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AS 9120 Rev B

Quality Management Systems Documentation

Quality Manual / Documented Information

Document No. QM-9120-B

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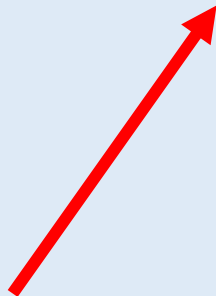
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Blue text throughout the manual highlight areas for customization

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Introduction

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Section B References

a. Normative reference

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Any text may be edited.

Quality Management System Requirements

Blue text provides examples of what you may want of use.

Section C Document Information

Black text describes the QMS.

a. Distribution Control List

b. Revision Status

c. Quality Policy, Quality Objective, Strategic Direction,

d. Organization Chart

e. Company Background - Products and Services

f. Process Flow Diagram

Section D List of Documented Information for the ISO standard clauses 4 through 10

Clause 4 Context of the Organization

Clause 5 Leadership

Clause 6 Planning

Clause 7 Support

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Section A Scope or the Quality Management System Provides general purpose and description of Quality Manual
General

To determine and establish the scope of the QMS, **Your Company** determined the boundaries and applicability of the QMS and considered the external and internal issues, the requirements of relevant interested parties, and the products and services of the company. The scope is available and maintained as documented information stating the products and services covered by the QMS.

Your Company applies all the requirements of AS 9120 Rev B when they are applicable within the determined scope of the QMS.

As developed with procedure P-400 for Organizational context, include the scope of your QMS here:

 **Blue text gives guidance for customization.**

For example, if you are a distributor of landing gear tires, the scope of the Quality Management System includes the major product and service categories associated with the distribution of landing gear tires from the Main Street warehouse location to regional, national, and international aviation, space, and defense customers.

Conformity to the standard may only be claimed if the requirements determined as not being applicable do not affect the organization’s ability or responsibility to ensure the conformity of its products and services and the enhancement of customer satisfaction.

In the event that any requirement is not applicable at **Your Company**, justification for any instance where a requirement cannot be applied is documented.

Your Company has determined that the following requirement(s) is/are not applicable to the operations at this site:

As determined with procedure **P-400**, identify the requirement(s) that do not apply and document the justification here:

Related documents are referenced.

For example, if you are a distributor of aircraft tires, a requirement that does not apply:

Clause 8.3 for design and development does not apply to the company. The product is designed and developed and meets requirements through the designer and provider of landing gear tires.

Section B References

a. Normative reference

- 9100:2016 Quality Management Systems – Requirements for aviation, space, and defense organizations,
- ISO 9000:2015 Quality Management Systems – Fundamentals and vocabulary.
- ISO 9001:2015 Quality Management Systems – Requirements

b. Definitions Applicable definitions are included in documented procedures and instructions at par 3.0 to enhance the understanding of the process.

Control of Monitoring and Measuring Equipment

1.0 Purpose/Scope

- 1.1 To outline the requirements for control of measuring and monitoring equipment at [Your Company](#).
- 1.2 The procedure applies to equipment where monitoring or measuring is used for evidence of conformity of [products and services](#)

2.0 Responsibilities and Authorities

- 2.1 The [Quality assurance manager / Management representative](#) has the prime responsibility and approval authority for this procedure.
- 2.2 In support of the [Quality assurance manager](#) and where monitoring or measuring is used for evidence of conformity of products and services, the [Quality team / AS steering committee](#) is responsible for determining the resources needed to ensure valid and reliable monitoring and measuring results.
- 2.3 The [Quality team / AS steering committee](#) is responsible to designate the [Equipment coordinator](#), and to assign responsibility for calibration and maintenance of the equipment.

3.0 References and Definitions

- 3.1 Reference: This document addresses clause 7.1.5 of the AS 9120 B standard, covering monitoring and measuring resources.
- 3.2 No definitions

4.0 Resources

- 4.1 None, ([unless an electronic equipment calibration tracking system is used](#)).

