

Comments: ACF specifically requests comments on (a) whether the proposed collection of information is necessary for the proper performance of the functions of the agency, including whether the information shall have practical utility; (b) the accuracy of the agency's estimate of the burden of the proposed collection of information; (c) the quality, utility, and clarity of the information to be collected; and (d) ways to minimize the burden of the collection of information on respondents, including through the use of automated collection techniques or other forms of information technology. Consideration will be given to comments and suggestions submitted within 60 days of this publication.

Authority: This supplement is sponsored by the ACF OCSE and is authorized by Title IV–D of the Social Security Act. This supplement will be conducted by the U.S. Census Bureau and is authorized by 13 U.S.C. 182 which states, “the Secretary may make surveys deemed necessary to furnish annual and other interim current data on the subjects covered by the censuses provided for in this title.”

Mary C. Jones,
ACF/OPRE Certifying Officer.

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DEPARTMENT OF HEALTH AND HUMAN SERVICES

Food and Drug Administration

[Docket No. FDA–2025–N–0183]

Agency Information Collection Activities; Submission for Office of Management and Budget Review; Comment Request; Establishing and Maintaining Lists of United States Establishments With Interest in Exporting Human Food Program-Regulated Products

AGENCY: Food and Drug Administration, HHS.

ACTION: Notice.

SUMMARY: The Food and Drug Administration (FDA) is announcing that a proposed collection of information has been submitted to the Office of Management and Budget (OMB) for review and clearance under the Paperwork Reduction Act of 1995.

DATES: Submit written comments (including recommendations) on the collection of information by September 22, 2025.

ADDRESSES: To ensure that comments on the information collection are received,

OMB recommends that written comments be submitted to <https://www.reginfo.gov/public/do/PRAMain>. Find this particular information collection by selecting “Currently under Review—Open for Public Comments” or by using the search function. The OMB control number for this information collection is 0910–0509. Also include the FDA docket number found in brackets in the heading of this document.

FOR FURTHER INFORMATION CONTACT:

Domini Bean, Office of Operations, Food and Drug Administration, Three White Flint North, 10A–12M, 11601 Landsdown St., North Bethesda, MD 20852, 301–796–5733, PRASStaff@fda.hhs.gov.

SUPPLEMENTARY INFORMATION: In compliance with 44 U.S.C. 3507, FDA has submitted the following proposed collection of information to OMB for review and clearance.

Establishing and Maintaining Lists of U.S. Establishments With Interest in Exporting HFP-Regulated Products

OMB Control Number 0910–0509—Extension

This information collection supports Agency export programs and associated guidance. The United States exports a large volume and variety of foods in international trade. Foreign governments often require official certification from the responsible authority of the country of origin about imported foods and establishments involved in their production, storage, or distribution. Some foreign governments establish additional requirements with which exporters are required to comply and ask for additional assurances from the responsible authority. Importing countries may require, and FDA may provide, official certification or assurances for food products in different forms, including certificates that accompany specific products or lists of establishments and products that comply with certain requirements.

To facilitate exports of food subject to importing country listing requirements, FDA has historically provided official certification in the form of country- and product-specific export lists that include establishments and their products when: (1) the establishment has expressed interest in exporting their products to these countries; (2) the establishment and the products are subject to FDA's jurisdiction; and (3) the establishment can demonstrate that it is in good regulatory standing for the products it intends to export, and the products are expected to comply with

applicable FDA requirements. As we advised in the guidance document “Establishing and Maintaining a List of U.S. Milk and Milk Product, Seafood, Infant Formula, and Formula for Young Children Manufacturers/Processors with Interest in Exporting to China,” FDA considers “good regulatory standing” as meaning that an establishment is in substantial compliance with applicable FDA requirements and is not the subject of a pending enforcement action (e.g., an injunction or seizure) or pending administrative action (e.g., a warning letter).

FDA has generally published guidance documents for these country and product-specific lists under the authority of section 701(h) of the Federal Food, Drug, and Cosmetic Act (FD&C Act) (21 U.S.C. 371(h)), which authorizes the Secretary of Health and Human Services (the Secretary) to develop guidance documents with public participation presenting the views of the Secretary on matters under the jurisdiction of FDA. The guidance documents generally explain what information establishments should submit to FDA to be considered for inclusion on the lists and what criteria FDA intends to use to determine eligibility for placement on the lists. The guidance documents also explain how FDA intends to update the lists and communicate any new information to the governments that requested the lists. Finally, the guidance documents note that the information is provided voluntarily by establishments with the understanding that it may be posted on FDA's external website and that it will be communicated to, and possibly further disseminated by, the government that requested the list; thus, FDA considers the information on the lists to be information that is not protected from disclosure under 5 U.S.C. 552(b)(4). The guidance documents include “Establishing and Maintaining a List of U.S. Dairy Product Manufacturers/Processors with Interest in Exporting to Chile” (November 2018) and “Establishing and Maintaining a List of U.S. Milk and Milk Product, Seafood, Infant Formula, and Formula for Young Children Manufacturers/Processors with Interest in Exporting to China” (November 2018) available at <https://www.fda.gov/food/guidance-regulation-food-and-dietary-supplements/guidance-documents-regulatory-information-topic-food-and-dietary-supplements>. Additional information about FDA's Food Export Lists program is available at <https://www.fda.gov/food/exporting-food-products-united-states/food-export-lists>.

