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REDACTED

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ASA-100 QUALITY MANUAL

Origination Date: (your origination date)

Document Identifier:	QMS-00 Quality Manual
Date:	Latest Revision Date
Document Revision:	Orig

Abstract:

This quality manual describes (your Company's) quality management system policies and procedures according to ASA-100 Rev: 5.

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Issue	Date	Comment	Author
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DOCUMENT CHANGE RECORD

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NOTE: Company policies herein are expressed from the perspective of "As-a-Matter-of-Fact". To apply this perspective, mentally add the phrase to the beginning of each paragraph herein. Delete this note prior to release of quality manual.

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1.0 Quality System and Quality Manual

(Your Company's) quality management system (QMS) quality manual summarizes [REDACTED]

The Company ensures [REDACTED] ensures requirements for products and services are defined according to the *QMS-07 Proposal Development and Contract Review Procedure*.

2.0 Self-Audit/Evaluation

2.1 Internal audit

The Company conducts internal audits [REDACTED] according to the following procedures: *QMS-12 Internal Auditing*, *QMS-04 Management Process*, *QMS-14 Control of Nonconformances* and *QMS-13 Corrective Action*.

2.2 Audit requirements

The Company assigns Responsible Authorities [REDACTED] according to the *QMS-12 Internal Auditing Procedure*.

3.0 Facilities

The Company determines and provides [REDACTED] QC stamps or registered [REDACTED] may be used to [REDACTED].

The Company maintains [REDACTED].

The Company determines, provides and maintains [REDACTED] according to the *QMS-04 Management Process Procedure*.

4.0 Training and Authorized Personnel

4.1 People

The Company determines and provides [REDACTED] according to the *QMS-04 Management Process Procedure*, *QMS-05 Responsibilities and Authorities* and *QMS-06 Training Procedure*. See Appendix A for Responsible Authority Chart.

4.2 Competence

The Company ensures [REDACTED] which includes [REDACTED] Action is taken [REDACTED].

The Company evaluates [REDACTED] according to the *QMS-04 Management Process Procedure*, *QMS-06 Training Procedure* and *QMS-03*

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Records Control Procedure. The Company maintains [REDACTED]
[REDACTED]

5.0 Procurement

5.1 General

The Company ensures that drop shipments and externally provided products are [REDACTED] according to the *QMS-08 Purchasing Procedure* and *QMS-09 Receiving Procedure*. The Company determines and applies [REDACTED] according to requirements and the *QMS-08 Purchasing Procedure*. The Company maintains a list of approved Suppliers and [REDACTED]

5.2 Type and extent of control

The Company ensures [REDACTED] according to the *QMS-08 Purchasing Procedure* and *QMS-09 Receiving Procedure*.

5.3 Information for external providers

The Company ensures [REDACTED] according to the *QMS-08 Purchasing Procedure*.

6.0 Receiving Inspection

Incoming supplies are [REDACTED] which is defined in the *QMS-09 Receiving Procedure*.

7.0 Measuring and Test Equipment

Measuring equipment is [REDACTED] according to the *QMS-15 Calibration Procedure*.

8.0 Material Control

The Company uses [REDACTED] The inspection/approval status of supplies is [REDACTED] according to the *QMS-10 Production Procedure*. The Company controls [REDACTED] split-lot/batch [REDACTED] for [REDACTED]

9.0 Shelf Life Control

The Company reviews [REDACTED] according to *QMS-08-1 Purchase Order Review Procedure* then identifies and controls [REDACTED] according to *QMS-09 Receiving Procedure*.

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10.0 Certification and Release of Materials

The Company prepares [REDACTED] according to the *QMS-11-Shipping Procedure*, which includes [REDACTED]. Traceability [REDACTED] is controlled according to the *QMS-10 Production Procedure*.

