

Please find attached the AAF International - Air Filter Products Supplier Questionnaire. As part of our ISO 9001 Quality Certification, we maintain a continuously evaluate Supplier information. This questionnaire is an important part of our qualifying process and Approved Suppliers Rating System.

Upon review of the questionnaire you will find several key areas that AAF focuses on when qualifying suppliers.

**Quality System**: Consistent quality management requires a documented quality system, a written quality philosophy, quality goals and objectives against which progress is tracked and an organizational structure which clearly defines lines of authority. The system should include formalized root cause, corrective and preventative action procedures to non-conformance issues.

**Design Information Control**: AAF uses detailed manufacturing specifications, keyed against AAF part numbers, for the products and services we purchase. Our suppliers adherence to these specifications is the foundation of our quality policy to our customers and I the basis for material returns. Specification with revision dated are noted on our Purchase Orders. If there is any question or discrepancy in information or revision dates, and immediate call to AAF Purchasing should be made before continuing with the order. Our Supplier's ability to maintain, and keep current, a file of AAF specifications and the verification of each order to meeting these requirements is essential AAF requires the marking of our part number, and Purchase Order number on all cartons, packaging, bills of lading, packing lists and invoices.

**Customer Service**: Customer Service function should be clearly defined and provide a direct channel of communication to assure timely material delivery, prompt response to inquiries/ quotations, and accurate shipping and invoice documents. Resolution to material returns and non-conformance's is and important part of customer service.

**Continuous Improvement**: A continuous quality improvement process and cost containment program should be actively working and should provide direct long-term benefits relative to all phases of the organization.

Please complete the questionnaire and return it to the address below. We value our relationship with our Suppliers, and feel that with regular communication and review of each others expectations, an environment of mutual growth and continuous improvement is created.

AF Purchasing P.O. Box 35690 Louisville, KY 40232 (502) 637-0603 phone (502) 637-0315 fax



### **General Business Section**

Company Name:			
Address			
Telephone Number:	Fax Number:		
Interment address:			
Type of Business: Partnership Corporation	Sole Proprietorship		
If a Corporation, please indicate state where inco	rporated:		
Date of Incorporation: Publicly	Traded Symbol:		
Years in Business: Number of Emplo	yees: Manufacturer:		
Distributor:			
Parent Company (if applicable):			
Is the organizational structure documented? If so provide a copy.			
• President			
Vice President-Sales			
Quality Manager			
Customer Service Manager			
General Plant Manager			
Primary Contact			
Primary Contact Telephone Number			
Union: Affiliated with:			



Contact Expiration Date:  Vacation Shut Down Period
Is your Company:
<ul> <li>Minority Business</li> <li>Minority Classification</li> <li>A Small Business Concern</li> <li>A Woman Owned Business Enterprise</li> </ul>
If so please provide copy of MBE, WBE, or Small Business Concern registration or certification.
Do you have written Standards of Business Ethics? If so, Please provide a copy.
What do you see as your company's core competencies:
Quality System
Quality management requires a documented quality system, a written quality philosophy, quality goals and objectives against which progress is tracked and an organizational structure which clearly defines lines of authority.
Is there a written quality mission statement or quality policy statement?  If so please provide a copy.
Is your company registered to ISO 9000?
Do you have other recognized quality registration?
If yes, Please provide details:
If not ISO registered, does a quality manual exist? If so please provide a copy.
Do you have a separate quality department?
How is it staffed?
Are statistically based procedures used to control the manufacturing process?
Do you have a formalized root cause, corrective and preventive procedures for non-conformance's?



### **Design Information Control**

A system should be in place to ensure that operating personnel have accurate, current and complete technical instructions for the manufacture and inspection, packaging and shipping of AAF products. This information includes drawings, specifications, engineering change order, inspection instructions, special purchase order instructions and other special information.