5.0 Instructions

- 5.1 The [Quality team / AS steering committee](#) determines and provides the resources needed to ensure valid and reliable results when monitoring and measuring is used to verify conformity to requirements.
- 5.1.1 With procedures P-810 for Operational planning and control, P-851 for Control of production and service provision, and P-910 for Monitoring, measurement, analysis and evaluation, consideration is given to monitoring and measuring resources to ensure that they are:
- Suitable for the specific type of monitoring and measuring activities undertaken,
 - Maintained to ensure their continuing fitness for their purpose and documented information maintained as evidence of fitness for purpose.
 - Calibrated or verified in suitable environmental conditions.
- 5.2 The [Quality team / AS steering committee](#) ensures that measuring instruments are calibrated when measurement traceability is considered to be an essential part of providing confidence in valid measurement results, [or is a statutory or regulatory requirement, or is customer or interested party expectations](#).

You can search and replace "your company" with your own company name.

INSERT COMPANY NAME/LOGO HERE

A-840-001

Provider Selection Guidelines

GUIDELINES – Evaluation and Selection of External Providers	Date Approved	Data Form A-840-001
<p>Providers are evaluated and selected by one of the following methods:</p> <p>Review methods listed below at par 1.1 to 1.6 and select one or more that are appropriate for your company.</p> <p>If you have goods or services that vary in its impact on quality you may want to set up categories, the higher the impact the more comprehensive the method. You may need to combine more than one method, for example an audit and samples for inspection and test.</p> <p>1.1 The provider is, at a minimum, registered to ISO 9001:2015.</p> <ul style="list-style-type: none">• Purchasing department staff reviews and maintains a copy of their certificate and quality manual on file.▪ Purchasing / Quality management staff performs quality system development with the objective of provider conformance to ISO 9001:2015 and leading to AS 9120 B. <p>1.2 The provider provides graded or classed material, and provides certificate of analysis with the material or item.</p> <p>1.3 Samples of the materials or items are provided for inspection and test, with satisfactory results.</p> <ul style="list-style-type: none">• The person requesting the purchase documents the sample size required and the inspection and test to be performed on the purchasing documents.• Completed inspection and test records show the criteria for acceptance and the actual results. If they are acceptable, the requisitioner sends them to purchasing to be kept in the provider's file. <p>1.4 An audit of the provider confirms that required elements of a quality system are in place and results documented in the provider assessment report F-840-001.</p> <ul style="list-style-type: none">• The Quality manager assigns an individual or team to perform the audit.• The Quality manager reviews the completed audit checklist, and determines if the supplier meets requirements.• If the provider meets requirements, the purchasing manager indicates acceptance on the provider assessment report and keeps the audit checklist in the provider's file.• The approved provider is added to the List of acceptable sources, form F-840-002. <p>1.5 The provider is specified by the customer contract. The use of customer designated providers does not relieve Your Company of the responsibility to ensure quality.</p> <p>1.6 The Purchasing department places a trial order.</p> <ul style="list-style-type: none">• Purchasing department orders the material or item, and the requisitioner uses the material, and measures the results.• If the results are not acceptable, the product that it was used for is controlled according to the control of nonconforming product procedure, P-870.• If the results are acceptable, they are documented and kept in the provider's file.		

Blue text throughout the manual highlight areas for customization

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INSERT YOUR COMPANY LOGO/NAME HERE

**F-710-001
Equipment Problem Report**

EQUIPMENT PROBLEM REPORT

EQUIPMENT DESCRIPTION: _____

LAST TASK PERFORMED: _____

JOB NUMBER: _____

DATE: _____ TIME: _____

OPERATOR: _____

REPORTED BY: _____

DESCRIPTION OF PROBLEM:

ACTION TAKEN

PROBLEM INVESTIGATED BY: _____

PROBLEM RESOLUTION DATE: _____

INSERT COMPANY NAME/LOGO HERE

AS 9120 Rev B - Quality Management Systems – The Internal Audit Checklist

This internal audit checklist is based on the information provided in the Nov 2016 revision of the AS 9120 Rev B, SAE international aerospace standard. The checklist is best used by trained and practicing auditors to evaluate or assess Quality Management Systems requirements based on the standard. You will see questions on the checklist that refer to the standard and for each clause provisions are made for additional questions.

