ANDREA SYSTEMS LLC SUPPLIER QUALITY SYSTEM SURVEY

Supplier: Address: City and State: Country: Telephone: Fax:		Date: Survey Format: On-Site Self Self Survey By: Title:			
PRODUCTS MANU	JFACTURED:				
FACILITIES:	Total Plant Area (Sq. Feet):	Number of Buildings:			
EMPLOYEE INFOR	RMATION				
Number of Employe	es: In Production: In	Quality Assurance: Total:			
REGISTERED:					
ISO 9001 □ ISO 9002 □ ISO 9003 □	Mil-I-45208 FAR 21 FAR 145	Registrar: Certificate Number: Date of Last Audit:			
If not registered, plea	ase state if you are compliant				
NOTE: This survey	needs to be completed only to this point	if supplier is ISO 9000 Registered.			
CLASSIFICATION	OF SUPPLIER BASED ON SURVEY:				
Approved:	Provisional:	Working Toward Conformance: □ Approval Withheld: □			
THIS SECTION TO	D BE COMPLETED BY ANDREA SYS	TEMS LLC QUALITY ASSURANCE			
NAMES AND TITL	ES OF INDIVIDUALS CONTACTED:				
OBSERVATIONS /	OBSERVATIONS / RECOMMENDATIONS:				

1.0 1.1	MANAGEMENT RESPONSIBILITY Does the person responsible for Quality Control at the plant report to the Plant Manager level or higher? a. Is quality their sole responsibility? Comments:	YES	
1.2	Does management have periodic reviews concerning quality? a. With all employees? b. Is there a Quality Policy? c. Is it well communicated? Comments:		
2.0	QUALITY SYSTEM		
2.1	Has the quality system been documented in a quality manual?		
	Comments:		
2.2	Is the quality manual current and controlled? Comments:		
2.3	Are all the relevant elements of this survey included in the quality system documentation? Comments:		
3.0	CONTRACT REVIEW		
3.1	Does the company have a written system for incorporating customer requirements into specifications?		
	Comments:		

		<u>YES</u>	<u>NO</u>	<u>N/A</u>
3.2	Are changes to the accepted order effectively communicated to affected departments within the organization?			
	Comments:			
4.0	DESIGN CONTROL			
4.1	Are there written procedures for control of product design?			
	Comments:			
4.2	Do all design changes require review and approval?			
	Comments:			
4.3	Is there a plan/system for testing new designs for reliability?			
	Comments:			
4.4	Is there immediate access to current ASTM, ANSI, etc. specifications as referenced in customer agreements?			
	Comments:			
5.0	DOCUMENT CONTROL			
5.1	Are there written procedures to control quality documents and data?			
	a. Drawings?			
	b. Work Instructions?			
	c. Specifications?			
	d. Test & Inspection Results?			Ц
	Comments:			

5.2	Are documents available to	personnel o	operating the	processes?
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- a. Do drawings exist where needed?
- b. Do work instructions exist where needed?
- c. Are procedures available where needed?
- d. Are all documents controlled?

Comments:

5.3 Are changes in the following documents reviewed and approved?

- a. Drawings?
- b. Work Instructions?
- c. Specifications?
- d. Procedures?

Comments:

6.0 PURCHASING

6.1	Does the supplier selection process involve supplier evaluations?a. Are Site Audits performed?b. Are Supplier Quality Surveys utilized?c. Are suppliers rated by a common measurement?		
6.2	Comments: Is there a list of approved suppliers? Comments:		
6.3	Is the quality performance of suppliers recorded and reviewed on regular intervals with the supplier? Comments:		

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7.0 CUSTOMER SUPPLIED PRODUCT

