combined screener and topical, for a total of 126,703 respondents.

Average Hours per Response: 5 minutes per screener response and 35–36 minutes per topical response, which in total is approximately 40–41 minutes for households with eligible children.

Burden Hours: 42,863.

Needs and Uses: The National Survey of Children's Health (NSCH) enables the Maternal and Child Health Bureau (MCHB) of the Health Resources and Services Administration (HRSA) of the U.S. Department of Health and Human Services (HHS) along with supplemental sponsoring agencies, states, and other data users to produce national and statebased estimates on the health and wellbeing of children, their families, and their communities as well as estimates of the prevalence and impact of children with special health care needs.

Data will be collected using one of two modes. The first mode is a web instrument (Centurion) survey that contains the screener and topical instruments. The web instrument first will take the respondent through the screener questions. If the household screens into the study, the respondent will be taken directly into one of the three age-based topical sets of questions. The second mode is a mailout/mailback of a self-administered paper-and-pencil interviewing (PAPI) screener instrument followed by a separate mailout/mailback of a PAPI age-based topical instrument.

The National Survey of Children's Health (NSCH) is a large-scale (sample size is approximately 375,000 addresses) national survey with approximately 73,000 of those addresses included as part of twelve separate state-based or region-based oversamples. As in prior cycles of the NSCH, there remain two key, non-experimental design elements. The first nonexperimental design element is the use of an unconditional incentive (\$5) in the initial screener and topical invitations. For the initial screener invitation, 90% of sampled addresses receive the cash incentive; the remaining 10% (the control) do not receive an incentive. This approach is used to consistently monitor the effectiveness of the cash incentive each cycle. The second nonexperimental design element is a data collection procedure based on the block group-level paper-only response probability used to identify households (30% of the sample) that would be more likely to respond by paper and send them a paper questionnaire in the initial mailing and every nonresponse followup mailing.

The 2025 NSCH will also include a web targeted secondary unconditional screener incentive envelope test that

will be sent to a subset of the sample that started the web questionnaire but did not finish. Prior cycles of the survey have included a \$5 unconditional cash incentive with both the initial screener mailing as well as the initial paper topical mailing as outlined in the paragraph above. The incentive has proven to be a cost-effective intervention for increasing survey response and reducing nonresponse bias. The test will be used to evaluate envelope and delivery method effectiveness (FedEx vs. Visible *Incentive*). Preliminary results of other Demographic household surveys indicate the use of visual incentive envelopes (USPS) also helps to boost response and is less costly than FedEx.

Affected Public: Individuals or households.

Frequency: The 2025 collection is the tenth administration of the NSCH. It is an annual survey, with a new sample drawn for each administration.

Respondent's Obligation: Voluntary.

Legal Authority: Census Authority: Title 13, United States Code (U.S.C.), section 8(b) (13 U.S.C. 8(b)).

HRSA MCHB Authority: Section 501(a)(2) of the Social Security Act (42 U.S.C. 701).

United States Department of Health and Human Services' Centers for Disease Control and Prevention, National Center on Birth Defects and Developmental Disabilities; Division of Nutrition, Physical Activity, and Obesity; and Division of Environmental Health Science and Practice Authority: Public Health Service Act, Section 301, 42 U.S.C. 241 and 301(a), 307, and 399G of the PHS 42 U.S.C. 241A, 242l, 280e–11, as amended.

United States Department of Agriculture Authority: Richard B. Russell National School Lunch Act, 42 U.S.C. 1755 (a)(3).

This information collection request may be viewed at *www.reginfo.gov*. Follow the instructions to view the Department of Commerce collections currently under review by OMB.

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 and

entering either the title of the collection or the OMB Control Number 0607–0990.

Sheleen Dumas,

Departmental PRA Compliance Officer, Office of the Under Secretary for Economic Affairs, Commerce Department.

[FR Doc. 2025-05460 Filed 3-28-25; 8:45 am]

BILLING CODE 3510-07-P

DEPARTMENT OF COMMERCE

Office of the Secretary

Agency Information Collection Activities; Submission to the Office of Management and Budget (OMB) for Review and Approval; Comment Request; Foreign National Request Form

AGENCY: Office of Security, Insider Risk and Continuity, Office of the Secretary, Commerce.

ACTION: Notice of information collection, request for comment.

SUMMARY: The Department of Commerce, in accordance with the Paperwork Reduction Act of 1995 (PRA), invites the general public and other federal agencies to comment on proposed and continuing information collections, which helps us assess the impact of our information collection requirements and minimize the public's reporting burden. The purpose of this notice is to allow for 60 days of public comment preceding submission of the collection to OMB.

