

Audit Report

FSNS Beef Trim CCP Addendum

Caviness Beef Packers - Hereford 3255 U.S. 60

Hereford, Texas 79045

Audit Date: November 11, 2022 Auditor: Scott Devitt



Audit Summary

Company Name:	Caviness Beef Packers - Hereford	Company ID:	AUCAVHER
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Auditor Name:	Scott Devitt
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Definitions for the purpose of this Addendum:

Validation - Data that demonstrates there is a pathogen kill when an intervention is operating within specified parameters. Verification - Demonstration of a microbiological reduction by an intervention when operating in validated parameter(s). Monitoring - Checking / reading of intervention parameters / measurements (ex. Temperature, concentration, etc.).

PLEASE NOTE: A "NO" answer does not necessarily represent a deficiency in a facility's programs or processes.



Beef Trim - CCP Addendum

1 HACCP

		Result
1.1	Adequacy of the HACCP plan is reassessed by the establishment on an annual basis or whenever changes occur that could affect the hazard analysis or alter the HACCP plan. Review the establishment's HACCP reassessment log to identify the last reassessment.	Yes
Comment:	HACCP plans were reassessed annually at a minimum or as required for process changes. The most recent reassessment occurred on 11/7/22.	
1.2	The establishment maintains records to demonstrate that responsible personnel have been trained in monitoring activities as described in their HACCP plan.	Yes
Comment:	CCP monitors received training annually and as needed; training records reviewed for 2022 evidenced program compliance.	
1.3	The establishment maintains records that confirm corrective actions are taken when there is a deviation from a critical limit.	Yes
Comment:	Corrective actions for CCP deviations met requirements of 9 CFR 417.3(a). Corrective actions from CCP zero tolerance deviations from 10/14/22 and 8/19/22 were provided and met regulatory requirements.	

2 Interventions/Process Aids - Steam Vacuum

		Result
2.1	The establishment uses the steam vacuum intervention method.	No
Comment:	Steam vacuums were not used.	
2.2	The establishment identified this intervention as a CCP.	Not Applicable
Comment:	Steam vacuums were not used.	
2.3	If the Steam Vacuum is a CCP, can the line run if this intervention is not operational or not in specification.	Not Applicable
Comment:	Steam vacuums were not used.	
2.4.1	None	Not Applicable
Comment:	Steam vacuums were not used.	
2.4.2	Validated Third Party Challenge Study or Validation Study	Not Applicable
Comment:	Steam vacuums were not used.	
2.4.3	In-house Challenge Study or Validation Study	Not Applicable
Comment:	Steam vacuums were not used.	
2.4.4	Third Party review of in-house challenge study or validation. List the name of the Third Party in Comments.	Not Applicable



Comment:	Steam vacuums were not used.	
2.4.5	Resource white paper (Published Journal Article)	Not Applicable
Comment:	Steam vacuums were not used.	
2.4.6	Resource white paper with third party review (peer reviewed paper - not published)	Not Applicable
Comment:	Steam vacuums were not used.	
2.4.7	Other List in comments	Not Applicable
Comment:	Steam vacuums were not used.	
2.5.1	A specific set of samples were chosen to support the validation hypothesis (objective).	Not Applicable
Comment:	Steam vacuums were not used.	
2.5.2	Statistical parameters were used in the validation hypothesis and/or the analysis to support the conclusion.	Not Applicable
Comment:	Steam vacuums were not used.	
2.5.3	Scientific support documentation.	Not Applicable
Comment:	Steam vacuums were not used.	
2.5.4	Validation study was prepared by a third party. List the name of the third party in comments.	Not Applicable
Comment:	Steam vacuums were not used.	
2.5.5	Other List in comments	Not Applicable
Comment:	Steam vacuums were not used.	
2.6	The establishment has records demonstrating on-going verification activities for this intervention. List the Frequency in comments.	Not Applicable
Comment:	Steam vacuums were not used.	
2.7.1	The establishment has documented procedures that include the following:	Not Applicable
	Operation of this intervention method	
Comment:	Steam vacuums were not used.	
2.7.2	Temperature monitoring	Not Applicable
Comment:	Steam vacuums were not used.	
2.7.3	Vacuum monitoring	Not Applicable
Comment:	Steam vacuums were not used.	
2.7.4	Steam pressure monitoring	Not Applicable
Comment:	Steam vacuums were not used.	
2.7.5	Removal of contamination (Must follow regulatory guidelines of 'less than one inch')	Not Applicable
Comment:	Steam vacuums were not used.	
2.7.6	Maintenance of the intervention equipment	Not Applicable



Comment:	Steam vacuums were not used.	
2.7.7	Observation of the intervention in operation	Not Applicable
Comment:	Steam vacuums were not used.	
2.7.8	None of the above.	Not Applicable
Comment:	Steam vacuums were not used.	
2.8	Operators of the steam vacuum(s) are following documented procedures as written for this intervention. If no, list findings in comments.	Not Applicable
Comment:	Steam vacuums were not used.	
2.9	The establishment's intervention operating parameters fall within the validation supporting documentation parameters	Not Applicable
Comment:	Steam vacuums were not used.	