11.0 Shipping

Products are released for drop shipment [REDACTED]. The Company retains [REDACTED]. Hazardous materials and deliverable supplies [REDACTED] according to the *QMS-11 Shipping Procedure* and the *QMS-16 Hazardous Material Handling Procedure*.

12.0 Records

The Company controls records to [REDACTED]. Records are suitable for [REDACTED] according to the *QMS-03 Records Control Procedure*.

13.0 Technical Data Control

Documents required for production of deliverable items are [REDACTED]. The Company controls [REDACTED] according to the *QMS-01 Document Control Procedure* and/or *QMS-03 Records Control Procedure*. Internal and external documents used [REDACTED] controlled according to the *QMS-02 Configuration Management Procedure*, which addresses [REDACTED].

14.0 Corrective Action Process

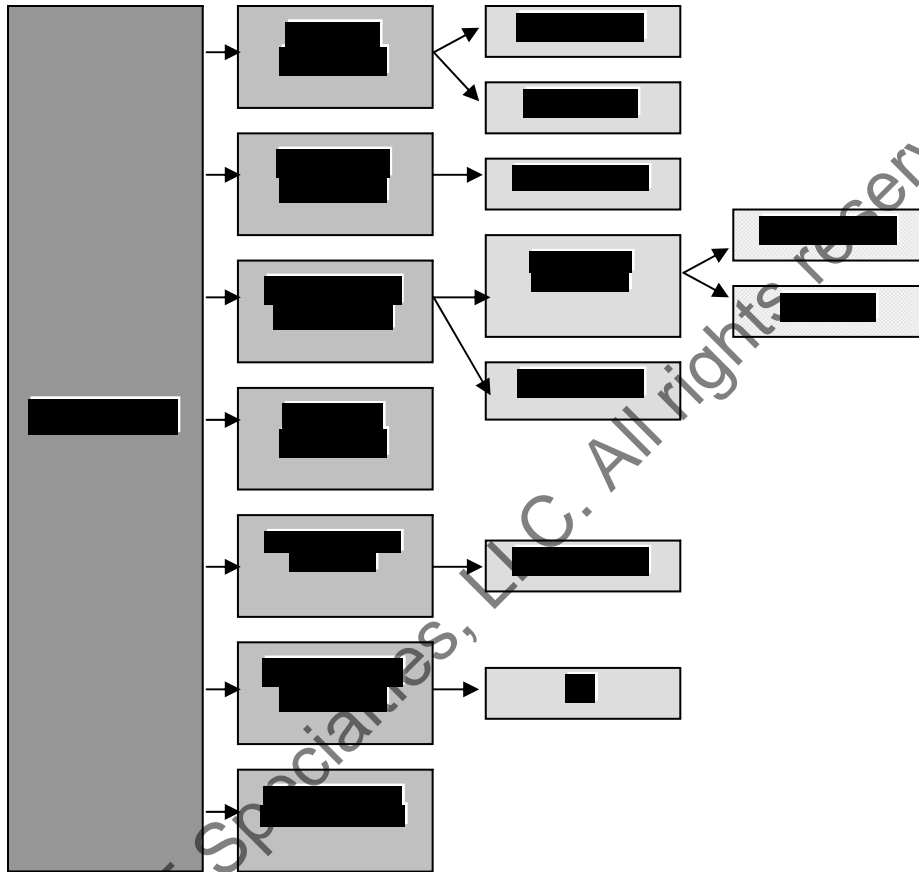
When a nonconformance occurs, including complaints and discovery of nonconformity(s) after delivery to Customer, the Company [REDACTED] according to the *QMS-13 Corrective Action Procedure* and *QMS-14 Nonconformance Control Procedure*. The Company [REDACTED] if [REDACTED]. The Company implements [REDACTED] and implements [REDACTED]. The Company reviews [REDACTED]. The Company ensures [REDACTED].

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14.1 Required records for nonconformities

The Company retains and maintains records [REDACTED] according to the *QMS-03 Records Control Procedure*.

Appendix A: Responsible Authorities



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