The auditors are expected to keep in mind that the standard does not requires mandatory procedures for the various QMS processes; however, the auditors will expect documented information to be available because in the clauses of the standard, the phrase 'documented procedures' is used to specify that a process, a method, a system, a work instruction, or an arrangement be documented.

The auditors must use a great deal of discretion and therefore must be careful and thoughtful prior to establishing a deficiency against a requirement. Evidence for visible top management leadership, commitment and quality management action must be looked for.

The **bold** numbers and titles used in the first two columns of the checklist indicate the "Requirements" and may be referred to on nonconformity reports prepared by the auditor.

During assessment of each requirement, auditors record the status of the evaluation by indicating in the right-hand column a

Yes - for Acceptable Condition or **No** - for Deficient Condition

---	QUALITY MANAGEMENT SYSTEMS	OBSERVATIONS / COMMENTS	STATUS OK Yes / No
4	CONTEXT OF THE ORGANIZATION		
4.1	Understanding the organization and its context		
	Does your company determine the external and internal issues that are relevant to your purpose and strategic direction?		
	Do you consider the relevant issues that affect your ability to achieve the intended results of the Quality Management System (QMS)?		

INSERT COMPANY NAME/LOGO HERE

AS 9120 Rev B - Quality Management Systems – The Gap Analysis Checklist

This gap analysis checklist is prepared for use in evaluating a Quality Management System (QMS) against the requirements of the new Aerospace standard AS 9120 Rev B. Each requirement is expressed as a question that the user (auditor / assessor) can use to evaluate your QMS capabilities. You will need to have a copy of the AS 9120 B standard to use along with this checklist so that you can refer to the requirements and the clarification sections of Annex A. The intent of the main clauses of the new standard is shown in **blue font**.

After you have prepared an audit schedule, and assigned responsibility to your auditors for different areas or processes to audit, copy each section of the checklist for the auditors working with that section. As you work through the checklist take notes on what is in place, and what needs to be developed.

In the space for 'currently in place', list or reference the procedures or other documents, or evidence that you have reviewed and that will provide information for the new QMS. Take notes on the status of the documents, that is, will they need to be revised for the new system, or can they be used as is? Also, note where processes are in place, but documentation is needed. Focus on what is in place, and what needs to be developed.

While you do want to know if documented information is in place and if procedures and processes are being complied with, compliance is not your focus for this audit. Remember that the outcome of this audit should be a list of things that your company needs to do to comply with the AS 9120 Rev B standard.

---	QUALITY MANAGEMENT SYSTEMS REQUIREMENTS	Currently in Place	Compliant YES / NO?	If No - % Completed	Items Needed
4	CONTEXT OF THE ORGANIZATION				
Intend of clause	This first clause introduces two sub-clauses relating to the context of the organization, (1) understanding the organization and its context and (2) understanding the needs and expectations of interested parties. Together they require that you determine the issues and requirements that can impact on the planning of the Quality Management System (QMS). In addition, the scope of the QMS and the QMS processes along with their applicability and interactions need to be determined.				
4.1	Understanding the organization and its context				
	Does your company determine the external and internal issues that are relevant to your purpose and strategic direction?				

Risk Management

Every version of the AS 9120 standard has advocated risk avoidance and risk management. The new AS 9120 Rev B standard continues to expect organizations to identify and address risks affecting compliance of products and services, resulting in improved customer satisfaction.

Besides identifying the risks, organizations should address opportunities for improvements and corrective actions based on the risk analysis.

