### A-4 PRODUCT/PROCESS QUALITY CHECKLIST

Customer or Internal Part No.

Revision Level

	Customer of internal Lart No.		,	Hevision Level							
	Question	Yes	No	N/A	Comment / Action Required	Person Responsible	Due Date				
	Is customer assistance or approval required										
1	for the development of the control plan?										
	Has the organization identified who will be the										
2	quality liaison with the customer?										
	Has the organization identified who will be the										
3	quality liaison with its suppliers?										
	Has the quality management system been										
	reviewed and approved per customer specific										
4	requirements?										
	Are there sufficient personnel identified to										
5	cover:										
а	Control plan requirements?										
b	Layout inspection?										
С	Engineering performance testing?										
d	<ul> <li>Problem reaction and resolution analysis?</li> </ul>										
6	Is there a documented training program that:										
а	Includes all employees?										
b	Lists whose been trained?										
С	Provides a training schedule?										
7	Has training been completed for:										
а	Statistical process control?										
b	Capability studies?										
С	Problem solving?										
d	Mistake proofing?										
е	Reaction Plans?										
f	Other topics as identified?										
	Is each operation provided with process										
8	instructions that are keyed to the control plan?										
	Are standard operator instructions accessible										
9	at each work station?										

	IDs an austru in atmosticus in alcula mietoma and	 1		
1,,	Do operator instructions include pictures and			
10	diagrams?			
	Were operator/team leaders involved in			
	developing standard operator instructions?			
12	Do inspection instructions include:			
	Easily understood engineering performance			
а	specifications?			
b	Test frequencies?			
С	Sample sizes?			
d	Reaction plans?			
е	Documentation requirements?			
13	Are visual aids:			
а	<ul> <li>Appropriate, easily understood and legible?</li> </ul>			
b	Available?			
С	Accessible?			
d	Approved?			
е	Dated and current?			
	Is there a procedure to implement, maintain,			
	and establish reaction plans, for issues such			
	as out of control conditions based on statistical			
14	process control?			
	Is there an identified problem solving process			
15	that includes root cause analysis?			
	Are the latest drawings and specification			
	available for the operator, in particular at the			
16	points of the inspection?			
	Have engineering tests (dimensional,			
	material, appearance, and performance) been			
	completed and documented as required in			
la	accordance with customer requirements?			
Ĕ	Are the current forms/logs available for	$\dashv$		
	appropriate personnel to record inspection			
17	results?			
<u> </u>	Are the following available and placed at the		+	
18	appropriate points of the operation?			
0	appropriate points of the operation.		l l	

SOP-102-SH-CR4.3-169 Effective date: 09/01/14

	L. Manifestan and accommunity design of	 1		1	<u> </u>
a	Monitoring and measurement devices?				
b	Gage instructions?				
С	Reference samples?				
d					
	Have provisions been made to certify and				
	calibrate gages and test equipment at a				
19	defined frequency that is appropriate?				
	Have required measurement system capability				
20	studies been:				
а	Completed?				
b	Accepted?				
	have initial process capability studies been				
21	conducted per customer requirements?				
	Are layout inspection equipment and facilities				
	adequate to provide initial and ongoing layout				
	of all details and components in accordance				
22	with customer requirements?				
	Is there a documented procedure for				
	controlling incoming material that may include,				
23	for example, the following items:				
а	Characteristics to be inspected?				
b	Frequency of inspection?				
С	Samplesize?				
d	Designated location for approved product?			 	
е	Disposition of nonconforming products?				
	Have sample production parts been provided		 		
24	per customer requirements?				
	Is there a procedure to identify, segregate, and				
	control nonconforming products to prevent				
25	shipment?				
	Are rework/repair procedures available to				
26	assure conforming product?				
	Is there a procedure to requalify				
27	repaired/reworked material?				
				•	

SOP-102-SH-CR4.3-169 Effective date: 09/01/14

	Has a master sample, if required, been retained as part of the part approval process?					
	Is there an appropriate lot traceability procedure?					
	Are periodic audits of outgoing products planned and implemented?					
31	Are periodic assessments of the quality system planned and implemented?					
	Has the customer approved the packaging and the packaging specification?					



Revision Date	
_	
Prepared by:	