

**TEXAS DEPARTMENT OF STATE HEALTH SERVICES
MEAT SAFETY ASSURANCE
AUSTIN, TEXAS**

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| <h1 style="margin:0;">MSA NOTICE</h1> | 06-18 | 1/25/2018 |
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**ADDITIONAL INFORMATION REGARDING INSTRUCTIONAL LABELING
STATEMENTS FOR RAW BEEF PRODUCTS SHIPPED TO INTERMEDIARY
OFFICIAL ESTABLISHMENTS PRIOR TO DELIVERY TO AN OFFICIAL
ESTABLISHMENT FOR FULL LETHALITY TREATMENT TO ADDRESS SHIGA
TOXIN-PRODUCING *ESCHERICHIA COLI* (STEC)**

NOTE: An instructional statement concerning STEC is a statement that addresses how the raw product is to be prepared or handled to ensure that the pathogen is eliminated or reduced to an undetectable level (e.g., “for cooking only” or “for full lethality treatment”).

I. PURPOSE

A. This notice clarifies instructions to inspection program personnel (IPP) for verifying an establishment’s use of instructional labeling statements concerning STEC when raw beef products are shipped to one or more official establishments before delivery to the official establishment for cooking or other full lethality treatment.

B. IPP are to continue to follow the instructions in MSA Directive 10,010.2, *Verification Activities for Shiga Toxin-Producing Escherichia coli in Raw Beef Products*, for products labeled with an instructional statement and shipped directly to an official establishment for cooking or other full lethality treatment.

C. This notice only applies to product that has not been tested or has tested negative for STEC. IPP are to follow verification procedures in MSA Directive 10,010.2 for product that is presumptive or confirmed positive for STEC and labeled with an instructional statement.

II. BACKGROUND

Products labeled with instructional statements may be produced and labeled at one establishment and undergo further processing (e.g., repackaging, grinding) at an intermediate, non-cooking official establishment prior to being sent to another official establishment for cooking or other full lethality treatment. If the product is to undergo further processing at an intermediary establishment, the intermediary establishment is to address the potential for cross-contamination in its HACCP system. ID warehouses and brokers are not official establishments, and cannot re-

box or further process product labeled with instructional or disclaimer statements. ID warehouses may store product only (9 CFR 412.1(e)).

III. VERIFICATION ACTIVITIES AT PRODUCING ESTABLISHMENTS

A. When conducting a HACCP verification task, IPP at an establishment that applies instructional statements are to follow the instructions in MSA Directive 10,010.2, to verify the producing establishment maintains and implements sanitation procedures to prevent cross-contamination and maintains records adequate to demonstrate the product was sent to an official establishment for a full lethality process.

B. When IPP identify noncompliance, they are to document the noncompliance on a noncompliance record (NR) as described in MSA Directive 5000.1, using the HACCP verification task and the appropriate regulatory citation (usually 9 CFR 417.5).

IV. VERIFICATION ACTIVITIES AT INTERMEDIARY ESTABLISHMENTS

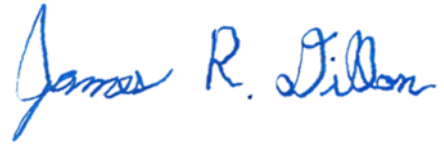
A. A producing establishment may ship product to one or more intermediary establishments before the product is delivered to the official establishment that applies the full lethality treatment.

B. Intermediary establishments that receive product labeled with an instructional statement and further process the product may reapply (i.e., “carry forward”) the instructional statement without label approval. IPP at intermediary establishments that carry forward labeling of product with an instructional statement are to:

1. Follow the instructions in MSA Directive 10,010.2 to verify the establishment is appropriately using the instructional statement. The HACCP system for establishments that carry forward the instructional statement do not need to include a validated intervention for STEC as the product is intended for cooking or other full lethality treatment;
2. Verify the establishment's hazard analysis (9 CFR 417.2) and decision-making documents (9 CFR 417.5) meet the criteria in MSA Directive 10,010.2 when performing the HACCP verification task;
3. Verify the establishment tracks and facilitates communication between the supplying establishments and receiving establishments to ensure records are available showing each lot of product was sent to an establishment for cooking or other full lethality treatment; and
4. IPP are to document noncompliance on an NR as described in MSA Directive 5000.1 using the HACCP verification task and the appropriate regulatory citation (usually 9 CFR 417.5) when they find that the intermediate establishment has not met the criteria above.

V. QUESTIONS

Refer questions through supervisory channels.

A handwritten signature in blue ink that reads "James R. Dillon". The signature is written in a cursive style with a large initial 'J' and 'D'.

James R. Dillon, DVM, MPH
Director, Texas State Meat and Poultry Inspection Program
Department of State Health Services