NewGenAerospace
Integrity • Commitment • Solutions

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SUPPLIER AUDIT FORM

In order for your firm to be placed on our Approved Supplier List, it is necessary that the responsible person in your firm fill out this audit form and return it to us via mail, fax, or e-mail. Please include copies of any Certificates attesting to the quality system in use.

Company	
Address	
City	
State	
Zip Code	
Country	

Name	
Title	
Phone	
Fax	
E-mail	

Quality System in use	

I certify that the information contained within this document is true and correct.

Signature:	**Date:**

Approved	Not Approved
Comments	
By:	
Date:	

New Gen Aerospace Corp. Supplier Audit Form

	Y	N	N/A
1. Quality System and Manual			
A. Is there an established quality system and a quality manual?			
B. Is the quality manual available to appropriate personnel?			
C. Is the quality system documentation kept current and readily available to employees, customers, auditors or designee(s)?			
D. Does the quality control manual include a detailed description of:			
1) the organization and relationship of the QC department to the rest of the organization?			
2) the assignment of personnel by title, for specific functions within the quality system?			
3) the revision control system for the quality system documentation?			
4) record keeping system?			
5) training requirements and records?			
6) shelf life control system?			
7) control of incoming discrepant parts and supplies?			
8) receiving inspection procedures?			
9) test and inspection equipment calibration program?			
10) storage facilities and specifications?			
11) part identification system?			
12) environmental controls?			
13) inspection stamp control?			
14) self-audit/evaluation program?			
2. Self-Audit/Evaluation Program			
A. Is there an established documented self-audit/evaluation program, which identifies who within the company is responsible for conducting self-audits, the frequency of audits, audit documentation and corrective action?			
3. Facilities			
A. Does the storage areas provide:			
1. adequate space and appropriate racks to prevent damage or mishandling?			
2. adequate security from unauthorized access?			
3. segregation of aircraft from non-aircraft functions?			
4. segregation of serviceable from non-serviceable parts?			
4. Training and Authorized Personnel			
A. Are personnel who perform inspection, shipping and receiving functions properly trained?			
B. Are inspection personnel properly authorized?			
C. Are both formal classroom and on-the-job training documented and maintained?			
D. Is a roster of personnel authorized to perform inspection functions maintained?			
E. Is there a training program in place for personnel involved in procurement, receiving inspection, shipping inspection and material control, that addresses unapproved parts, and counterfeit parts and materials?			

	Y	N	N/A
5. Procurement			
A. Does the system assure that parts procured conform to the customer's documentation requirements?			
B. Does the quality system assure that parts conform to the customer's purchase request and that deviations are disclosed and approved by the customer?			
C. Does the system require the distributor/dealer to maintain a list of approved suppliers and a quality history for each source?			
D. Does the quality system assure that parts procured for sale:			
1) which are known to have been subjected to conditions of extreme stress, heat or environment are identified?			
2) that all represented Airworthiness Directives (AD's) which have been accomplished are documented?			
3) that are identified as overhauled, repaired or modified have all appropriate signed and dated documentation?			
6. Receiving Inspection			
A. Does the quality system provide for a visual inspection of all items received and accompanying documentation?			
B. Is there a procedure for reporting unapproved parts in accordance with FAA Advisory Circular 21-29?			
C. Is there an accountability system in place to control stamp issuance, usage and replacement?			
7. Measuring and Test Equipment			
A. Is there an effective calibration program for test equipment?			
8. Material Control			
A. Is material handled in an appropriate manner and is the material protected from damage & deterioration?			
B. Is batch/lot control maintained for parts so identified by the manufacturer?			
C. Is there a system in place for recall control which ensures that parts shipped can be traced and recalled?			
D. Whenever practical, is material stored & delivered in the manufacturer's original packaging?			
E. Does the system specify material control requirements for material subject to damage by electrostatic discharge?			
F. Does the system assure that serviceable parts/components are adequately protected against the environment?			
G. Does the system assure that no part number ambiguity exists?			
H. Does a closed loop system exist to implement corrective action following detection of substandard or nonconforming parts?			
1) are aircraft parts being segregated from non-aircraft parts?			
I. Is there a documented procedure in place to mutilate scrapped parts to prevent the possibility of their being restored and returned to service?			
J. Are suspected unapproved parts reported to the FAA according to AC 21-29 or to the appropriate CAA?			

	Y	N	N/A
9. Shelf Life Control			
A. Does the distributor have a system for identifying and controlling shelf life limited parts?			
10. Certification and Release of Materials			
A. Does the system call for providing the customer with appropriate documentation?			
B. Does the system provide for the issuance of a certified statement disclosing that the material or parts were or were not:			
1) subjected to conditions of extreme stress, heat or environment;			
2) obtained from any government or military source			
11. Shipping			
A. Does the quality system require shipments in ATA-300 containers or equivalent as appropriate for the unit being shipped, or as specified by the customer?			
B. Does the quality system provide for a visual inspection of all items and accompanying documentation prior to shipping?			
12. Records			
A. Does the record system require record retention for at least 7 years from the date of sale to the customer?			
B. Does the system require all life-limited parts have records confirming current life limited status?			
C. Are records protected against damage, alteration, deterioration and loss?			
13. Technical Data Control			
A. Does the quality system provide for maintaining technical data in a manner which ensures such data is up-to-date and accessible?			