



Supplier Survey		
Record Number QR-0840-02	Record Revision 1	Date of Revision 07/08/2020
Approved by: Jake Enger	Date Approved: 07/08/2020	Page 1 of 13

## SUPPLIER QUALITY SYSTEM QUESTIONNAIRE

Prepared By: (Name) \_\_\_\_\_ (Title) \_\_\_\_\_

Date: \_\_\_\_\_

Company Name: \_\_\_\_\_

Address: \_\_\_\_\_

City: \_\_\_\_\_ State: \_\_\_\_\_ Zip Code: \_\_\_\_\_

Phone: \_\_\_\_\_ Fax: \_\_\_\_\_ Email: \_\_\_\_\_

Major product or service: \_\_\_\_\_

Major Customers: \_\_\_\_\_

\_\_\_\_\_

\_\_\_\_\_

Total Plant area: \_\_\_\_\_

### Key Supplier Personnel

Senior Company Official: Name: \_\_\_\_\_ Title: \_\_\_\_\_

Senior Quality official: Name: \_\_\_\_\_ Title: \_\_\_\_\_

Other Key Personnel: Name: \_\_\_\_\_ Title: \_\_\_\_\_

Number of Quality personnel: \_\_\_\_\_ Total Number of Employees: \_\_\_\_\_

### General Information

**Manufacturing percentage:** % Military/Aerospace products: \_\_\_\_\_ % Commercial products: \_\_\_\_\_

Is your facility under Government QA cognizance? \_\_\_\_\_

Name and address of Gov't agency: \_\_\_\_\_

Quality Manual revision level and/or date: \_\_\_\_\_

**What Specification is your quality assurance system auditable to?**

**Please attach a copy of your Quality System Manual and Certifications from an Accredited Certifying Body. If you are certified to AS9100 or ISO9001, complete only Sections I, III, V, VI.**

<input type="checkbox"/> ISO9001	<input type="checkbox"/> ISO9002	<input type="checkbox"/> FAR 145
<input type="checkbox"/> AS9000	<input type="checkbox"/> AS9100	<input type="checkbox"/> Nadcap
<input type="checkbox"/> Other: _____		



### I. Special Processes

*Please list special processes or testing that your facility is approved to perform, (Include applicable specifications): Please attach any approvals, i.e. Nadcap, OEM, Etc.*