Note that while nonconformity and corrective action are requirements of AS 9120 Rev B, the concept of preventive action can be addressed through a risk-based approach where risks are determined and actions to address risks and opportunities are taken.

This risk analysis exercise is intended to outline several approaches / options for the management of risk at your company.

To prepare for the change, it is time to begin understanding Risk-Based-Thinking and begin looking at your processes in terms of risks.

Risk is defined as the combination of the probability of occurrence of harm and the severity of that harm.

When evaluating risk, it is helpful to address it using two (2) metrics or parameters:

1. Severity (if harm happens, how serious is the event)
2. Likelihood (what is the probability of a harmful event occurring)

Because this topic is so important, it will have an impact on your QMS.

Course Materials

The supplies you will need are:

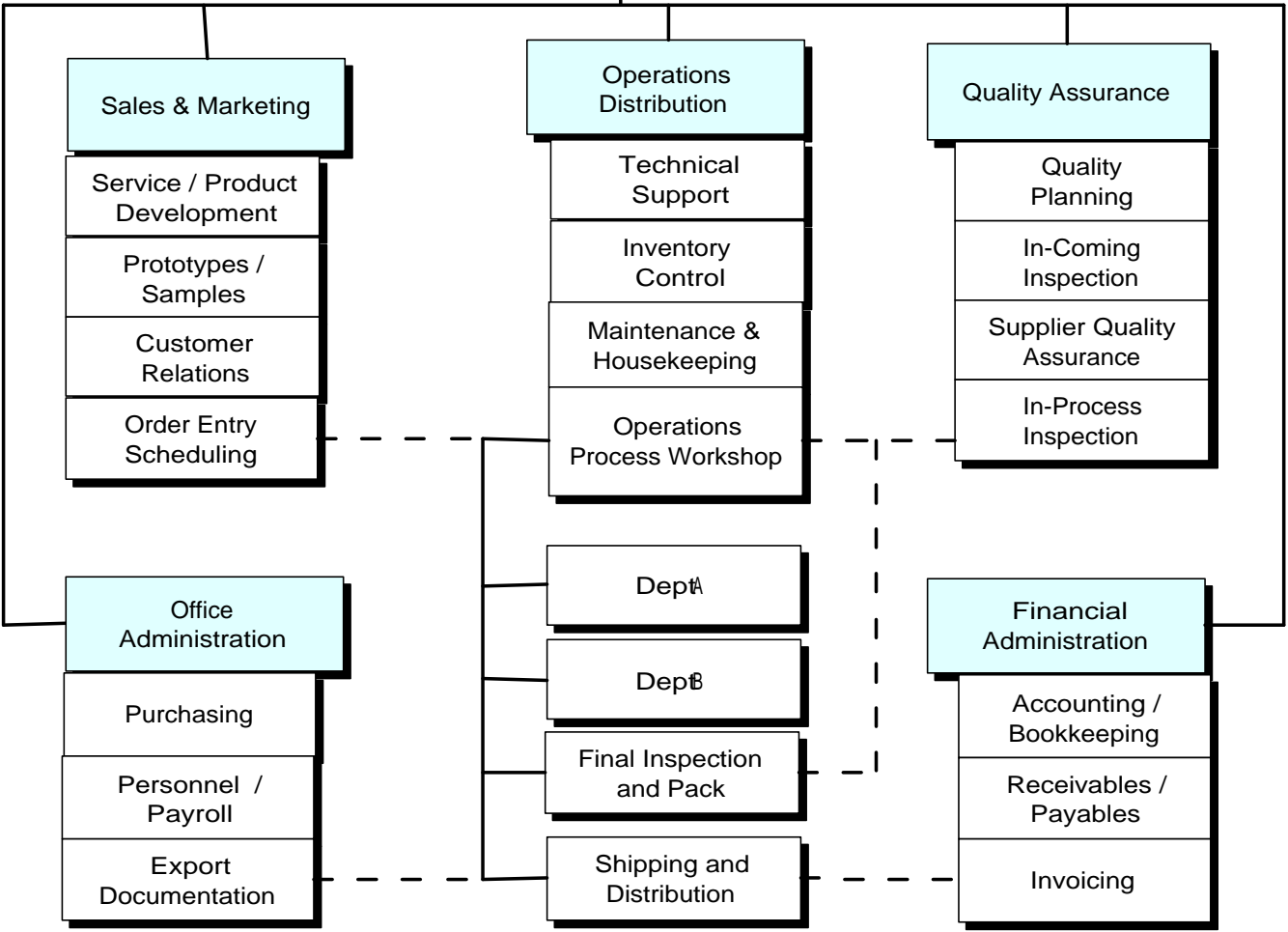
- PowerPoint: **Guide to Internal Audits** (included).
- PowerPoint: **Requirements of AS 9120 REV B** (included).
 - A complete version with Speaker Notes is in this Trainer's Guide
- PowerPoint: **Steps of Internal Audit** (included).
 - A complete version with Speaker Notes is in this Trainer's Guide
- Student Manual (included).
 - Print one copy for **each student**
 - You may wish to have extra copies of the CAR form
 - It includes reduced versions of all the PowerPoints.
- AeroSource Company - Documented Information (included).
 - Print one copy for **each team** of two or three students.
 - See next page for list of contents.
 - Note that for this training, it is not possible to bring all documents from a fictitious company in the classroom.
 - However, documents relevant to the audit and non-conformances observed are included. In the list of documented information, the relevant manual and procedures are highlighted in **brown font**.
- The AS 9120 REV B Standard (**NOT Included***)
 - You will need one copy for every 2-3 students.
 - Standards are available electronically from <http://www.techstreet.com/products>

