



Fuji Autotech USA, LLC
Walton, KY 41094

DATE: _____

SUPPLIER QUALITY SELF AUDIT & QUESTIONNAIRE FORM

DUNS# _____
Supplier Name _____
Complete Mailing Address _____
Complete Shipping Address _____
(if different then mailing
address) _____
Main Phone Number _____
Main Fax Number _____
Commodity to supply FAU _____
FAU Purchasing Contact _____
Company Web Page Address _____

	Name/Title	e-mail Address	Phone Number	Fax Number
Main Contact Person/Title				
Plant Manager/or Title				
Plant Quality Manager				
Designated QR/CAR				
Contact & Title				
Backup QR/CAR Contact				
& Title				

Quality Certifications &/or Awards	Date of Original Certification	Last Re-Certification Date	Next Re-Certification Date	If not Certified to TS16949, ISO 9001:2008 or ISO14001, list projected certification date
ISO/TS16949				
ISO 9001:2008				
QS9000				
ISO14001				
Other				

Return completed form to: Fuji Autotech USA, LLC
70 Precision Drive
Walton, KY 41094

Attention: Purchasing Department

SUPPLIER QUALITY SELF AUDIT & QUESTIONNAIRE FORM

The purpose of this questionnaire is to provide a general understanding of the nature of the supplier's quality organization and approach to quality. The responses to these questions should be provided by the facility that will be supplying the product. Please answer each question by typing, or printing in ink, an "X" in the appropriate box, or by filling in the blanks. Return the completed questionnaire to Fuji Autotech USA, LLC. The questionnaire will be distributed to the appropriate personnel for evaluation and possible follow-up audit.

Plant Information (For this location):

1. Basic products:

2. Customers:

3. Major material suppliers:

4. Do you utilize just in time (JIT) delivery? Y or N

5. Do you utilize FIFO? Y or N

6. Do you have product liability insurance, and at what amount? \$_____

7. Total facilities area: _____sq. ft.

8. What percent is utilized capacity?_____%

9. Personnel:

Total employees:_____

Mfg. hourly:_____

Total salaried:_____

Quality hourly:_____

Quality salaried:_____

Engineering:_____

10. Union affiliation:_____

Date contract expires:_____

11. List laboratory/test facilities:

12. Describe prototype capabilities:

13. Do you use CAD/CAM? Y or N

14. Organizational structure: (Attach a copy of both the Plant & Quality Organization charts)

15. Please attach copy of all Third Party Registrations

SUPPLIER QUALITY SELF AUDIT & QUESTIONNAIRE FORM

		YES	NO
A.	General Systems		
1.	Is the Quality Department a separate/distinct part of your organization?	_____	_____
2.	Do you have a Quality Manual?	_____	_____
3.	Are there written procedures defining the quality-related functions?	_____	_____
4.	Does the Quality Department have the authority to control nonconforming material?	_____	_____
5.	Do you have a documented quality related training program?	_____	_____
6.	Do you have a system for customer complaints and rejections?	_____	_____
7.	Do you have a specific department, or organization responsible for advanced quality planning on new products?	_____	_____
8.	Do you have a procedure for approval of new & rebuilt machinery/equipment?	_____	_____
9.	Do you have a quality cost reporting system?	_____	_____
10.	Do you regularly perform internal quality system audits to assure compliance to your existing quality procedures?	_____	_____
11.	Does your facility have third party quality registration to QS9000, ISO 9001:2008, or TS 16949?	_____	_____
12.	Does your procedure require customer notification of continued registration and / or "loss of third party registration"?	_____	_____
13.	Does your procedure require that "loss of third party registration" be communicated to your customer?	_____	_____
B.	Statistical Process Control		
1.	Do you have an ongoing program to train employees in SPC?	_____	_____
2.	Do you have documented SPC for significant product characteristics & process parameters?	_____	_____
3.	Is manufacturing responsible for SPC?	_____	_____
4.	Does your SPC program require (min) 1.33 Cpk & 1.67 Ppk?	_____	_____
5.	In instances where process capability has not yet been demonstrated, do you require a documented action plan to improve the process?	_____	_____
6.	Do you use statistical techniques such as Design Of Experiments, Pareto analysis, regression and correlation analysis, etc.?	_____	_____
C.	Drawing and Specification Control		
1.	Does the quality function review related drawings and specifications for new products before acceptance?	_____	_____
2.	Are controls in place to assure the latest drawings, change notices and specifications are in use by manufacturing and quality personnel at the point of production and inspection?	_____	_____
3.	Are there documented procedures for drawing & specification control and data control procedures?	_____	_____
4.	Is there a documented record retention procedure? (Ex., Life of Product & Service + 1 year)	_____	_____
D.	Measuring and Test Equipment Control		
1.	Is all measuring and test equipment uniquely identified?	_____	_____
2.	Do you have a documented calibration and control system in place?	_____	_____
3.	Are calibration standards traceable to the National Institute of Standards and Technologies (NIST) or appropriate international standards?	_____	_____
4.	Are GR&R's performed on all types of measuring devices?	_____	_____

