## **Supplier Quality Questionnaire**

**Company Information** 



Please provide copies of all ISO certificates and all Classification Society Approvals to **Supplier.Quality@bergenengines.com** 

	Company Name		
	Address		
	Telephone Number		
	Fax Number		
	Principle Products		
	Timelple Froducts		
	Number of Employees		
	Name	Email Address	Phone Number
Quality Manager			
Managing Director			
Technical Contact			
Commercial Contact			
	le copies of certificates		
Quality System (	Certification Held by C	Company	
	Certification Held by C	Company	Approval Body
Quality System (	Certification Held by C 2015)	Company	Approval Body
Quality System (	Certification Held by C 2015)	Company	Approval Body
Quality System (	Certification Held by C 2015)	Company	Approval Body
Quality System (	Certification Held by C 2015)	Company	Approval Body
Quality System (ex: BS EN ISO 9001:2	Certification Held by Control  Standard  Vals Held by Company		Approval Body
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Quality System (ex: BS EN ISO 9001:2	Certification Held by Control  Standard  Vals Held by Company Oprovals held		Approval Body
Quality System (ex: BS EN ISO 9001:2	Certification Held by Control  Standard  Vals Held by Company Oprovals held		Approval Body





## **SECTION A - Quality**

Al. Do you maintain a manual/procedure describing your quality management system(s)?
A2. Do you maintain an internal audit program?
A3. Do you determine the necessary competencies for personnel performing work affecting product quality and maintain records of education, training, skills and experience?
A4. Do you conduct a formal review of order requirements ensuring that the requirements are fully understood and adequately communicated throughout the organization and supply chain?
A5. Are written instructions used which list the manufacturing sequence and describe the methods to be used on each job including inspection requirements?
A6. Is the inspection status of material and part identified throughout the receiving and manufacturing process?
A7. Is inspection equipment used subject to calibration controls?
A8. Are non-conforming materials and parts identified, segregated and stored in a designated area?
A9. Do you maintain a procedure for recording, analyzing, and prioritizing the management of problems (e.g., non-conformance)?



Alo. Do you continually improve the effectiveness of the quality management system through the use of quality policy, quality objectives, audit results, analysis of data, corrective and preventative actions and management review?
All. Do you have a documents procedure within your management system which controls source and method change?
SECTION B - Procurement
B1. Do your purchase orders flow down all requirements, including contract specific quality requirements, specifaction and revisions to sub-tier suppliers?
B2. Do you inspect purchased items upon receipt to verify that they meet purchase order requirements?
B3. Do you have a process for the selection, approval and maintenance of sub-tier suppliers?
B4. Are sub-tier suppliers subject to quality surveillance audit to validate systems that have an effect on product quality?
B4. Are sub-tier suppliers subject to quality surveillance audit to validate systems that have an effect on product quality?



## **SECTION C - Drawing and document change control**

C1. Are there documented procedures to ensure that the correct drawings, specifications and related documents are available for the procurement, manufacturing and acceptance of your products? **SECTION D - Health, Safety and Environment** D1. Is your management system certified to the Environmental Management System (EMS) standard ISO 14001 or registered under EMAS (Eco Management and Audit Scheme) #? D2. How is your environmental management system deployed? D3. Is your management system certified to the Health and Safety Management System standard OHSAS 18001 or equivalent #? D4. Do you assess your Health, Safety and Environmental risks and put appropriate controls in place?

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