**Finishing/Coating:** \_\_\_\_\_  
\_\_\_\_\_  
\_\_\_\_\_

**Plating/Chemical Processing:** \_\_\_\_\_  
\_\_\_\_\_  
\_\_\_\_\_

**Welding/Brazing:** \_\_\_\_\_  
\_\_\_\_\_

**Heat Treating:** \_\_\_\_\_  
\_\_\_\_\_  
\_\_\_\_\_

**Non-Destructive Inspection:** \_\_\_\_\_  
\_\_\_\_\_  
\_\_\_\_\_

**Laboratory Testing:** \_\_\_\_\_  
\_\_\_\_\_  
\_\_\_\_\_

**Other (Special Processes):** \_\_\_\_\_  
\_\_\_\_\_  
\_\_\_\_\_  
\_\_\_\_\_  
\_\_\_\_\_

## II. Contract Responsibilities

<p>Is the system of quality assurance adequately described in management approved written procedures, instruction, and/or policies?</p>	<input type="checkbox"/> Yes	<input type="checkbox"/> No	<input type="checkbox"/> N/A
<p>Comments:</p>			
<p>Are such procedures, instructions, and /or policies maintained and made available to all affected personnel?</p>	<input type="checkbox"/> Yes	<input type="checkbox"/> No	<input type="checkbox"/> N/A
<p>Comments:</p>			
<p>Are quality procedures reviewed and upgraded at predetermined levels?</p>	<input type="checkbox"/> Yes	<input type="checkbox"/> No	<input type="checkbox"/> N/A
<p>Comments:</p>			
<p>Has the responsibility for quality assurance/inspection been formally established?</p>	<input type="checkbox"/> Yes	<input type="checkbox"/> No	<input type="checkbox"/> N/A
<p>Comments:</p>			
<p>Has the responsibility and authority for product quality been assigned to personnel with sufficient management stature to resolve problems effectively?</p>	<input type="checkbox"/> Yes	<input type="checkbox"/> No	<input type="checkbox"/> N/A
<p>Comments:</p>			
<p>Does quality assurance participate in contract reviews to identify and evaluate quality requirements prior to and during production?</p>	<input type="checkbox"/> Yes	<input type="checkbox"/> No	<input type="checkbox"/> N/A
<p>Comments:</p>			
<p>Does quality assurance review manufacturing, processing, packaging, and inspection/test instructions or procedures to assure their adequacy?</p>	<input type="checkbox"/> Yes	<input type="checkbox"/> No	<input type="checkbox"/> N/A
<p>Comments:</p>			
<p>Is there a provision for quality assurance to qualify production and laboratory acceptance tests?</p>	<input type="checkbox"/> Yes	<input type="checkbox"/> No	<input type="checkbox"/> N/A
<p>Comments:</p>			
<p>When tests are conducted by production or laboratory personnel, are results verified by quality assurance personnel to perform required inspection?</p>	<input type="checkbox"/> Yes	<input type="checkbox"/> No	<input type="checkbox"/> N/A
<p>Comments:</p>			
<p>Does the Supplier provide, through written instructions, procedures to notify customer of any changes to their inspection system or procedures?</p>	<input type="checkbox"/> Yes	<input type="checkbox"/> No	<input type="checkbox"/> N/A
<p>Comments:</p>			
	<input type="checkbox"/> Yes	<input type="checkbox"/> No	<input type="checkbox"/> N/A

Are flow down requirements i.e. drawings, and specifications adequately defined for jobs that require these items?

Comments: \_\_\_\_\_

### III. Documentation, Records, and Corrective Action

#### A. Inspection and Testing Documentation:

Are written instructions, in sufficient detail, provided for in-process and final inspection?  Yes  No  N/A

Comments: \_\_\_\_\_

Are written instructions, in sufficient detail, provided to cover all work affecting quality (handling, machining, assembly, fabrication, processing, cleaning, inspect/test, etc.)?  Yes  No  N/A

Comments: \_\_\_\_\_

Does quality assurance participate in the review of work instructions to verify contract requirements and specifications are included?  Yes  No  N/A

Comments: \_\_\_\_\_

Are in-process inspections documented in such a manner as to provide a positive inspection status of the material or parts?  Yes  No  N/A

Comments: \_\_\_\_\_

Are assembly and inspection operations and test results documented and validated by quality assurance on a traveler, work order, or other identifying document?  Yes  No  N/A

Comments: \_\_\_\_\_

Does Supplier have procedures outlining the acceptance criteria of inspection results?  Yes  No  N/A

Comments: \_\_\_\_\_

Are written instructions provided as a guide to preservation, packaging, marking, and shipping methods?  Yes  No  N/A

Comments: \_\_\_\_\_

#### B. Records:

Are all inspections and tests recorded and maintained on file for at least 11 years or more as required by contract?  Yes  No  N/A

Comments: \_\_\_\_\_

Yes  No  N/A

Do records indicate lot size, sample size, deficiencies found, quantities accepted and quantities rejected?

Comments: \_\_\_\_\_

Does the product and/or its associated inspection records identify operator and/or inspector performing the detailed operations?

Yes  No  N/A

Comments: \_\_\_\_\_

Do records indicate nature of deficiencies and corrective action taken?

Yes  No  N/A

Comments: \_\_\_\_\_

Are records of final inspection and test data maintained?

Yes  No  N/A

Comments: \_\_\_\_\_

Do inspection records indicate actual dimensions checked?

Yes  No  N/A

Comments: \_\_\_\_\_

**C. Corrective Action:**

Does the Supplier maintain an adequate corrective action system including verification that corrective actions have been implemented?

Yes  No  N/A

Comments: \_\_\_\_\_

Does the system ensure prompt responses to requests for corrective action?

Yes  No  N/A

Comments: \_\_\_\_\_

Does the Supplier have a system for corrective actions from subcontractors and Suppliers?

Yes  No  N/A

Comments: \_\_\_\_\_

**D. Drawings and Changes:**

Does the Supplier have a written procedure describing drawing change control necessary to meet customer requirements?