SUPPLIER QUALITY SELF AUDIT & QUESTIONNAIRE FORM

	YES	NO
E. Purchased Productive Material and Service Control		
1. Is a list of approved suppliers maintained?	<input type="text"/>	<input type="text"/>
2. Are supplier quality audits/surveys conducted?	<input type="text"/>	<input type="text"/>
3. Are suppliers required to use SPC and are records/data available?	<input type="text"/>	<input type="text"/>
4. Are in-house or external heat treat sources approved to specific or specified Heat Treat specifications?	<input type="text"/>	<input type="text"/>
5. Do you require supplier material certifications & routinely verify same?	<input type="text"/>	<input type="text"/>
6. Are inspected materials uniquely identified from those awaiting inspection?	<input type="text"/>	<input type="text"/>
7. Do you have controls to prevent movement of rejected incoming material to storage or point of use?	<input type="text"/>	<input type="text"/>
8. Do you have AIAG APPROVED bar coding capabilities?	<input type="text"/>	<input type="text"/>
9. Do you have product traceability from incoming through shipped product?	<input type="text"/>	<input type="text"/>
10. Do you have documented procedures for contract review?	<input type="text"/>	<input type="text"/>
11. Did you read the terms and conditions agreement at http://www.fujiautotec.com/supplier/termsandconditions.html and do you agree with the terms and conditions?	<input type="text"/>	<input type="text"/>
12. Did you read the FAU Supplier Requirements Manual located at http://www.fujiautotec.com/supplier/fausupplierrequirement.html and can you meet all the requirements identified in the FAU Supplier Requirements manual?	<input type="text"/>	<input type="text"/>
F. Internal Non-Conforming Material Control		
1. Are controls in effect to prevent movement of nonconforming or suspect material back into the normal production flow?	<input type="text"/>	<input type="text"/>
2. Is nonconforming product analyzed to determine cause and extent?	<input type="text"/>	<input type="text"/>
3. Do you take corrective action to prevent recurrence of nonconformity?	<input type="text"/>	<input type="text"/>
4. Is there a procedure for timely disposition for non-conforming material?	<input type="text"/>	<input type="text"/>
5. Do you have segregated hold areas for nonconforming material?	<input type="text"/>	<input type="text"/>
6. Is reworked or sorted material resubmitted for inspection?	<input type="text"/>	<input type="text"/>
G. In-Process Control		
1. Is first piece (heat, batch, lot, ladle analysis) inspection approval required?	<input type="text"/>	<input type="text"/>
2. Do you perform in-process product audits?	<input type="text"/>	<input type="text"/>
3. Is material identified throughout the process, with unique identification?	<input type="text"/>	<input type="text"/>
4. Are control plans and/or operator instructions available at each operation?	<input type="text"/>	<input type="text"/>
H. Finished Product Verification		
1. Does finished product receive a final inspection by the Quality Department?	<input type="text"/>	<input type="text"/>
2. Are inspection/test procedures or instruction sheets used?	<input type="text"/>	<input type="text"/>
3. Do you perform PPAP's submissions for all customers per the current edition of the AIAG PPAP Manual?	<input type="text"/>	<input type="text"/>
4. Does your PPAP process require customer notification and concurrence (designated using facility) prior to shipping product?	<input type="text"/>	<input type="text"/>
5. Does your PPAP process include provisions for yearly customer PPAP submission (including sub-components if applicable)?	<input type="text"/>	<input type="text"/>
I. Customer Deviation / Concession		
1. Do you have a deviation/concession procedure?	<input type="text"/>	<input type="text"/>
2. Does it include customer approval authority for the deviation / concession before the shipment of product?	<input type="text"/>	<input type="text"/>
3. Do you have a procedure that requires notification to FAU for a Change Request? (Changes to Machines, Processes, Materials, Design ect. including Sub-Supplier changes)	<input type="text"/>	<input type="text"/>
J. Customer Complaints / Quality Rejections		
1. When Customer complaints are received, Do you have a method of ensuring all departments of the organization are notified (including Upper Management)?	<input type="text"/>	<input type="text"/>
2. Do procedures require that your response to nonconforming material include (D3) Containment actions and response within 24 hours, (D6) Permanent Corrective Actions and response within 7 days, and (D8) Completion within 30 Days?	<input type="text"/>	<input type="text"/>
3. Do procedures require Customer concurrence with closure of completed corrective actions?	<input type="text"/>	<input type="text"/>

Person completing this questionnaire, please complete following:

Print Name/Title: _____ Date: _____

Fuji Autotech USA, LLC Review

Quality (Name): _____ Date: _____

Purchasing (Name): _____ Date: _____

Is follow-up visit required? YES NO

Revision History

[illegible]