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## **SUPPLIER EVALUATION QUESTIONNAIRE**

## **GENERAL INFORMATION**

Company Name:	Contact Name:			Title:	Title:				
Email:	nail: Phone #:				Fax #:			DUN #:	
Address: City:			State:				Zip Code:		
Are there other locations? Yes  No If yes, please lists them on a separate sheet.									
Is there any intention to from an off shore locati	-	-		vices to	es to II-VI If yes, please indicate whic			te which location.	
			KEY	PERSC	ONNEI	L			
President: General Ma			er: Purchasing:				Sales Contact:		
QA Manager:	nager's e	email address:				QA Manager Reports to:			
Tech support available   References (customers II-VI A&D can contact):   Yes   No									
REASON FOR SURVEY (Please check appropriate box):									
New Supplier: New Product:			Certification: Re-evaluation			luation:	Other:		
FACILITIES									
Total Number of Employees: Ma			cturing: Quality: Er			ngineering:		Administrative:	
Year Established:	MANUFACTURING/SERVICES PROVIDED to II-VI:					II-VI:			
сом	IPANY Q	UALITY S	YSTEM I				CORDAN	ICE W	тн
MIL-I-45208A	9001 🗌	AS910	0 🗌 📗	ISO 134	l85 🗌	Other	:		ITAR Registered? Yes ☐ No ☐



BUSINESS CLASSIFICATION (please check appropriate box):										
Small Business	Small Business HubZone Women Owned Veteran Owned Business Small Disadvantaged Business					Large Business				
PROVIDE A	IF YOUR COMPANY IS CERTIFIED TO ANY THE QMS STANDARD (ISO 9001, AS9100 or ISO 13485), STOP HERE AND PROVIDE A COPY OF YOUR CERTIFICATE.  IF NO, CONTINUE, AND COMPLETE QMS REQUIREMENTS LISTED BELOW.									
Signature of Person that Completed this Survey:				Title:		Date:				

	QMS Requirements	Yes	No	N/A	Comments
Ш	Quality Management System				
a	Do you have a quality management system, establish, document, implement, and maintain in accordance with quality standards?				
b	Do you have a Quality Assurance Manual?				
С	Do the employees have access to quality management system documentation and are aware of relevant procedures?				
Ш	Management Responsibility				
а	Do you have documented statements of quality policy and quality objectives?				
b	Does the management regularly review quality performance data?				
С	Is the management review output documented and maintained?				
IV	Control of Documents				
a	Is there a procedure describing control of documents?				
b	Are the obsolete drawings, specifications, and procedures removed from the locations where they might be mistakenly used?				
С	Do you have a system of controlling incoming documentation changes?				
d	Are the latest drawings, specifications and procedures available at point of use?				
٧	Control of Records				

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	QMS Requirements	Yes	No	N/A	Comments
а	Is there a procedure describing control of records of quality management system such as work order router, receiving inspection data, process inspection data, test results, calibration records, audit reports, training records, etc.?				
b	Are the records of quality activities maintained per procedure?				
VI	Identification and Traceability				
а	Is identification and traceability maintained throughout manufacturing processing?				
VII	Purchasing				
а	Do you have a procedure for supplier selection?				
b	Do your purchasing documents describe all requirements, inspection, testing, and packaging?				
С	Does the quality assurance review the purchase orders for the inclusion of applicable requirements?				
d	Do you periodically review and evaluate your subcontractors?				
VIII	Customer Property				
а	Do you have a system to ensure proper identification, storage, and segregation of customer supplied product?				
IX	Control of Monitoring and Measuring Devices				
а	Do you have a calibration procedure?				
b	Are calibration records maintained for measuring and test equipment?				
С	Are measuring and test equipment not calibrated identified and stored to preclude their use pending calibration?				
Х	Internal Quality Audit				
а	Do you have internal audit procedures?				
b	Do you maintain the results of internal audits?				
С	Is corrective action taken on nonconformance?				
XI	Process Control				
а	Do manufacturing process sheets exist, identifying operations, tests, and inspections?				
b	Is suitable equipment maintenance being performed to ensure continuing process capability?				
С	What are the requirements of the company to prevent Foreign Object Damage (FOD)?				
XII	Control of Nonconforming Product				

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	QMS Requirements	Yes	No	N/A	Comments
а	Is there any procedure for control of nonconforming material?				
b	Are the records of nonconforming material kept?				
С	Are all nonconforming material identified and segregated in assigned holding areas?				
XIII	Corrective Action				
а	Is there any corrective action procedure?				
b	Is corrective action system documented?				
С	Is follow-up taken and documented concerning effectiveness of corrective action?				
XIV	Handling, Storage, and Shipping				
а	Are there written procedures for handling, storage, and packaging?				
b	Do you verify final packaging and marking prior to shipping?				
С	Do you have the methods of proper handling to prevent damage or deterioration in stores?				
d	Do all materials stored have evidence of inspection?				

## APPROVAL STATUS (to be filled out by II-VI A&D)

Approved? Yes No (May require two sig	(May require two signatures by II-VI A&D at the discretion of QMS Manager)				
Signature:	Title:	Date:			
Signature:	Title:	Date:			
Comments:					

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