3 Interventions/Process Aids - Thermal Intervention

		Result
3.1	The establishment uses the Thermal (hot water or steam pasteurization) intervention method.	Yes
Comment:	The site utilized 180F pre-evisceration and hot water pasteurization cabinets.	
3.2	The establishment identified this intervention as a CCP.	Yes
Comment:	The final pasteurization cabinet was identified as an either/or CCP with lactic acid.	
3.3	If the Thermal (hot water or steam pasteurization) intervention is a CCP, can the line run if this intervention is not operational or not in specification.	Yes
Comment:	The line could run without hot water pasteurization if lactic acid was operational. The line could not run without one of the interventions functioning properly.	
3.4.1	None	Not Applicable
3.4.2	Validated Third Party Challenge Study or Validation Study	No
3.4.3	In-house Challenge Study or Validation Study	Yes
Comment:	Hot Water Wash Cabinet Stand Alone Intervention Validation - 9/1/2021	
3.4.4	Third Party review of in-house challenge study or validation. List the name of the Third Party in Comments.	No
3.4.5	Resource white paper (Published Journal Article)	Yes



Comment: Effects of Steam-Vacuuming and Hot Water Spray Wash on the Microflora of Refrigerated Beef Carcass Surface Tissue Inoculated with *Escherichia coli* O157:H7, *Listeria innocula* and *Clostridium sporogenes* Dorsa 1997. Journal of Food Protection. Vol. 60, No. 2, Pages 114-119.

Use of Hot Water for Beef Carcass Decontamination. Journal of Food Protection. Vol. 60, Pages 19-25. Castillo 1998

Treatment Using Hot Water Instead of Lactic Acid to Reduce Levels of Aerobic Bacteria and Enterobacteriaceae and Reduce the Prevalence of *Escherichia coli* O157:H7 on Pre-evisceration Beef Carcasses. Journal of Food Protection. Vol. 69, No.8. Pages 1808-1803. Bosilevac 2006

Evaluation of Commonly Used Antimicrobial Interventions for Fresh Beef Innoculated with Shiga-Toxin Producing *Escherichia coli* Serotypes O26, O45, O103, O111, O121, O145, and O157:H7. Journal of Food Protection. Vol. 75, No. 7, 2012. Pages 1207-1212. Kalchayanand 2012

3.4.6	Resource white paper with third party review (peer reviewed paper - not published)	No
3.4.7	Other List in comments	No
3.5.1	A specific set of samples were chosen to support the validation hypothesis (objective).	Yes
Comment:	Specific sample sets were selected for the validation.	
3.5.2	Statistical parameters were used in the validation hypothesis and/or the analysis to support the conclusion.	Yes
Comment:	Log reduction of APC, coliforms, and generic E. coli supported the conclusion.	
3.5.3	Scientific support documentation.	Yes
Comment:	Microbiological testing data supported the conclusion.	
3.5.4	Validation study was prepared by a third party. List the name of the third party in comments.	No
Comment:	Validation study was prepared in-house.	
3.5.5	Other List in comments	Not Applicable
3.5.5 3.6	Other List in comments The establishment has records demonstrating on-going verification activities for this intervention. List the Frequency in comments.	Not Applicable Yes
	The establishment has records demonstrating on-going verification activities for this	
3.6	 The establishment has records demonstrating on-going verification activities for this intervention. List the Frequency in comments. One out of every 300 carcasses processed was swabbed for generic <i>E. coli</i> per regulatory requirements. Products intended for raw ground use were sampled and tested for <i>E. coli</i> O157:H7. Sampling and testing of finished products for non-O157 STEC, Salmonella, and indicator organisms including APC, coliforms, and generic <i>E. coli</i> was based on customer-specific requirements. Quarterly process validation swabs were collected on ten carcasses pre and post interventions; swabs were tested for APC, coliforms, and generic <i>E.</i> 	
3.6 Comment:	The establishment has records demonstrating on-going verification activities for this intervention. List the Frequency in comments. One out of every 300 carcasses processed was swabbed for generic <i>E. coli</i> per regulatory requirements. Products intended for raw ground use were sampled and tested for <i>E. coli</i> O157:H7. Sampling and testing of finished products for non-O157 STEC, Salmonella, and indicator organisms including APC, coliforms, and generic <i>E. coli</i> was based on customer-specific requirements. Quarterly process validation swabs were collected on ten carcasses pre and post interventions; swabs were tested for APC, coliforms, and generic <i>E. coli</i> .	Yes
3.6 Comment: 3.7.1	 The establishment has records demonstrating on-going verification activities for this intervention. List the Frequency in comments. One out of every 300 carcasses processed was swabbed for generic <i>E. coli</i> per regulatory requirements. Products intended for raw ground use were sampled and tested for <i>E. coli</i> O157:H7. Sampling and testing of finished products for non-O157 STEC, Salmonella, and indicator organisms including APC, coliforms, and generic <i>E. coli</i> was based on customer-specific requirements. Quarterly process validation swabs were collected on ten carcasses pre and post interventions; swabs were tested for APC, coliforms, and generic <i>E. coli</i>. Operation of this intervention method. 	Yes



3.7.3	Checking the nozzles to ensure that they are not plugged and that they are all functioning.	Yes
Comment:	Nozzle function was verified during CCP monitoring.	
3.7.4	Checking the position of the arbors (are they moving correctly, or if stationary, are they aimed correctly).	Yes
Comment:	Arbor operation was verified during CCP monitoring.	
3.7.5	Start-up and shut-down procedures.	Yes
Comment:	Start up and shut down procedures were defined within maintenance PMs and the CHAD owner's manual.	
3.7.6	There is documentation of a monitoring process that assures that the water or steam is as least 160°F at the carcass surface.	Yes
Comment:	Carcass surface temperature was verified twice hourly using a surface thermometer.	
3.7.7	The establishment monitors dwell time.	No
Comment:	Dwell time was not monitored.	
3.7.8	The establishment ensures that all areas and/or surfaces of the carcass are adequately covered by water or steam.	Yes
Comment:	Carcass coverage was visually verified during CCP monitoring.	
3.7.8	The establishment documents monitoring of start-up and shut-down.	Yes
Comment:	Start up and shut down procedures were documented through maintenance PMs.	
3.8	The establishment's intervention operating parameters fall within the validation supporting documentation parameters.	Yes
Comment:	Operating parameters fell within supporting validation documentation.	
1 Interve	ntions / Process Aids Chemical Applications	Result
4.1	The establishment uses Chemical Application(s) as an intervention method.	Yes
	The site utilized hypobromous acid, lactic acid, and ASC (acidified sodium chlorite) as chemical interventions.	100
	List each intervention chemical (ex. Lactic acid, peracetic acid, chlorine, Sanova, SYNTRx) bein and the location of use. Verify that the establishment has FSIS Regulatory approval or other re approval for the chemical(s) in use. Identify CCPs with parentheses.	
	ASC (500-1200PPM) was applied to applied to carcass mid-line at hide opening, bung area por removal and bunging, to the neck area of the carcass post-hide removal, to the inside cavity a underneath the inside skirt post-evisceration, to railed-out carcasses prior to re-entry to the mat through a pre-fabrication cabinet. Heads, tongues, tails, hearts, kidneys, and livers (CCP), on trim belts, and offline combo filling stations. Lactic acid (2%-10%) was applied after hot water pasteurization (either/or CCP) and on (2%-5 fabrication trim belts	nd ain rail, and fabrication %)

Hypobromous acid (Bovibrom, 75-900ppm) was applied to carcasses through the spray chill system.

Lactic acid was approved as available for use on the variety meats. Intervention chemicals were approved for use through FSIS Directive 7120.1.



4.2	NOTE: Answer the following questions for each designated CCP.	Yes
	The establishment identified this intervention as a CCP. If YES, identify the location of the application (ex. Post-evis lactic acid).	
Comment:	Final carcass (lactic acid) and variety meat application (ASC or lactic) were identified as CCPs. The site primarily used ASC for variety meats.	
4.3	If the Chemical Application is a CCP, can the line run if this intervention is not operational or not in specification.	Yes
Comment:	The line could run if the hot water carcass final wash was operational and the ASC or lactic acid with hand application spray for variety meat application was operational. Otherwise, the line could not run.	
4.4.1	None	Not Applicable
4.4.2	Validated Third Party Challenge Study or Validation Study	Yes
Comment:	Antimicrobial Spray Treatments for Red Meat Carcasses Processed in Very Small Establishments - Penn State University 2005.	
4.4.3	In-house Challenge Study or Validation Study	Yes
Comment:	ASC Vs. Lactic Acid - July 2020 Lactic Acid Validation January 2020	
4.4.4	Third Party review of in-house challenge study or validation. List the name of the Third Party in Comments.	No
4.4.5	Resource white paper (Published Journal Article)	Not Applicable



Comment:	Efficacy of Organic Acids Against Escherichia coli O157:H7 Attached to Beef Carcass
	Tissue Using a Pilot Scale Model Carcass Washer. Journal of Food Protection. Vol. 57, No.
	2, Pages 97-103. Cutter 1994

Evaluation of Various Antimicrobial Interventions for the Reduction of Escherichia coli O157:H7 on Bovine Heads During Processing. Journal of Food Protection. Vol. 71, No. 3, 2008. Pages 621-624. Kalchayanand 2008

Comparison of the Efficacy of a Sulfuric Acid Sodium Sulfate Blend and Lactic Acid for the Reduction of Salmonella on Pre-rigor Beef Carcass Surface Tissue. Journal of Food Protection. Vol. 80, No. 5, 2017. Pages 809-813. Yang 2017

Comparison of Water Wash, Trimming, and Combined Hot Water and Lactic Acid Treatments for Reducing Bacteria of Fecal Origin on Beef Carcasses. Journal of Food Protection. Vol. 61, No. 7, 1998. Pages 823-828. Castillo 1998

Lactic Acid Sprays Reduce Bacterial Pathogens on Cold Beef Carcass Surfaces and in Subsequently Produced Ground Beef. Journal of Food Protection. Vol. 64, No. 1, 2001. Pages 58-62. Castillo 2001

Effects of Cetylpyridinium Chloride, Acidified Sodium Chlorite and Potassium Sorbate on Populations of Escherichia coli O157:H7, Listeria monocytogenes, and Staphylococcus aureus on Fresh Beef. Journal of Food Protection. Vol. 67, No. 2, 2004. Pages 310-315. Lim 2004

Decreased Dosage of Acidified Sodium Chlorite Reduces Microbial Contamination and Maintains Organoleptic Qualities of Ground Beef Products. Journal of Food Protection. Vol. 67, No. 10, 2004. Pages 2248-2254.

Efficacy of Antimicrobial Compounds on Surface Decontamination of Seven Shiga Toxin-Producing Escherichia coli and Salmonella Inoculated onto Fresh Beef. Journal of Food Protection. Vol. 78, No. 3, 2015. Pages 503-510.

4.4.6	Resource white paper with third party review (peer reviewed paper - not published)	No
4.4.7	Other List in comments	Yes
Comment:	FSIS Directive 7120.1.	
1	A specific set of samples were chosen to support the validation hypothesis (objective).	Yes
Comment:	Specific sample sets were selected for the validation.	
2	Statistical parameters were used in the validation hypothesis and/or the analysis to support the conclusion.	Yes
Comment:	Log reduction of APC, coliforms, and generic <i>E. coli</i> supported the conclusion.	
3	Scientific support documentation.	Yes
Comment:	Microbiological testing data supported the conclusion.	
4	Validation study was prepared by a third party. List the name of the third party in comments.	No
Comment:	Validation study was prepared in-house.	
5	Other List in comments	Not Applicable



