge Gen 5	Operations	11/17/2021
SOP 8.5.1.3	Heatframes	Operations	11/17/2021
SOP 8.5.1.4	Ejector Kits	Operations	11/17/2021
SOP 8.5.1.5	Final Inspection	Operations	11/17/2021
SOP 8.5.1.7	FOD Prevention Procedure	Operations	11/17/2021
F-852-001	Identification tag	Operations	11/17/2021
F-852-002	Traceability serial number log	Operations	11/17/2021
F-852-003	Traceability label	Operations	11/17/2021
F-854-001	Storage inspection report	Operations	11/17/2021
F-870-001	Non conforming	Operations	11/17/2021
F-870-002	Non Conforming Output Analysis Form	Operations	11/17/2021
F-870-003	Discrepant Material Report Form	Operations	11/17/2021
F-870-004	Scrap Log	Operations	11/17/2021
F-870-005	Final Inspection. Rework Log	Operations	11/17/2021
F-870-006	Final Inspection Pass/ Fail Log	Operations	11/17/2021
A-870-001	FOD Awareness Sign	Operations	11/17/2021
F-910-001	Production – Monitoring, measuring, and analysis table	Operations	11/17/2021
F-910-002	QMS – Monitoring, measuring, and analysis table	Operations	11/17/2021
F-910-004	Inspection report	Operations	11/17/2021
F-912-001	Customer Satisfaction Form	Sales	11/17/2021
A-912-001	Delivery Quality Performance	Operations	11/17/2021
F-920-001	Applicable procedure by work area	Operations	11/17/2021
F-820-002	Internal audit checklist	Quality Manager	11/17/2021
F-920-003	Audit plan	Quality Manager	11/17/2021
F-920-004	Audit report	Quality Manager	11/17/2021
P-930-001	Management review agenda	Quality Manager	11/17/2021
F-930-002	Management review output report	Quality Manager	11/17/2021

A-930-001	2018 12-6 6 Month Management Review	Quality Manager	11/17/2021
A-930-002	2019 12-11 6 Month Mgt Review	Quality Manager	11/17/2021
F-1010-001	Data analysis worksheet	Operations / Engineering	11/17/2021
A-1010-001	Quality Objectives RMA chart	Operations / Engineering	11/17/2021
F-1020-001	Corrective Action Request - CAR	Operations / Engineering	11/17/2021
A-1020-001	Corrective Action Log 2017	Operations / Engineering	11/17/2021
A-1020-002	Corrective Action Log 2018	Operations / Engineering	11/17/2021
A-1020-003	Corrective Action Log 2019	Operations / Engineering	11/17/2021
A-1020-004	Corrective Action Log 2020	Operations / Engineering	11/17/2021
A-1020-005	2018 CARS	Operations / Engineering	11/17/2021
A-1020-006	2019 CARS	Operations / Engineering	11/17/2021
A-1020-007	2020 CARS	Operations / Engineering	11/17/2021
A-1020-008	2021 CARS	Operations / Engineering	11/17/2021