Yes  No  N/A

Comments: \_\_\_\_\_

Will Supplier's system assure that all manufacturing and processing is performed in accordance with the applicable revision of drawings & specifications as required by the contract?  Yes  No  N/A

Comments:

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Will this system serve to prevent the use of marked, illegible, or obsolete drawings which are not formally approved and controlled?  Yes  No  N/A

Comments:

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Is the system adequate to control customer furnished drawings and specifications?  Yes  No  N/A

Comments:

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Does the system assure that applicable technical documents, and changes thereto, are provided to subcontractors and Suppliers?  Yes  No  N/A

Comments:

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#### IV. Measuring and Test Equipment

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Are detailed procedures used for inspection and calibration of tools, gages, and test equipment?  Yes  No  N/A

Comments:

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Does the system adequately provide for mandatory recall of all calibrated tools, gages, and test equipment?  Yes  No  N/A

Comments:

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Is production tooling calibrated according to procedures at established intervals?  Yes  No  N/A

Comments:

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Are customer furnished tools, gages, and test equipment adequately controlled?  Yes  No  N/A

Comments:

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Are employee-owned tools and gages subject to the same controls as company-owned tools and gages?  Yes  No  N/A

Comments:

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Do standards, currently in calibration, have certifications on file that are traceable to NIST?  Yes  No  N/A



Comments:

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Are all tools, gages, and test equipment adequately stored and maintained?  Yes  No  N/A

Comments:

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Are uncalibrated or out-of-calibration items identified and/or stored in such a manner as to preclude their use prior to calibration?  Yes  No  N/A

Comments:

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Are tools, gages, and test equipment identified in a manner to reflect:

A. Date calibration/inspected?  Yes  No  N/A

B. Date due for recalibration?  Yes  No  N/A

C. Item identity or serial number?  Yes  No  N/A

Comments:

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## V. Process Control

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Are written procedures adequate to control all processes?  Yes  No  N/A

Comments:

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Are requirements as stated in the contract and/or specifications included in written procedures?  Yes  No  N/A

Comments:

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Are the written procedures an integral part of the inspection system?  Yes  No  N/A

Comments:

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Is there a procedure in place for Counterfeit Part avoidance?  Yes  No  N/A

Comments:

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Does the supplier have an established Foreign Object Damage (FOD) program in place that meets NAS412?  Yes  No  N/A

Comments:

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## VI. Indication of Inspection Status

Is the inspection status of products and materials indicated by stamps, tags, routing tickets, or other control methods as appropriate?  Yes  No  N/A

Methods: \_\_\_\_\_

Comments: \_\_\_\_\_

Enter typical inspection status stamps below or attach a copy of procedures that identify the quality assurance stamps used by your company.

## VII. Government/Customer Furnished Material

Is government/customer furnished material:

- A. Checked for damage upon receipt?  Yes  No  N/A
- B. Inspected to applicable drawings and specifications?  Yes  No  N/A
- C. Stored to preclude damage?  Yes  No  N/A
- D. Functionally tested as required by contract?  Yes  No  N/A
- E. Controlled by identification and segregation?  Yes  No  N/A
- F. Verified for quantity received?  Yes  No  N/A

Comments: \_\_\_\_\_

## VIII. Damaged Government/Customer Furnished Material

Yes  No  N/A



Is a written system in place to report damaged government/customer furnished material to the government or customer?

Comments:

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Does the system provide for the identification and segregation of damaged material?  Yes  No  N/A

Comments:

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Does the system provide for the determination of probable cause and necessity for withholding material for use?  Yes  No  N/A

Comments:

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### IX. Nonconforming Material or Product

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Does the supplier have formal procedures and an established system for:

A. Detection of nonconforming items?  Yes  No  N/A

B. Segregation of Nonconforming items?  Yes  No  N/A

C. Identification and/or marking of nonconforming items?  Yes  No  N/A

Comments:

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Is a holding area used for nonconforming materials or products?  Yes  No  N/A

Comments:

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Are deviations that affect the customer's requirements referred to the customer for disposition?  Yes  No  N/A

Comments:

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Is there a documented procedure for the repair or rework of nonconforming material or product?  Yes  No  N/A

Comments:

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Is nonconforming material or product identified to the document used for rejection?  Yes  No  N/A

Comments:

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Is Quality Assurance represented in the materials review board (MRB) activity?  Yes  No  N/A

Comments:

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Do records of nonconformance and MRB action reflect adequate descriptions of deficiencies and subsequent corrective action?

