

he purpose of this questionnaire is to ensure consistent evaluation of a supplier's capabilities and to determine the caliber of the Quality Program in place. Because of this, as a supplier or potential supplier to Teledyne Controls, your prompt response to the following questionnaire is critical.

Please answer as completely as possible. Where the question does not apply to your company, check-off the box identified N/A. Thank you for your cooperation.

## TELEDYNE CONTROLS SUPPLIER DEVELOPMENT

Supplier Code:

## 1. GENERAL INFORMATION

	Name of Company: Street Address:		Telephone Fax:		
	Mailing Address:		City:		
	State:		Zip Code:		
2.	TYPE OF BUSINESS				
		ssembly Shop 🔲 Distributor		] Special Process 🛛 Maintenance	
3.	If applicable, list the name of y	our parent company or any subsidiary	below:		
	Subsidiary:			Principal Owner	
4.	When was your company established:				
-		y ear/month			
5.		Id manufacture or repair product proc			
6. 7		Indicate the approximate square footage of each Office: Warehouse: Manufacturing: What type of measuring units are used by your company: Metric 🗌 Imperial 🗌 Both 🗌			
7. 8.	Total number of employees:		Imperial 🗌 🛛 Both 🗌	J	
•	Administration:	Engineering:	Production:	QA/QC:	
9.	Key Management Positions:	(please attach a copy of your org			
	President/General Manager:		-		
	Vice President:				
	QA/QC Manager:				
	Engineering Manager:				
	Production:				
10.	Does your facility shut down for If yes, please state normal vac			Yes 🗌 No 🗌	
11.	List the principal products or se Product/Services:	ervices sold assembled or distributed b	by your company:	% ofBusiness	
12.	List major customers, including	GovernmentAgencies, for which you	r company has or is currently per	forming work:	

\_\_\_% Military



## Supplier Evaluation Questionnaire

% Commercial

13.		es your company have an established Quality Program? es, which standard(s) does your program conform to:	Yes 🗌	No 🗆	N/A 🗆
		AS9100C 🗌 ISO 9001:2008 🗌 MIL-I-45208A	□ FAR 14	5	
		AS9100B 🗌 ISO 9003 🗌 MIL-Q9858A	□ FAR 21		
		AS9120A 🗌 ISO 9002 🔲 FAA-STD-016			
		NADCAP List			
		v long has your Quality Program been established? Years	_	_	
	lf ye	s your Quality Program ever been audited and approved by any of your customers or an accredited registrar? es, by whom?	Yes 🗌	No 🗆	N/A 🗆
16.	Doy	you have an FAA approved Drug& Alcohol Program at your facility?	Yes 🗌	No 🗖	N/A 🗆
17.		es your sub-tier suppliers' conducting safety sensitive work for the repair stations have an FAA approved Drug Icohol Program?	g Yes 🗌	No 🗆	N/A 🗆
18.	Plea	ase provide a copy of your current FAA and or EASA Repair Station Certificate, Ops Spec and evidence of an	active Drug a	and Alcoho	Program:
	Plar	Number: Comments:			
19.	CO	UNTERFEIT PART AVOIDANCE:			
		es your company comply with the following standards to ensure counterfeit part avoidance?			
		ndard: 2021 (Counterfait Bart Aveidance)		Yes 🗌	No 🗆
		3081 (Counterfeit Part Avoidance) 3553 (Counterfeit Part Avoidance- Electronics)		Yes 🗆	
		6174 (Counterfeit Part Avoidance - Materials)		Yes	
		12931 (Authentication standards for brand owners, manufacturers, and packagers)		Yes 🗆	
		thers: Specify:		Yes 🗆	
	A.	Does your organization have documented procedures/plans addressing the avoidance of counterfeit parts?		Yes 🗌	No 🗆
	л. В.	Does your organization have documented procedures for the containment of suspect / confirmed counterf	eit narts and	Yes	
	_	customer notification?			
	C.	Does your organization provide counterfeit part avoidance training to employees?		Yes 🗌	No 🗆
	D.	Does your organization procure Electrical, Electronics, and Electromechanical (EEE) parts, electronic parts, and equipment only through Original Equipment Manufacturers (OEMs) or an authorized distributor ever from an unauthorized supplier achieves costsavings?		Yes 🗌	No 🗆
	E.	Does your organization flow counterfeit part avoidance and other purchasing requirements through the supp	lier chains?	Yes 🗆	No 🗖
	F.	Does your organization notify Customers when counterfeit parts are detected?		Yes 🗆	No 🗆
	G.	Does your organization maintain a method of item traceability that ensures tracking of the supply chain	back to the	Yes 🗆	No 🗆
		original manufacturer of all Electrical, Electronics, and Electromechanical (EEE) components and devices assemblies and subassemblies being delivered?			
	H.	Does your organization approve and retain part Certificates of Conformance (C of C) for all Electrical, Electromechanical (EEE) components and devices included in assemblies and subassemblies being deliver		Yes 🗌	No 🗆
	I.	Will Certificates of Conformance (C of C) for all Electrical, Electronic, and Electromechanical (EEE) com devices be available upon Teledyne Controls request?	ponents and	Yes 🗆	No 🗆
	J.	Does your organization retain manufacturer's certifications per the record retention requirements dir Teledyne Controls Purchase Orders?	ected within	Yes 🗆	No 🗆
	K.	Does your organization utilize incoming inspection and test methods to assure the detection of potential cou and materials?	nterfeit parts	Yes 🗌	No 🗆



Supplier Evaluation Questionnaire

	IF YOUR COMPANY HAS A 3 <sup>RD</sup> PARTY QUALITY MANAGEMENT SYSTEM CERTIFICATION ACCREDITED BODY (SUBMIT COPY OF CERTIFCATION), YOU DO NOT NEED TO COMP BELOW. YOUR SIGNATURE IS REQUIRED ON PAGE 5 (ELECTRONIC SIGNATURE PERMIS	LETE QUE		
20.	Which of the following standards is your company compliant with:			
	□       MIL-STD-785       □       MIL-STD-701       □       MIL-STD-2000       □         □       IPC-610       □       MIL-P-50884       □       MIL-P-55110       □         □       MIL-STD-105       □       MIL-STD-1535       □       AS5553       □         □       ANSI/ASQC Z1.4 & Z1.9       □       MIL-M-55565       □       RS-471       □	AS6174 ISO100 IPC-A-6 IPC-A-6 IPC-A-6 J-STD-0 J-STD-0	12 500 510 VHMA-620 001	
21.	Do any of your major customers perform source inspection at your company?	Yes 🗌	No 🗌	N/A 🗆
22.	How many inspectors does your company employ?			
23.	Does your organization have a Quality Program Manual?	Yes 🗌	No 🗆	N/A 🗆
	If yes, what is the date of issue and current revision level? Date: Revision Level:			
	(Please send copy of your Quality Manual with this questionnaire)			
24.	Does your Quality Program Manual have written procedures for the following:			
		lentification & n, Measuring e Action		
25.	Does your company maintain a program to capture quality costs, such as prevention, detection and failure?	Yes 🗆	No 🗆	N/A 🗆
	Does your company review all contracts (including purchase orders)?	Yes 🗆	No 🗆	N/A 🗆
	Are documents such as drawings, specifications, procedures and QA forms controlled?	Yes 🗆	No 🗆	N/A 🗆
28.	Are company policies and procedures periodically reviewed and revised?	Yes 🗆	No 🗆	N/A 🗆
29.	Are records of changes documented and maintained?	Yes 🗆	No 🗆	N/A 🗆
30.	Are customer document changes processed in a controlled manner?	Yes 🗆	No 🗆	N/A 🗆
31.	Does QA/QC approve company suppliers/subcontractors?	Yes 🗆	No 🗌	N/A 🗆
32.	Is a list maintained of approved suppliers/subcontractors?	Yes 🗆	No 🗆	N/A 🗆
33.	Are company purchase orders reviewed by QA/QC prior to release?	Yes 🗆	No 🗆	N/A 🗆
34.	Are QA requirements (Quality Clauses) included within the purchase order?	Yes 🗆	No 🗆	N/A 🗆
35.	Are only approved suppliers/subcontractors used?	Yes 🗆	No 🗆	N/A 🗆
	When applicable, are source inspections performed at supplier/subcontractor facilities?	Yes 🗖	No 🗖	N/A 🗆
37.	When applicable, are certifications and/or test reports requested to accompany shipments from suppliers/subcontractors?	Yes 🗆	No 🗆	N/A 🗆
38.	Does your company ever receive customer supplied materials?	Yes 🗆	No 🗌	N/A 🗌
39.	Are procedures in place for the storage and maintenance of customer supplied material?	Yes 🗆	No 🗆	N/A 🗆
40.	Is customer supplied material identified and segregated?	Yes 🗆	No 🗆	N/A 🗆
41.	Are shop orders, travelers, work instructions, etc. used during manufacturing?	Yes 🗆	No 🗆	N/A 🗆
42.	Does your company maintain a system for material and partidentification?	Yes 🗆	No 🗆	N/A 🗆
43.	When applicable, does your company maintain raw material and component traceability? List all special processes performed by your company such as welding, plating, sandblasting, painting, soldering, e	Yes 🗆 etc.	No 🗆	N/A 🗆
лл	Does your company maintain records of personnel training/certification for special processes?		No	

44.	Does your company maintain records of personnel training/certification for special processes?	Yes 🗌	No 🗆	N/A 🗆
45.	Are written procedures available and used by processing and inspection personnel?	Yes 🗌	No 🗖	N/A 🗆
	TCF 1083 (Rev. J 9/10//15)			



46.	Does your company monitor special processes performed by subcontractors?	Yes 🗆	No 🗖	N/A 🗆
	Are all incoming parts, raw materials and assemblies inspected?	Yes 🗆	No 🗆	N/A 🗆
	Is rejected material tagged, segregated and bonded?	Yes 🗆	No 🗖	N/A 🗆
	Does your company perform in-process inspection?	Yes 🗆	No 🗆	N/A 🗆
	Are final acceptance inspection and tests performed on completed items prior to submittal to customer?	Yes 🗆	No 🗆	N/A 🗆
	Are first article inspections performed? If yes, please specify: Company Form AS9102A	Yes 🗆	No 🗆	N/A 🗆
52.		Yes 🗆	No 🗖	N/A 🗆
53.	Are written workmanship standards maintained?	Yes 🗆	No 🗆	N/A 🗆
	Does your company maintain a system for periodic calibration of inspection, test, measuring and	Yes 🗆	No 🗆	N/A 🗆
	manufacturing equipment? If yes, what standard does this system conform to:			
55.	Does your company perform their own calibration in-house? If yes, please list below each inspection device calibrated in-house and the specification used to calibrate that specific inspection device:	Yes 🗌	No 🗖	N/A 🗆
56.	Is calibration performed on a scheduled basis?	Yes 🗆	No 🗆	N/A 🗆
57.	Are written procedures used for calibrating equipment?	Yes 🗆	No 🗖	N/A 🗆
58.	Are calibration standards traceable to nationally recognized standards?	Yes 🗆	No 🗆	N/A 🗆
59.	Is equipment identified as to due date for next calibration?	Yes 🗆	No 🗆	N/A 🗆
60.	Are calibration frequencies established for each type of equipment?	Yes 🗆	No 🗖	N/A 🗆
61.	Is there a system to calibration recall?	Yes 🗆	No 🗆	N/A 🗆
62.	Does the calibration system include employee's personal tools?	Yes 🗆	No 🗖	N/A 🗆
63.	Is subcontracted calibration performed by qualified sources?	Yes 🗆	No 🗆	N/A 🗆
64.	Is a system maintained for identifying inspection status throughout all operations?	Yes 🗆	No 🗖	N/A 🗆
65.	Are identification tags, stickers, etc. used to indicate inspection status? If yes, specify:	Yes 🗆	No 🗆	N/A 🗆
66.	Are inspection stamps used? If no, do you maintain an Initial log? Yes 🗌 No 🗔	Yes 🗆	No 🗖	N/A 🗆
67.	Are nonconforming items identified and segregated?	Yes 🗆	No 🗆	N/A 🗆
68.	Who dispositions nonconforming items?	Yes 🗆	No 🗖	N/A 🗆
69.	Are dispositions documented?	Yes 🗆	No 🗖	N/A 🗆
70.	Are records maintained of all non-conformances?	Yes 🗆	No 🗖	N/A 🗆
71.	Is material dispositioned for rework or repair subject to re-inspection?	Yes 🗆	No 🗖	N/A 🗆
72.	Are repetitive nonconformance's identified and analyzed?	Yes 🗆	No 🗖	N/A 🗆
73.	Is there a formal corrective action program?	Yes 🗆	No 🗖	N/A 🗆
74.	Who initiates corrective action?	Yes 🗆	No 🗖	N/A 🗆
75.	Does your company have a Material Review Board?	Yes 🗆	No 🗖	N/A 🗆
76.	Are MRB dispositions fully documented and signed by authorized members?	Yes 🗆	No 🗖	N/A 🗆
77.	Is there a system for processing customer rejects?	Yes 🗆	No 🗆	N/A 🗆
78.	Does your company have a formal failure analysis program?	Yes 🗆	No 🗖	N/A 🗆
79.	What are your packaging capabilities?  Military  Commercial  Other: Specify:			
80.	Is there a system of control for storing products having a defined shelf life?	Yes 🗆	No 🗆	N/A 🗆
81.	Are audits conducted and documented of storage areas at established intervals?	Yes 🗆	No 🗆	N/A 🗆
82.	Does your company have maintenance and repair program for equipment that affects product conformity?	Yes 🗆	No 🗆	N/A 🗆
83.	Are inspections performed during packaging and prior to shipping?	Yes 🗌	No 🗖	N/A 🗆
84.	Does your organization maintain all inspection records generated?	Yes 🗆	No 🗆	N/A 🗆
	If yes, how long are these records retained on file?	Yes 🗆	No 🗖	N/A 🗆
85.	Are quality records analyzed by management to assess quality program effectiveness?	Yes 🗆	No 🗆	N/A 🗆
86.	Does your organization maintain a quality audit program? Internal:	Yes 🗆	No 🗖	N/A 🗆



## Supplier Evaluation Questionnaire

		External:	Yes 🗌	No 🗖	N/A 🗖
87.	Is an annual audit schedule maintained?		Yes 🗌	No 🗆	N/A 🗆
88.	Are all quality audits documented?		Yes 🗌	No 🗖	N/A 🗆
89.	Does your company have a Total Quality Management/Quality Improvement Program?		Yes 🗌	No 🗆	N/A 🗆
90.	Does your company employ SQC and/or SPC techniques?		Yes 🗌	No 🗆	N/A 🗆
91.	Is inspection performed by sampling?		Yes 🗌	No 🗆	N/A 🗆
	If yes, what sampling standards used:		Yes 🗌	No 🗆	N/A 🗆
92.	Are ESDS parts handled at your facility?		Yes 🗆	No 🗆	N/A 🗆
93.	Is ESD protective clothing used?		Yes 🗌	No 🗆	N/A 🗆
94.	Are all ESDS parts handled at ESD work stations?		Yes 🗌	No 🗆	N/A 🗆
95.	Are ionizers used?		Yes 🗌	No 🗆	N/A 🗆
96.	Do written procedures exist for the storage, handling and packaging of ESDS parts?		Yes 🗌	No 🗆	N/A 🗆
97.	Are good housekeeping practices followed?		Yes 🗌	No 🗖	N/A 🗆
	The above statements are certified to be true:				

Name

Title

Date