

Process/Product Audit Checklist

Cust #: <u>10</u>		Customer: <u>HEIL FAB</u>		GRP/Size/GRD/Width: <u>HRS 19E/11530/19.44 x 159.07/1525</u>	
PWC: <u>GPI</u>		W/O#: <u>155898</u>		Date: <u>5/13/22</u> Part #(s): <u>137-0670</u> Auditor: <u>C. Martinez</u>	
Gauge Range: <u>.1520-.1620</u>		Actual Gauge: <u>.152</u>		Width Range: <u>19.31-19.50</u> Width Actual: <u>19.44</u>	
Length Range: <u>159.01-159.11</u>		Length Actual: <u>159.070</u>		Other: <u>1.720-1.780</u> Other Actual: <u>1.750</u>	
Other: <u>N/A</u>		Other Actual: <u>N/A</u>		Other: <u>N/A</u> Other Actual: <u>N/A</u>	
Item	YES	NO	N/A	Comments/Action Taken (Required for NO)	
Process Inspection Sheets filled out according to frequency and sampling required?	✓				
Correct raw material type and size?	✓			Tag(s) to use: <u>300 710</u> Tag(s) used: <u>300 710</u>	
Setup performed according to W/O?	✓				
Product is acceptable according to customer-specific requirements? [Fab: Is the Part Print Present & the correct Revision? Are required measurements documented?]	✓			[Fab: Print Rev: <u>0</u> , W/O Rev: <u>0</u> , Part Spec Rev: <u>0</u>] (Leave blank if non-Fab audit)	
Packaging is acceptable according to customer-specific requirements?	✓				
Visual Inspection performed and product meets requirements?	✓				
Out of spec noted, with actions taken?		✓			
Non-conforming material put into reject warehouse and physically put into non-conforming area?		✓			
Required gages available & functional?	✓				
All Gages Calibrated (List in Comments)	✓			Gages Observed (list last calibration and when due) <u>1205 2/4/21 Due 2/4/23</u> <u>2094 1/18/21 Due 1/18/23</u>	
Housekeeping: Machine/Floor clean? Loose tags & paperwork cleaned up?	✓				
Required PPE being worn?	✓				
Forms are the latest revision per Quality Intranet?	✓			List Forms (Observed Rev vs Intranet Rev) <u>Rev 0 vs Rev 0</u>	
Hardcopy Controlled Documents are listed on Quality Intranet by location?	✓			List Documents and their Location: <u>Brook Room, Plasma, Front Office</u>	