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K.J. Karr

Curtis L. Kastner

James L. Marsden

See next page for additional authors

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Control of Escherichia coli 0157:H7 in large-diameter, lebanon-style bologna

Abstract

Lebanon bologna raw batter was mixed with a five-strain mixture of Escherichia coli O157:H7 to achieve average inoculum levels of 7.79, 7.77, and 7.92 log CFU/g as deter mined on MSA, 202, and PRSA media, respectively. Treatment 1 consisted of a fermentation cycle of 8 hrs at an internal temperature (I.T.) of 80EF then 24 hrs at 100EF I.T., followed by 24 hrs at 110EF I.T. Treatments 2, 3, and 4 included additional heating at 115EF I.T. for 1, 2, and 5 hrs, respectively. All heat treatments resulted in product that was negative (<1.9 log CFU/g detection limit) on all culture media and negative after enrichment on mEC selective medium. This study validates that a five-log reduction of E. coli O157:H7 can be achieved using the described protocol, thus meeting USDA/FSIS requirements.

Keywords

Kansas Agricultural Experiment Station contribution; no. 97-309-S; Report of progress (Kansas State University. Agricultural Experiment Station and Cooperative Extension Service); 804; Cattlemen's Day, 1998; Beef; E. coli O157:H7; Food safety; Fermented beef; Sausage

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Authors

K.J. Karr, Curtis L. Kastner, James L. Marsden, and Randall K. Phebus

CONTROL OF *ESCHERICHIA COLI* O157:H7 IN LARGE-DIAMETER, LEBANON-STYLE BOLOGNA

K. J. Karr, C. L. Kastner, J. L. Marsden, and R. K. Phebus

Summary

Lebanon bologna raw batter was mixed with a five-strain mixture of Escherichia coli O157:H7 to achieve average inoculum levels of 7.79, 7.77, and 7.92 log CFU/g as determined on MSA, 202, and PRSA media, respectively. Treatment 1 consisted of a fermentation cycle of 8 hrs at an internal temperature (I.T.) of 80°F then 24 hrs at 100°F I.T., followed by 24 hrs at 110°F I.T. Treatments 2, 3, and 4 included additional heating at 115°F I.T. for 1, 2, and 5 hrs, respectively. All heat treatments resulted in product that was negative (<1.9 log CFU/g detection limit) on all culture media and negative after enrichment on mEC selective medium. This study validates that a five-log reduction of E. coli O157:H7 can be achieved using the described protocol, thus meeting USDA/FSIS requirements.

(Key Words: *E. coli* O157:H7, Food Safety, Fermented Beef, Sausage.)

Introduction

In December of 1994, an outbreak of *Escherichia coli* O157:H7 was linked to the consumption of dry cured salami. This outbreak caused USDA/FSIS to require that fermented sausage processes achieve a fivelog reduction of *E. coli* O157:H7 in a test situation when starting with at least 7.3 CFU.

Lebanon bologna is a fermented beef sausage that utilizes a low-temperature, long-time fermentation process, but is susceptible to *E. coli* O157:H7 contamination. An available medium is sensitive to recovering heatinjured cells. Therefore, the objectives of this study were: 1) to determine the effects of typical thermal processing temperatures and times for Lebanon bologna on reducing *E. coli* O157:H7 and 2) to evaluate the effectiveness of MacConkey Sorbitol Agar (MSA), 202 agar, and Phenol Red Agar with 1% sorbitol (PRSA) for detecting *E. coli* O157:H7.

Experimental Procedures

Five different isolates of *E. coli* O157:H7 were used. Two were human isolates, and the others were of meat origin, one being implicated in the 1995 salami outbreak. Isolates were incubated on tryptic soy agar slants at $98^{\circ}F$ for 20 ± 2 hrs and maintained at $40^{\circ}F$ until needed. After further inoculation, cells were harvested by centrifugation, resuspended, centrifuged again, and then held at $40^{\circ}F$ until needed (less than 2 hrs).

Commercially prepared beef meat batter (90% lean) containing salt; sucrose; dextrose; spices; potassium nitrate; sodium nitrite; and starter culture (*Pediococcus*, *Lactobacillus*, and *Micrococcus* spp.) was received overnight from the manufacturer. Upon receipt, the raw batter was at $45 \pm 4^{\circ}$ F.

For the inoculated treatments, 55 lb of meat batter was spread evenly (1 to 1.5 in. thick) onto a flat surface to allow for even distribution. The inoculum was intermittently pipetted drop-wise over the meat surface and thoroughly mixed.

The meat batters (control and inoculated) were transferred to a hand stuffer and stuffed into prestuck, presoaked, 4½ in.-diameter casings. Each chub weighed approximately 6.6 lb and was about 10 in. long.

Chubs were hung vertically on racks. Inoculated and control chubs were placed randomly in a commercial smoke house (Alkar, Lodi, WI). Fermentation included 8 hrs at an internal temperature (I.T.) of 80°F, then 24 hrs at 100°F I.T., followed by 24 hrs at 110°F I.T. Natural smoke was applied during the last 2 hrs of the 110°F cycle. Heat treatments 2, 3, and 4 included additional heating at 115°F I.T. for 1, 2, and 5 hrs, respectively. For each internal temperature, an appropirate time was allowed for that temperature to be reached. The relative humidity (RH) for the 80°F stage was 90%. For the 110, 110, and 115°F stages, the relative humidity was 60 to 65%. In the commercial Lebanon bologna process, the RH is maintained at 90% throughout the process, and the specified moisture to protein ratio is 3.1:1. Because of the lower relative humidity, our product is referred to as "Lebanon-style bologna."

MacConkey Sorbitol Agar, 202 agar, and PRSA were used for enumerating *E. coli* O157:H7. All plates were spiral plated and incubated at 107°F for 24 hrs. Modified *E. coli* broth was used for enrichment of *E. coli* O157:H7. Identification was confirmed with API 20# and RIM *E. coli* O157:H7 latex agglutination test.

A special medium (APT) was used for lactic acid bacteria (LAB) enumeration. All LAB plates were incubated at 95°F for 24 hrs in a CO₂ chamber with 20% CO₂.

Both raw and heat-treated samples were analyzed for moisture, fat, salt, protein, ash, water activity, pH, and titratable acidity.

Results and Discussion

For all heat treatments, the log (CFU/g) reduction values were 5.89, 5.87, and 6.07 on MSA, 202, PRSA media, respectively. A 6 log reduction means a kill of 99.9999% of original *E. coli* O157:H7 organisms. All heat treatment samples were also negative after enrichment on mEC selective medium. The LAB counts were between 7.2 and 7.4 log CFU/g for the raw batter and 6.8 to 6.9 log CFU/g for all of the heat treatments. A 7 log population is 10 million organisms.

Minimal variation was found for all product characteristics both within and between treatments. Overall pH was 4.4 after fermentation. Moisture was 60.8%; protein, 22.5%; fat, 10.6%; and salt, 4.8%. The moisture to protein ratio was 2.7, with water activity at 0.94. All heat treatments on all media resulted in a product that was negative (<1.9 log CFU/g detection limit) for *E. coli* O157:H7 and negative after enrichment on mEC selective medium. This study validates that a five-log reduction of *E. coli* O157:H7 can be achieved using the described heating protocol, thus meeting USDA/FSIS requirements.