# **Notices**

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This section of the FEDERAL REGISTER contains documents other than rules or proposed rules that are applicable to the public. Notices of hearings and investigations, committee meetings, agency decisions and rulings, delegations of authority, filing of petitions and applications and agency statements of organization and functions are examples of documents appearing in this section.

## **DEPARTMENT OF AGRICULTURE**

## Submission for OMB Review; Comment Request

July 28, 2021.

The Department of Agriculture has submitted the following information collection requirement(s) to OMB for review and clearance under the Paperwork Reduction Act of 1995, Public Law 104–13. Comments are requested regarding whether the collection of information is necessary for the proper performance of the functions of the agency, including whether the information will have practical utility; the accuracy of the agency's estimate of burden including the validity of the methodology and assumptions used; ways to enhance the quality, utility and clarity of the information to be collected; and ways to minimize the burden of the collection of information on those who are to respond, including through the use of appropriate automated, electronic, mechanical, or other technological collection techniques or other forms of information technology.

Comments regarding this information collection received by September 1, 2021 will be considered. Written comments and recommendations for the proposed information collection should be submitted within 30 days of the publication of this notice on the following website www.reginfo.gov/public/do/PRAMain. Find this particular information collection by selecting "Currently under 30-day Review—Open for Public Comments" or by using the search function.

An agency may not conduct or sponsor a collection of information unless the collection of information displays a currently valid OMB control number and the agency informs potential persons who are to respond to the collection of information that such persons are not required to respond to the collection of information unless it

displays a currently valid OMB control number.

## **Food Safety and Inspection Service**

*Title:* Consumer Complaint Monitoring System.

OMB Control Number: 0583-0133. Summary of Collection: The Food Safety and Inspection Service (FSIS) has been delegated the authority to exercise the functions of the Secretary as provided in the Federal Meat Inspection Act (FMIA) (21 U.S.C. 601 et seq.), the Poultry Products Inspection Act (PPIA) (21 U.S.C. 451 et seq.), and the Egg Product Inspection Act (EPIA) (21 U.S.C. 1031 et seq.). These statutes mandate that FSIS protect the public by ensuring that meat and poultry products are safe, wholesome, unadulterated, and properly labeled and packaged. FSIS tracks consumer complaints about meat, poultry, and egg products. Consumer complaints are usually filed because food made the consumer sick, caused an allergic reaction, was not properly labeled (misbranded).

Need and Use of the Information: The Consumer Complaint Monitoring System web portal is used primarily to track consumer complaints regarding meat, poultry, and egg products. FSIS will use the information collected from the web portal. To not collect the information from the web portal would reduce the effectiveness of the meat, poultry, and egg products inspection program.

Description of Respondents: Individuals or households.

Number of Respondents: 3,000. Frequency of Responses: Reporting: On occasion.

Total Burden Hours: 750.

## **Food Safety and Inspection Service**

*Title:* Voluntary Recalls of Meat and Poultry Products.

OMB Control Number: 0583–0135. Summary of Collection: The Food Safety and Inspection Service (FSIS) has been delegated the authority to exercise the functions of the Secretary as provided in the Federal Meat Inspection Act (FMIA) (21 U.S.C. 601 et seq.) and the Poultry Products Inspection Act (PPIA) (21 U.S.C. 451 et seq.) These statutes mandate that FSIS protect the public by ensuring that meat and poultry products are safe, wholesome, unadulterated, and properly labeled and packaged. A firm that has produced or imported meat or poultry that is

adulterated or misbranded and is being distributed in commerce, may voluntarily recall the product in question. When a firm voluntarily recalls a product, FSIS will conduct a recall effectiveness check.

Need and Use of the Information: In conducting a recall, the establishment will be asks to provide FSIS with some basic information, including the identity of the recalled product, the reason for the recall, and information about the distributors and customers of the product. Industry representatives use the FSIS Form 5020–3 FSIS Preliminary Inquiry Worksheet to provide firm contact information and specific details regarding adulterated or misbranded product in commerce, including product identifiers, product amounts and supplemental information. Recalling firms and distributors then use the FSIS Form 5020-4 FSIS Recall Distribution Information Template to provide the location and contact information of consignees who received recalled product. FSIS will check on the effectiveness of the recall to ensure that all products subject to recall are accounted for. FSIS field personnel will use FSIS form 8400-4 A to determine (1) if the retail consignee received notification of the recall and (2) the number of recalled products received. FSIS field personnel will also use FSIS form 8400-4 B to verify that product held by the retail consignee was properly disposed.

Description of Respondents: Business or other for-profit.

Number of Respondents: 6,090. Frequency of responses: Reporting: On Occasion.

Total Burden Hours: 6,600.

## **Food Safety and Inspection Service**

*Title:* Animal Disposition Reporting System.

OMB Control Number: 0583–0139.

Summary of Collection: The Food
Safety and Inspection Service (FSIS) has
been delegated the authority to exercise
the functions of the Secretary as
provided in the Federal Meat Inspection
Act (FMIA) (21 U.S.C. 601 et seq.) and
the Poultry Products Inspection Act
(PPIA) (21 U.S.C. 451 et seq.). These
statutes mandate that FSIS protect the
public by ensuring that meat and
poultry products are safe, wholesome,
unadulterated, and properly labeled and
packaged. FSIS also inspects exotic
animals and rabbits under the authority

of the Agricultural Marketing Act of 1946, as amended (7 U.S.C. 1621 et seq.). In accordance with 9 CFR 320.6, 381.180, 352.15, and 354.91, establishments that slaughter meat, poultry, exotic animals, and rabbits are required to maintain certain records regarding their business operations and to report this information to the Agency as required.

Need and Use of the Information: FSIS will collect information from establishments using FSIS Form 6510-7, Poultry Lot Information. Poultry establishments complete the form after each shift and submit it to the agency. FSIS uses this information to plan inspection activities, to develop sampling plans, to target establishments for testing, to develop Agency budget, and to develop reports to Congress.

Description of Respondents: Business

or other for-profit.

Number of Respondents: 1,159. Frequency of Responses: Reporting: Other (daily).

Total Burden Hours: 23.180.

## Food Safety and Inspection Service

*Title:* Requirements to Notify FSIS of Adulterated or Misbranded Product, Prepare and Maintain Written Recall Procedures, and Document Certain HACCP Plan Reassessments.

OMB Control Number: 0583–0144. Summary of Collection: The Food Safety and Inspection Service (FSIS) has been delegated the authority to exercise the functions of the Secretary as provided in the Federal Meat Inspection Act (FMIA) (21 U.S.C. 601 et seq.) and the Poultry Products Inspection Act (PPIA) (21 U.S.C. 451 et seq.). These statutes mandate that FSIS protect the public by verifying that meat and poultry products are safe, wholesome, unadulterated, and properly labeled and packaged. Section 11017 of the Food, Conservation, and Energy Act of 2008 (Pub. L. 110-246, 112 Stat 1651, 448-49), amended the FMIA and the PPIA by adding sections 12 and 13 to the FMIA and by amending section 10 of the PPIA (21 U.S.C. 459). These sections require official establishments that believe, they have shipped into commerce or received, misbranded, or adulterated products to notify the Secretary of Agriculture.

Need and Use of the Information: Official establishments are to document each time they reassess their HACCP plans and make the reassessments available to FSIS officials for review and copying. Official establishments are to notify the FSIS District Office that they have received or have shipped into commerce misbranded or adulterated product. The information collected will

permit FSIS officials to monitor closely establishments HACCP plan reassessments and to facilitate recalls or adulterated or misbranded product.

Description of Respondents: Business or other for-profit.

Number of Respondents: 6,300. Frequency of Responses: Reporting: On occasion.

Total Burden Hours: 9.960.

