

According to the Paperwork Reduction Act of 1995, an agency may not conduct or sponsor, and a person is not required to respond to, a collection of information unless it displays a valid OMB control number. The valid OMB control number for this information collection is 0583-0144. The time required to complete this information collection is estimated to average 30 minutes per response, including the time for reviewing instructions, searching existing data sources, gathering and maintaining the data needed, and completing and reviewing the collection of information.

## INDUSTRY REPORT OF ADULTERATION:

Report of adulterated or misbranded meat/poultry product received from or shipped to commerce by the official establishment  
*Attach any supporting or supplemental documentation*

**REPORTED DATE:** *(date reported to the Agency):*

**NOTIFIER INFORMATION** *(identify the establishment representative reporting the incident)*

First Name:	<input type="text"/>	Last Name:	<input type="text"/>
Telephone Number:	<input type="text"/>	Ext:	<input type="text"/>
		Email:	<input type="text"/>

**Notifying Establishment Role:** *(identify the establishment representative reporting the incident)*

<input type="checkbox"/> Shipping Establishment	<input type="checkbox"/> Receiving Establishment
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Notifying Establishment Name:	<input type="text"/>
Notifying Establishment Number:	<input type="text"/>

Notifier Information Additional Comments:

**INFORMATION FOR OTHER INVOLVED ESTABLISHMENTS:**

**Additional Establishment Role** *(provide information for other establishments involved):*

<input type="checkbox"/> Shipping Establishment	<input type="checkbox"/> Receiving Establishment
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Additional Establishment Name:	<input type="text"/>
Additional Establishment Number:	<input type="text"/>

**Comments for additional establishment(s):**

**PRODUCT INFORMATION:**

Date the Adulteration or Misbranding was identified: \_\_\_\_\_

Identified issue(s):

Pathogen

Extraneous Material

Undeclared Allergen

Mislabeling

Unapproved Substance

Undeclared Substance

Residue

Undeclared Substance

Produced Without Benefit of Inspection

SRM

Insanitary Conditions

Failure to Present Import Reinspection

**Issue(s) Description (include specific details to describe the issue and how and when the problem was discovered)**

**Likely Root Cause (describe how and when the issue occurred, including any production dates):**

Date Shipped: \_\_\_\_\_ Date Received: \_\_\_\_\_

Product Name: \_\_\_\_\_ Lot Code/identifier: \_\_\_\_\_

Establishment Name on Product: \_\_\_\_\_

Producer Name on Product: \_\_\_\_\_

HACCP Category: \_\_\_\_\_ Finished Products Type: \_\_\_\_\_

Species: \_\_\_\_\_ Product Group: \_\_\_\_\_

**PRODUCT QUANTITIES (provide in pounds):**

Implicated: \_\_\_\_\_ In Commerce: \_\_\_\_\_ Under Control: \_\_\_\_\_

**STATUS OF IMPLICATED PRODUCT (if applicable):**

Amount condemned: \_\_\_\_\_

Location(s) of: product under control: \_\_\_\_\_

**CARRIER INFORMATION (if applicable)** Carrier Name: \_\_\_\_\_

Carrier Phone Number: \_\_\_\_\_ Carrier Address: \_\_\_\_\_

**ADDITIONAL COMMENTS:**