7.1	Are there written procedures for verification, storage and preservation of customer supplied items?		
	Comments:		
8.0	PRODUCT IDENTIFICATION AND TRACEABILITY		
8.1	Have individual product or batches been uniquely identified (date code, serial number, etc.)?		
	Comments:		
9.0	PROCESS CONTROL		
9.1	Are Key Processes which directly affect quality identified?		
	Comments:		
9.2	Are there documented work instructions for each Key Process?		
	Comments:		
9.3	Have Key Product and Process Characteristics been defined?		
	Comments:		
10.0	INSPECTION AND TESTING		
10.1	Is incoming material inspected according to a plan?		
	Comments:		
10.2	Are there procedures for verification of incoming materials?		
	Comments:		

10.3	Are records kept to show acceptance and rejection of incoming material?		
	If so, how long are these records kept on file?		
	Comments:		
10.4	Are written procedures provided for in-process inspections and tests?		
	Comments:		
10.5	Is final inspection and testing done according to a documented plan?		
	Comments:		
10.6	Are there records to show that finished product has passed test?		
	Comments:		
10.7	Are test results available for customer review? Comments:		
	INSPECTION, MEASURING AND TEST EQUIPMENT		
11.1	Is there a procedure for calibration of inspection / test equipment?		
	Comments:		
11.2	Has all inspection, measuring and test equipment used for product acceptance been calibrated?		
	Comments:		

11.3	Are calibration records (including gage history and a recall file) available?		
	Comments:		
11.4	Is calibration status shown by tool marking and record?		
	Comments:		
11.5	Are calibration standards traceable to NIST?		
	Comments:		
12.0	INSPECTION AND TEST STATUS		
12.1	Are inspection stamps, markings, tags, labels or other suitable means for identifying the status of products limited and controlled?		
	Comments:		
13.0	CONTROL OF NONCONFORMING PRODUCT		
13.1	Is reworked product re-inspected according to documented procedures?		
	Comments:		
13.2	Are there written procedures for disposition of non-conforming product?		
	Comments:		
13.3	Are there written procedures for review of repetitive discrepancies?		
	Comments:		

14.0	CORRECTIVE AND PREVENTATIVE ACTION		
14.1	Is a documented system implemented to effectively handle customer complaints and reports of product nonconformities?		
	Comments:		
14.2	Are there implemented procedures that determine the corrective action needed to eliminate the cause of nonconformities?		
	Comments:		
15.0	HANDLING, STORAGE, PACKAGING, PRESERVATION AND DELIVERY		
15.1	Are there implemented procedures for handling, storage, and delivery of product?		
	Comments:		
15.2	Are documents available for controlling packing and packaging?		
	Comments:		
15.3	Are materials in stock adequately identified?		
	Comments:		
16.0	QUALITY RECORDS		
16.1	Are quality records legible?		
	Comments:		
16.2	Are quality records identifiable to the product? Comments:		

17.0 INTERNAL QUALITY AUDITS

17.1	Are there procedures covering scope of, plan for, frequency of, and responsibility for internal auditing?			
	Comments:			
17.2	Are all areas of the quality system being audited periodically?			
	Comments:			
17.3	Is timely corrective action taken to correct deficiencies?			
	Comments:			
18.0	TRAINING			
18.1	Is specialized job training given to key employees to assure quality?			
	Comments:			
18.2	Have various training needs been identified by job classification?			
	Comments:			
19.0	SERVICING			
19.1	Are there procedures for servicing customer-owned product as required by purchase contract?			
	Comments:			
20.0	STATISTICAL TECHNIQUES			
20.0	Has the need for statistical techniques been identified in order to control processes?	п	П	п
	Comments:			

20.2 Are there written procedures that define the application of statistical techniques?

Comments:

21.0 CERTIFICATION / TRACEABILITY AND MATERIAL CONTROL

- 21.1 Does your company's Certificate of Conformance (CoC) with each shipment contain the following:?
 - a. Andrea Systems LLC Part Number
 - b. Andrea Systems LLC Purchase Order Number
 - c. Revision Level
 - d. Name of Manufacturer
 - e. Lot / Batch Numbers
 - f. Manufacturers Part Number
 - g. List Mil, ASTM, SAE Standards appearing on Andrea Systems LLC Drawings or Purchase Order

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