Yes  No  N/A

Comments:

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## X. Purchasing & Supplier Quality

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Are adequate written procedures in use for the control of purchased materials and services?

Yes  No  N/A

Comments:

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Does quality assurance review procurement documents for the inclusion of quality requirements?

Yes  No  N/A

Comments:

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Are purchase orders and applicable drawings, specifications, inspection and test instructions, etc. available at the time and place of inspection?

Yes  No  N/A

Comments:

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Are supplier's quality programs evaluated by internal auditing?

Yes  No  N/A

Comments:

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Are the supplier's special processes monitored at adequate intervals by testing at receiving inspection, surveillance or process surveys?

Yes  No  N/A

Comments:

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## XI. Sampling Inspection

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Are sampling inspection instructions available to inspection personnel?

Yes  No  N/A

Comments:

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Do instructions provide for tightening or reducing inspection when previous inspection data warrants?

Yes  No  N/A

Comments: \_\_\_\_\_

Is sampling performed to Mil-STD-105, other approved sampling plan and/or customer requirements?     Yes     No     N/A

Comments: \_\_\_\_\_

Are different inspection indications (stamps, Inspection forms, etc.) used to identify sampling versus 100% inspection?     Yes     No     N/A

Comments: \_\_\_\_\_

Do inspection records include lot size, sample size, and lot identity?     Yes     No     N/A

Comments: \_\_\_\_\_

**XII. Inspection Provisions**

Is the measuring/testing equipment used for supplier's inspection adequate to control the products being inspected?     Yes     No     N/A

Comments: \_\_\_\_\_

Is the customer consulted in the event of changes to the inspection procedure?     Yes     No     N/A

Comments: \_\_\_\_\_

**XIII. Receiving Inspection**

Does receiving inspection check incoming shipments to requirements of the purchase order, applicable drawings, and specifications?     Yes     No     N/A

Comments: \_\_\_\_\_

Are incoming materials identified to the applicable purchase order or material certifications?     Yes     No     N/A

Comments: \_\_\_\_\_

Are Materials, which are accepted on the basis of test reports and /or certificates of conformance, subjected to verification testing?     Yes     No     N/A

Comments: \_\_\_\_\_

Yes     No     N/A



Are test reports or certificates of chemical or physical analysis requested when applicable and maintained on file?

Comments:

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Do receiving inspection records indicate acceptance or rejection of incoming materials or product?

Yes     No     N/A

Comments:

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Are records of receiving activities maintained for at least 7 years or more if specified on the contract?

      

Comments:

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Are there adequate controls for handling and protection of received goods in use?

Yes     No     N/A

Comments:

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Are inspected items properly segregated from material awaiting inspection?

Yes     No     N/A

Comments:

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Are materials properly handled and stored to prevent damage, contamination, and/or deterioration?

Yes     No     N/A

Comments:

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Is there a procedure in place for Counterfeit Part avoidance?

Yes     No     N/A

Comments:

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Does the supplier have an established Foreign Object Damage (FOD) program in place that meets NAS412?

Yes     No     N/A

Comments:

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**General Comments:**

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Please mail, email or fax this form upon completion to the following:



Mac Machine Company, Inc.  
7209 Rutherford Rd.  
Baltimore, MD 21244  
Attn: Amanda Tomaj  
Phone: (410)944-6171, Fax: (410)944-6832  
amanda@macmachine.com

<b>Result of Survey:</b> Approved <input type="checkbox"/>	Conditional Approval <input type="checkbox"/>	Disapproved <input type="checkbox"/>
<b>For Mac Machine use only</b> Reviewed by: _____ Date: ___ / ___ / ___		

REV	DESCRIPTION	DATE	AUTHORIZED BY
-	INITIAL RELEASE – RECREATED IAW AS9100 REV. D	08/28/2017	J. ENGER
1	UPDATED RESPONSE INFORMATION - DCR-0069	07/08/2020	J. ENGER