

Process/Product Audit Checklist

Cust #: <u>3418</u> Customer: <u>Freight Car</u> GRP/Size/GRD/Width: <u>HRS/312/A57260/42.5</u>				
PWC: <u>LL1</u> W/O#: <u>10053</u> Date: <u>3/18/19</u> Part #(s): <u>M112812-000</u> Auditor: <u>Z. Delp</u>				
Gauge Range: <u>.3025 - .3225</u> Actual Gauge: <u>.310</u> Width Range: <u>42.5⁺ - .06³</u> Width Actual: <u>42.5</u>				
Length Range: <u>107.5⁺ - .06³</u> Length Actual: <u>107.468"</u> Other: <u>/</u> Other Actual: <u>/</u>				
Other: <u>/</u> Other Actual: <u>/</u> Other: <u>/</u> Other Actual: <u>/</u>				
Item	YES	NO	N/A	Comments/Action Taken (Required for NO)
Process Inspection Sheets filled out according to <u>frequency</u> and <u>sampling</u> required?	✓			
Correct raw material type and size?	✓			Tag(s) to use: <u>20654</u> Tag(s) used: <u>20654</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>/</u> , W/O Rev: <u>/</u> , Part Spec Rev: <u>/</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) 050 Mils 11/18 due 5/19 046 Tape 11/18 due 5/19 16/156625 Calipers 11/18 due 5/19
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) Gad 059 obs. Rev. 0 intranet Rev. 0
Hardcopy Controlled Documents are listed on Quality Intranet by location?			✓	List Documents and their Location: No controlled docs @ LL1