#### Ruth Brown.

Departmental Information Collection Clearance Officer.

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## **DEPARTMENT OF AGRICULTURE**

## **U.S. Codex Office**

**Codex Alimentarius Commission:** Meeting of the Ad hoc Codex Intergovernmental Task Force on **Antimicrobial Resistance** 

AGENCY: U.S. Codex Office, USDA. **ACTION:** Notice of public meeting and request for comments.

**SUMMARY:** The U.S Codex Office is sponsoring a public meeting on September 9, 2021. The objective of the public meeting is to provide information and receive public comments on agenda items and draft United States (U.S.) positions to be discussed at the 8th Session of the Ad hoc Codex Intergovernmental Task Force on Antimicrobial Resistance (TFAMR) of the Codex Alimentarius Commission. which will convene virtually, October 4–9, 2021 with the report adoption on October 13, 2021. The U.S. Manager for Codex Alimentarius and the Acting Deputy Under Secretary for Trade and Foreign Agricultural Affairs recognize the importance of providing interested parties the opportunity to obtain background information on the 8th Session of the TFAMR and to address items on the agenda.

**DATES:** The public meeting is scheduled for September 9, 2021, from 10:00 a.m. -12:00 p.m. EDT.

ADDRESSES: The public meeting will take place via Video Teleconference only. Documents related to the 8th Session of the TFAMR will be accessible via the internet at the following address: http://www.fao.org/fao-who-codexali mentarius/meetings/detail/en/ ?meeting=TFAMR&session=8.

Dr. Donald A. Prater, U.S. Delegate to the 8th Session of the TFAMR, invites U.S. interested parties to submit their comments electronically to the following email address: donald.prater@ fda.hhs.gov.

Registration: Attendees must register to attend the public meeting here: https://www.zoomgov.com/meeting/ register/vJIscOqsqTIrG4XB3KA 9PuFQT5y28NJF068 by September 2, 2021. Early registration is encouraged.

FOR FURTHER INFORMATION CONTACT: 8th Session of the TFAMR, contact U.S. Delegate, Dr. Donald A. Prater, donald.prater@fda.hhs.gov, +1 (301) 348-3007. For Further Information about the public meeting Contact: U.S. Codex Office, 1400 Independence Avenue SW, Room 4861, South Agriculture Building, Washington, DC 20250. Phone (202) 720-7760, Fax: (202) 720-3157, Email: uscodex@usda.gov

## SUPPLEMENTARY INFORMATION:

#### **Background**

Codex was established in 1963 by two United Nations organizations, the Food and Agriculture Organization and the World Health Organization. Through adoption of food standards, codes of practice, and other guidelines developed by its committees, and by promoting their adoption and implementation by governments, Codex seeks to protect the health of consumers and ensure fair practices in the food

The Terms of Reference of the Ad hoc Codex Intergovernmental Task Force on Antimicrobial Resistance (TFAMR) are:

- (i) To review and revise, as appropriate, the *Code of Practice to* Minimize and Contain Antimicrobial Resistance (CXC 61-2005) to address the entire food chain, in line with the mandate of Codex.
- (ii) To consider the development of Guidance on Integrated Surveillance of Antimicrobial Resistance, taking into account the guidance developed by the WHO Advisory Group on Integrated Surveillance of Antimicrobial Resistance (AGISAR) and relevant OIE documents.

The TFAMR is hosted by the Republic of Korea. The United States attends the TFAMR as a member country of Codex.

Issues to be Discussed at the Public Meeting: The following items on the Agenda for the 8th Session of the TFAMR will be discussed during the public meeting:

- Proposed draft revision to the *Code* of Practice to Minimize and Contain Foodborne Antimicrobial Resistance (COP)(CXC 61-2005); and
- Proposed draft Guidelines on Integrated Monitoring and Surveillance of Foodborne Antimicrobial Resistance (GLIS).

Public Meeting: At the September 9, 2021, public meeting, draft U.S. positions on the agenda items will be