The AS 9120 REV B Standard is a copyrighted document and we are unable to include it.

Typical - AS 9120 B
Organization Chart

President

Secretary



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Easy to customize with
your company name!

Welcome to AS 9120 Rev B

Our Company is working on becoming AS 9120 B registered. This international standard provides for a (QMS), Quality Management System that outlines some basic good business practices that we need to have in place for our distribution business.

By implementing a Quality Management System (QMS) that complies with AS 9120 B we will be able to make our company run more efficiently, increase customer satisfaction, and communicate to potential customers that we have good quality processes in place.

Surveyed AS 9120 Registered Companies state that they have:

- Higher customer satisfaction
- Increased profitability because of efficiencies
- Market advantages
- Improved communications
- Higher job satisfaction

What will employees need to do for the AS 9120 B Quality Management System?

First Management will be determining both the internal and external issues that are relevant to the QMS and will identify our “Key Processes”.

Those are the processes that affect the quality of our product and our services. Then they will determine how we will control these processes to make sure that we are all doing them the same way, and the best way our organization has identified.

Controlling the process means having documented information for the quality management system, and training employees or finding other or best ways

to make sure that the process is done consistently no matter who is doing it. This means that employees may be required to have certain training, or to follow specific work instructions.

Employees will also need to be aware of how their job affects the quality of our products and customer satisfaction.

AS 9120 Highlights: Things that you will be hearing about as we proceed with this project....

Our Quality Policy

We will identify our Quality Policy, and will be communicating it to all employees. It is important that all of us are aware of what this statement says about our company’s vision is for quality and for meeting customers’ expectations.

Registration Audit

To become AS 9120 B registered, an independent Registrar will audit our quality system. This Registration Audit will be done after we have set up processes to meet all the requirements of AS 9120 B.

The Registrar will send an auditor or audit team to come in to our facilities and evaluate the processes we have in place.

They will check to see if the processes meet the requirements of the standard, and to see if we are following the processes. If everything looks good, we will be recommended for registration and be recognized globally!



Watch for our next newsletter for more introduction to AS 9120 B, what it will mean to you and your coworkers.