4.5.1	The establishment has records demonstrating on-going verification activities for this intervention. List the Frequency in comments.	Yes
Comment:	One out of every 300 carcasses processed was swabbed for generic <i>E. coli</i> per regulatory requirements. Products intended for raw ground use were sampled and tested for <i>E. coli</i> O157:H7. Sampling and testing of finished products for non-O157 STEC, Salmonella, and indicator organisms including APC, coliforms, and generic <i>E. coli</i> was based on customer-specific requirements. Quarterly process validation swabs were collected on ten carcasses pre and post interventions; swabs were tested for APC, coliforms, and generic <i>E. coli</i> .	
1	The establishment has documented procedures that include the following:	Yes
	Operation of this intervention method, including application of the treatment	
Comment:	Maintenance PMs defined operation and application requirements.	
2	Preparation of the treatment solution(s)	Yes
Comment:	Maintenance PMs defined solution preparation requirements.	
3	Start up of the intervention equipment	Yes
Comment:	Maintenance PMs defined start up and shut down requirements.	
4	Shut down of the intervention equipment	Yes
Comment:	Maintenance PMs defined start up and shut down requirements.	
4.6.1	The establishment monitors and has set lower limits on the concentration of the treatment solution. Specify in the comments if TITRATION or CONDUCTIVITY is used to monitor the solution concentration.	Yes
Comment:	Concentration of chemical interventions was verified through titration. Lower limits were established as 75 ppm for Bovibrom, 2% for lactic acid and 500 ppm for ASC.	
4.6.2	The establishment monitors the temperature of the treatment solutions.	Yes
Comment:	Lactic acid temperature was verified through CCP monitoring. Temperature of other chemical interventions was not monitored.	
4.6.3	The establishment monitors the flow / volume	Yes
Comment:	Application (flow) of chemical interventions was verified through CCP and control point monitoring.	
4.6.4	The establishment monitors the nozzle pressure.	Yes
Comment:	Nozzle pressure was verified for lactic acid through CCP monitoring. Nozzle pressure of other chemical interventions was not monitored.	
4.6.5	The establishment ensures all areas and/or surfaces of the carcass are adequately covered by the chemical application.	Yes
Comment:	Visual verification of coverage application was conducted through CCP and control point monitoring.	
4.6.6	The intervention method is implemented as written in the documented procedure.	Yes
Comment:	Chemical interventions were operating within defined operational requirements during the facility walk. Lactic was operating at 4.2%. ASC was observed at 1200PPM.	



4.7	The establishment's intervention operating parameters fall within the validation supporting documentation parameters.	Yes
Comment:	Operating parameters fell within and were compliant with defined validation parameters.	
4.8.1	Is / Are there alternative intervention methods(s) being utilized other than those listed in the previous pages	No
Comment:	Alternative interventions were not utilized.	
5 Dressir	ng Procedures / Critical Job Tasks	
		Result
5.1	Is there an intervention or process aid utilized upon entering or exiting the out rail.	Yes
Comment:	ASC was applied to carcasses upon exiting the out-rail.	
5.2	The establishment designates and has documented descriptions of critical job tasks (i.e., skinning line, evisceration, etc.).	Yes
Comment:	Slaughter SOP Job Tasks defined requirements for critical job tasks.	
5.3	The establishment uses hot water or chemical solution to sanitize equipment (i.e., knife, steel, hook, etc.) during operations.	Yes
Comment:	Hot water or bleach water was used to sanitize equipment after trimming contamination.	
5.4.1	The establishment uses the following to ensure that knives are in the sanitizer dip long enough to sanitize: List which methods are utilized in which process i.e. multiple knife rotation on skinning line, 1-2 second dip post skinning, etc.	Not Applicable
	Knife blade stays in the dip 1-2 seconds.	
Comment:	A 2-3 second dip was utilized for sanitizing equipment following evisceration tasks and post-evisceration trimming. A multiple knife rotation was utilized for sanitizing equipment on the skinning line to the point of evisceration.	
5.4.2	Knife blade stays in the dip 2-3 seconds.	Yes
Comment:	A 2-3 second dip was utilized for sanitizing equipment following evisceration tasks and post-evisceration trimming.	
5.4.3	Knife blade stays in the dip for 4-6 seconds.	No
Comment:	A 2-3 second dip was utilized for sanitizing equipment following evisceration tasks and post-evisceration trimming.	
5.4.4	Multiple knife rotation.	Yes
Comment:	A multiple knife rotation was utilized for sanitizing equipment on the skinning line to the point of evisceration.	
5.5	The establishment sanitizes all equipment (hooks and knives) between each use to reduce cross contamination in the process when trimming visible contamination (i.e., fecal, hair, or dirt.).	Yes
Comment:	Equipment was sanitized between each use following trimming of contamination.	



5.6	There is an auditing / observation process for monitoring of critical job tasks	Yes
Comment:	Sanitary dressing processes were monitored hourly by QA technicians and continuously by supervisors. Records from the week of 8/8/22 demonstrated compliance with the facility's procedure.	
5.7.1	Type(s) of monitoring at the establishment:	Yes
	Auditor	
Comment:	QA performed documented monitoring of sanitary dressing protocols hourly.	
5.7.2	Supervisor	Yes
Comment:	Supervisors performed continuous monitoring of sanitary dressing processes; monitoring was not documented.	
5.7.3	Video	No
Comment:	Video was not utilized	
5.7.4	Other List in Comments	Not Applicable
5.8	The Auditor declares that he/ she does not have a conflict of interest with this auditee and the audit has been carried out independently and impartially.	Yes
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Comment: I, Scott Devitt, do not have a conflict of interest with this auditee.