•	How do employees access design/manufacturing specification?
•	Are they computerized?
•	To what level in the organization is this information available on production floor?
•	Is there a documented procedure for reviewing, approving, and distributing specification changes?
•	Do procedures ensure that only current documents are in circulation?
•	Is there a master list showing who has controlled documents?
•	How are revisions properly tracked?
•	Does the operator have the authority to make necessary changes (up to shutting down machine if necessary)?
Fin.	al Acceptance Is there final acceptance procedures to ensure that all shipping units meet all specifications for product, packaging, markings, and documentation?
•	Do you provide certificates of analysis and or conformity with your products?
•	Who in authority signs certificates of conformity? Name: Title:
A c	ntinuous Improvement Process ontinuous quality improvement process should be actively working and should provide direct long-term benefits tive to all phases of the organization.
•	Do you have a formalized continuous improvement program?
•	Do you have a corporate statement endorsing continuous improvement?  If so please provide a copy.
•	Do you use a consumer inquiry/complaint data base?



### **Calibration Verification**

Good manufacturing practices should include a calibration program, verification program, preventive maintenance program, adequate cross checks, and documentation to show calibration and verification checks.

•	Are inspections and calibrations performed and documented at appropriate intervals for all process gauges and laboratory instruments?
•	Are calibration checks documented?
Pro	<b>aterial Control</b> curement control should ensure that material sources meet specified quality standards and provide an anterrupted source of supply.
•	Do you have mutually agreed upon specifications with your suppliers?
•	Do you have procedures to confirm that incoming materials meet specifications?
•	Is there a process to track and improve your supplier's performance?
•	Are "Non-Conforming" materials identified and segregated?
•	Are "Non-Conforming" materials stored in a designated place, separated from good materials?
•	Are provisions in place for customer approval of non-conforming materials disposition as "use as is"?
Fa	cilities & Equipment Capability
•	What is your current manufacturing schedule (2 shifts. 5 days per week)?
•	Do you have a contingency plan for operating during a work stoppage (strike)?
•	Do you have multiple manufacturing locations capable of producing the products used by AAF?
•	Do you have facilities for in house testing of tolerances, performance, materials, etc., on your products?
•	Is there a designated R & D Function?



### **Customer Service**

Customer Service function should be clearly defined and provide a direct channel of communi-cation to assure timely material delivery, prompt response to inquiries/quotations, and accurate shipping and invoice documents.

Do you monitor or track on-time delivery?	
What procedures exist to ensure and measure on-time delivery?	
Do procedures exist to ensure consistently accurate invoicing and documentation?	
Do procedures provide prompt resolution of rejected materials?	
Is there a procedure for inspecting and replacing non-conforming materials? If so please provide a copy.	
Who is designated to handle issues with non-conforming materials?	
Is this person authorized to issue return goods authorizations?	
Do you have procedures to track and respond to inquiries, complaints, requests?	
Is a person designated to handle requests and inquiries?	
Is there a designated response time? Is it tracked? Are inquiries and requests logged?	
Do you have EDI capabilities?currently used and what is the software capability?	What transits are
For example: transmission of text, quotes, order receipt, order acknowledgment, invoicing, EFT, etc	advanced ship notices
Are you capable of producing scannable codes to identify materials and products?	
Do you track lead-time?	



#### **Cost Control**

The supplier should be actively involved in cost control programs and should be able to demon-strate results of these activities.

Do you have programs to control and reduce costs in the areas of waste reduction, supplier cost comprovement, and technological advancement?	ntrol, productivity
Do you have specific cost reduction goals?  If so please provide a copy.	
Do you have a cost control program with your supplier?	
What method are used to control supplier cost?	
Do you use established cost standards?	
Supplier questionnaire must be reviewed and accepted by Purchasing Agent.	

### **Summary**

Thank you for the time and effort to complete this questionnaire. We do not take this effort lightly. The above information is considered confidential and is important to our Approved Supplier Certification. We view our suppliers as an extension of our own resources and an essential part of meeting our customer's requirements. If you have any questions please call.

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