FDA has also published guidance on export certification that contains useful information that applies to export lists: “FDA Export Certification” (August 2021) available at <https://www.fda.gov/regulatory-information/search-fda-guidance-documents/fda-export-certification>.

Foreign governments are increasingly relying on certification as a strategy for ensuring the safety of imported food products, and many countries have announced new requirements for lists of establishments and products certified to comply with certain food safety requirements. FDA is committed to facilitating compliance with new listing requirements for U.S. establishments that export FDA-regulated food products by establishing and maintaining country- and product-specific export lists.

Application for inclusion on all export lists will continue to be voluntary. However, some foreign governments may require inclusion on export lists as a precondition for market access or to satisfy other importing country registration or approval requirements. FDA uses the Export Listing Module (ELM), an electronic system (Form FDA 3972), to receive and process applications for inclusion on

export lists for HFP-regulated products. The ELM allows applicants to provide information about the products intended for export, the establishment that produces those products, evidence of the establishment’s compliance with applicable requirements for the products intended for export, and any additional data or information (such as third-party certifications) that foreign governments may require. We request that this information be updated every 2 years. Additional information and screenshots of the ELM are available at <https://www.fda.gov/food/exporting-food-products-united-states/food-export-lists>. If an establishment is unable to submit an application via the ELM, it may contact HFP and request assistance.

We use the information submitted by establishments to determine eligibility for certification and inclusion on the export lists, which may be published on our website or the websites of foreign governments. The purpose of the lists is to help HFP-regulated industries meet the import requirements of foreign governments. This collection of information is intended to cover all of HFP’s existing export lists, as well as any additional export lists established by the program.

FDA notes section 801 of the FD&C Act (21 U.S.C. 381) also provides that FDA may charge a fee of up to \$175 if the Agency issues export certification within 20 days of receipt of a complete request for such certification.

Description of Respondents: Respondents to this collection of information include U.S. establishments subject to FDA/HFP jurisdiction that wish to be included on export lists.

In the **Federal Register** of May 1, 2025 (90 FR 18688), FDA published a 60-day notice requesting public comment on the proposed collection of information. We received two comments, believing one to have been posted erroneously to docket FDA–2025–N–0183 because it discussed vaccines. The second comment offered general support for the information collection and suggested technical enhancements to FDA’s collection mechanisms. Because FDA developed the ELM to facilitate the export of food and avoid delay, we appreciate this comment. At the same time, we are only able to make updates to our automated systems as our limited resources allow. We made no adjustment to our estimate after review of the public comments.

FDA estimates the burden of this collection of information as follows:

TABLE 1—ESTIMATED ANNUAL REPORTING BURDEN¹

IC activity recommended by guidance	Number of respondents	Number of responses per respondent	Total annual responses	Average burden per response (hours)	Total hours
New request	167	5	835	1	835
New request + third-party certification	85	2	170	22	3,740
Biennial update	132	4	528	0.5 (30 minutes)	264
Biennial update + third-party certification	58	2	116	22	2,552
Occasional updates	60	2	120	0.5 (30 minutes)	60
Total			1,769		7,451

¹ There are no capital costs or operating and maintenance costs associated with this collection of information.

Based on a review of the information collection since our last request for OMB approval, we have made no adjustments to our burden estimate.

Dated: August 15, 2025.

Grace R. Graham,
Deputy Commissioner for Policy, Legislation, and International Affairs.
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DEPARTMENT OF HEALTH AND HUMAN SERVICES

Food and Drug Administration

[Docket No. FDA–2024–D–1242]

Animal Studies for Dental Bone Grafting Material Devices—Premarket Notification (510(k)) Submissions; Guidance for Industry and Food and Drug Administration Staff; Availability

AGENCY: Food and Drug Administration, HHS.

ACTION: Notice of availability.

SUMMARY: The Food and Drug Administration (FDA or Agency) is announcing the availability of a final guidance entitled “Animal Studies for Dental Bone Grafting Material Devices—

Premarket Notification (510(k)) Submissions.” This guidance document provides recommendations for animal study design and animal study information to include to support a 510(k) submission for dental bone grafting material devices. This guidance may help manufacturers comply with some special controls for dental bone grafting material devices. The recommendations reflect current review practices and are intended to promote consistency and facilitate efficient review of these submissions.

DATES: The announcement of the guidance is published in the **Federal Register** on August 22, 2025.

ADDRESSES: You may submit either electronic or written comments on Agency guidances at any time as follows: