

SUPPLIER SURVEY FORM Instructions

- 1. The following Supplier Survey was developed by Vishay Measurements Group, Inc. to assess and document the capability of its supplier base.
- 2. The Supplier Survey is based on the requirements of ISO-9001 and is a simple yes/no format. We realize that some questions may not apply to your particular operation. Exceptions or commentary may be attached.
- If you are ISO-9001 certified, or are certified to some other nationally recognized standard, please complete Section 1 of the survey and return it with a copy of your certificate or registration of approval. You do not need to complete Section 2.
- 4. If you are not certified to ISO-9001 or some other nationally recognized standard, please complete the entire survey.
- 5. Please return all information within 21 days, and direct any questions to:

Quality Program Manager VISHAY Measurements Group, Inc. PO Box 27777 Raleigh, NC 26711-7777

Phone: 919-365-3800 FAX: 919-365-3945 Email: survey@vishaypg.com

Approved:

(Electronic includes date and title)

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VISHAY MEASUREMENTS GROUP, INC. QUALITY ASSURANCE SUPPLIER SURVEY FORM

Supplier's Company Name:			
Address:			
City, State, Zip Code:			
Web-site:			
Contact Person:			
E-mail:			
Telephone #:	Fax #:		
Product(s) or service(s) provided:			
Survey completed by: Name: Ti	tle: Date:		
1. Is your company ISO certified? Yes 🗌 No	D Comment:		
2. If yes, name of Registrar:	Expiration Date:		
3. Are you considering ISO-9001 certification?	Yes 🗌 No 🗌 Comment:		
If yes, expected date of completion:			
4. Is your company ROHS Certified? Yes	No 🗌 Comment:		
5. Is your company Conflict Minerals Certified?	Yes 🗌 No 🗌 Comment:		
OTHER QUALITY-BASED CERTIFICATIONS			

OTHER QUALITI-BASED CERTIFICATIONS				
Туре	Regulatory Body	Date of Certification		

The information you provide in this questionnaire is important for the establishment and/or maintenance of your Approved Supplier status at VISHAY Measurements Group, Inc.

If your company is ISO-9000 certified, stop here and attach a copy of your certification.

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Section 2

	2.1 Management Responsibility					
1.	Do you have a documented quality policy?	Yes 🗌	N/A 🗌	No 🗌		
2.	Is there a system for dissemination of directives and instructions to all personnel within the organization?	Yes 🗌	N/A 🗌	No 🗌		
3.	Is the commitment to quality known and understood by all employees?	Yes 🗌	N/A 🗌	No 🗌		
4.	Do you have a formal documented process to measure customer satisfaction?	Yes 🗌	N/A 🗌	No 🗌		
	2.2 Quality System					
1.	Do you have documentation defining the Quality System in use?	Yes 🗌	N/A 🗌	No 🗌		
2.	Do you have an organizational chart that clearly establishes direct responsibility for the quality system? Please attach a copy	Yes 🗌	N/A 🗌	No 🗌		
3.	Does management routinely review the quality system to ensure effectiveness of the system?	Yes 🗌	N/A 🗌	No 🗌		
4.	Is management's review of the system documented?	Yes 🗌	N/A 🗌	No 🗌		
5.	Do you perform audits internally to review processes and procedures?	Yes 🗌	N/A 🗌	No 🗌		
	2.3 Contract Review					
1.	Does your Quality system have the capability to determine whether you can meet customer requirements prior to accepting an order?	Yes 🗌	N/A 🗌	No 🗌		
2.	Do you have written contracts that clearly spell out the requirements of your customers?	Yes 🗌	N/A 🗌	No 🗌		
3.	Do you have written procedures for handling amendments to contracts?	Yes 🗌	N/A 🗌	No 🗌		
4.	What is your process or procedure to notify customers and report risks or concerns?	Yes 🗌	N/A 🗌	No 🗌		
2.4 Design Control						
1.	Do you have procedures in place to address design control?	Yes 🗌	N/A 🗌	No 🗌		
2.	Do your procedures address design and development planning?	Yes 🗌	N/A 🗌	No 🗌		
3.	Do your procedures address design input and design output?	Yes 🗌	N/A 🗌	No 🗌		
4.	Do your procedures address design review and verification?	Yes 🗌	N/A 🗌	No 🗌		
5.	Do your procedures address design validation and design changes?	Yes 🗌	N/A 🗌	No 🗌		
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2.5 Document and Data Control

1.	Are there procedures for the creation, approval, and changes to documents and/or data?	Yes 🗌	N/A 🗌	No 🗌
2.	Are current copies of necessary documents accessible to your employees where needed?	Yes 🗌	N/A 🗌	No 🗌
3.	Is there a procedure for collecting and destroying obsolete documents?	Yes 🗌	N/A	No 🗌
	2.6 Purchasing			
1.	Do you have a list of approved suppliers?	Yes 🗌	N/A 🗌	No 🗌
2.	Do you review and approve purchasing documents for accuracy of specified requirements before release?	Yes 🗌	N/A 🗌	No 🗌
3.	Do you have a formal corrective action system for your suppliers?	Yes 🗌	N/A 🗌	No 🗌
	2.7 Customer Supplied Products			
1.	Do you have documented procedures for the control of verification, storage and maintenance of customer-supplied products?	Yes 🗌	N/A 🗌	No 🗌
2.	Are customer-supplied products that are lost, damaged, or otherwise unsuitable for use recorded and reported to the customer?	Yes 🗌	N/A 🗌	No 🗌
3.	Are retention samples approved and stored?	Yes 🗌	N/A 🗌	No 🗌
2.8 Product Identification and Traceability				
1.	Does each batch or lot have its own unique identification?	Yes 🗌	N/A 🗌	No 🗌
2.	Is the batch identification maintained throughout its production and delivery?	Yes 🗌	N/A 🗌	No 🗌
3.	Can the product be traced back from any point of manufacture or delivery?	Yes 🗌	N/A 🗌	No 🗌
2.9 Process Control				
1.	Do front line personnel do their jobs by following documented work instructions?	Yes 🗌	N/A 🗌	No 🗌
2.	Is there a system in place to review and approve proposed process changes?	Yes 🗌	N/A 🗌	No 🗌
3.	Are preventative maintenance practices and principles employed?	Yes 🗌	N/A 🗌	No 🗌

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2.10 Inspection and Testing

1.	Do you have a Receiving Inspection/Testing system that assures that incoming product is not used until it is verified as meeting specified requirements?	Yes 🗌	N/A 🗌	No 🗌
2.	Do you have a system to ensure that final product is not released until it is verified as meeting Final Inspection/Testing requirements?	Yes 🗌	N/A 🗌	No 🗌
3.	Do you document the results of inspection and testing?	Yes 🗌	N/A 🗌	No 🗌
4.	Do you have a system to identify and separate non-conforming/suspect material?	Yes 🗌	N/A	No 🗌
	2.11 Inspection, Measuring and Test Equipment			
1.	Do you uniquely identify all of your inspection, measuring, and test equipment that can affect product quality?	Yes 🗌	N/A	No 🗌
2.	Do you have procedures that detail how and when your measuring and test equipment calibrations are to be done?	Yes 🗌	N/A 🗌	No 🗌
3.	Can the standards used to calibrate your measuring and test equipment be traced to N.I.S.T. or other nationally recognized primary standards?	Yes 🗌	N/A 🗌	No 🗌
4	Do you have a method to identify the calibration status of measuring and test equipment?	Yes 🗌	N/A 🗌	No 🗌
2.12 Product Inspection and Test Status				
1.	Is the identification and test status of products identified by suitable means that indicate conformance or non-conformance?	Yes 🗌	N/A	No 🗌
2.	Do the Inspection and Test Records identify the inspection authority for the release of conforming product?	Yes 🗌	N/A 🗌	No 🗌
2.13 Non-Conforming Materials				
1.	Do you have documented procedures that outline the review and disposition of non- conforming materials?	Yes 🗌	N/A 🗌	No 🗌
2.	Do you re-inspect product that has been reworked or repaired?	Yes 🗌	N/A 🗌	No 🗌
3.	Do you advise the customer when reworking or repairing nonconforming product?	Yes 🗌	N/A	No 🗌

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	2.14 Corrective/Preventive Action			
1.	Do you have documented procedures for the handling of customer complaints?	Yes 🗌	N/A 🗌	No 🗌
2.	Do you have documented procedures for handling Corrective/Preventive Action?	Yes 🗌	N/A 🗌	No 🗌
3.	Do you use recognized problem solving methods to identify the root cause of problems?	Yes 🗌	N/A	No 🗌
	2.15 Handling, Storage, Packaging, Preservation and Deliver	У		
1.	Do you have documented procedures for handling, storage, packaging, preservation and delivery of product?	Yes 🗌	N/A	No 🗌
2.	Are you delivering at least 99% on time to your customers?	Yes 🗌	N/A 🗌	No 🗌
3.	Do you have a process to ensure that all required, correct, associated documents are provided with ordered product as requested on our purchase order?	Yes 🗌	N/A	No 🗌
	2.16 Quality Records			
1.	Are your product records for quality identified, readily retrievable and legible?	Yes 🗌	N/A 🗌	No 🗌
2.	Do you have established times for retention of quality records?	Yes 🗌	N/A 🗌	No 🗌
	2.17 Internal Quality Audits			
1.	Do you have an established and documented internal audit system to verify quality activities?	Yes 🗌	N/A 🗌	No 🗌
2.	Do you carry out audits as scheduled?	Yes 🗌	N/A 🗌	No 🗌
3.	Are records of audits maintained and reviewed by responsible management personnel?	Yes 🗌	N/A 🗌	No 🗌
2.18 Training				
1.	Have the personnel, whose work can directly affect product quality, received all the training necessary in accordance with your documented procedures?	Yes 🗌	N/A	No 🗌
2.	Do you consider training a strategic issue and is training periodically evaluated?	Yes 🗌	N/A 🗌	No 🗌
	2.19 Servicing			
1.	For organizations that provide for servicing, do your procedures document how servicing will meet the specified requirements of the customer?	Yes 🗌	N/A 🗌	No 🗌

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2.20 Statistical Techniques 4.20 Statistical Technintechnintechniques 4.20 Statistical Techniques 4.20 Statis

Thank you for the time and effort spent on completing this survey.

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