DATES: To ensure consideration, comments regarding this proposed information collection must be received on or before May 30, 2025.

ADDRESSES: Interested persons are invited to submit written comments by mail to *PRAcomments@doc.gov*. Please reference OMB Control Number 0690–0033 in the subject line of your comments. Do not submit Confidential Business Information or otherwise sensitive or protected information.

FOR FURTHER INFORMATION CONTACT:

Requests for additional information or specific questions related to collection activities should be directed to Mackenzie McGuire, Plans, Programs & Compliance Division, Office of Security, Department of Commerce, 1401 Constitution Ave. NW, Washington, DC 20230, email: mmcguire1@doc.gov or telephone: 202–615–6702.

SUPPLEMENTARY INFORMATION:

I. Abstract

The purpose of this collection is to gather information to mitigate variances in foreign access management program implementation and registration information requirements needed to reach risk-based determinations of physical and logical access by foreign national visitors and guests to Commerce facilities and resources. Due to the increasing diversity of foreign national participation in Departmental programs, considerable efforts have been made to baseline requirements as a means to define uniform program standards as well as to expand current guidance beyond foreign visitor control to manage emerging risks associated with physical and logical access to the Department's facilities and resources. This form has been revised to ensure that it adequately collects all information mandated by the Office of the Director of National Intelligence and aligns with updated guidance from the Office of Personnel Management.

II. Method of Collection

This information is collected in both paper form and electronically.

III. Data

OMB Control Number: 0690–0033. Form Number(s): 207–12–1.

Type of Review: Regular submission. Revision.

Affected Public: Individuals or households.

Estimated Number of Respondents: 16,000.

Estimated Time per Response: 15 minutes.

Estimated Total Annual Burden Hours: 4,000.

Estimated Total Annual Cost to Public: \$0.

Respondent's Obligation: Voluntary. Legal Authority: DOO 20–6 and DAO 207–12.

IV. Request for Comments

We are soliciting public comments to permit the Agency to: (a) Evaluate whether the proposed information collection is necessary for the proper functions of the Department, including whether the information will have practical utility; (b) Evaluate the accuracy of our estimate of the time and cost burden for this proposed collection, including the validity of the methodology and assumptions used; (c) Evaluate ways to enhance the quality, utility, and clarity of the information to be collected; and (d) Minimize the reporting burden on those who are to respond, including the use of automated collection techniques or other forms of information technology.

Comments that you submit in response to this notice are a matter of public record. We will include or summarize each comment in our request

to OMB to approve this ICR. Before including your address, phone number, email address, or other personal identifying information in your comment, you should be aware that your entire comment—including your personal identifying information—may be made publicly available at any time. While you may ask us in your comment to withhold your personal identifying information from public review, we cannot guarantee that we will be able to do so.

Sheleen Dumas,

Departmental PRA Compliance Officer, Office of the Under Secretary for Economic Affairs, Commerce Department.

[FR Doc. 2025-05496 Filed 3-28-25; 8:45 am]

BILLING CODE 3510-17-P

DEPARTMENT OF COMMERCE

International Trade Administration

[C-351-865]

Hard Empty Capsules From Brazil: Preliminary Affirmative Countervailing Duty Determination and Alignment of Final Determination With Final Antidumping Duty Determination

AGENCY: Enforcement and Compliance, International Trade Administration, Department of Commerce.

SUMMARY: The U.S. Department of Commerce (Commerce) preliminarily determines that countervailable subsidies are being provided to producers and exporters of hard empty capsules (capsules) from Brazil. The period of investigation is January 1, 2023, through December 31, 2023. Interested parties are invited to comment on this preliminary determination.

DATES: Applicable March 31, 2025. FOR FURTHER INFORMATION CONTACT: Samuel Evans, AD/CVD Operations, Office IX, Enforcement and Compliance, International Trade Administration, U.S. Department of Commerce, 1401 Constitution Avenue NW, Washington, DC 20230; telephone: (202) 482–2420.

SUPPLEMENTARY INFORMATION:

Background

On November 20, 2024, Commerce published the notice of initiation of this countervailing duty (CVD) investigation on capsules from Brazil.¹ On January 15, 2025, Commerce postponed the

preliminary determination of this investigation until March 24, 2025.² This preliminary determination is made in accordance with section 703(b) of the Tariff Act of 1930, as amended (the Act).

For a complete description of the events that followed the initiation of this investigation, see the Preliminary Decision Memorandum.³ A list of topics discussed in the Preliminary Decision Memorandum is included as Appendix II to this notice. The Preliminary Decision Memorandum is a public document and is on file electronically via Enforcement and Compliance's Antidumping and Countervailing Duty Centralized Electronic Service System (ACCESS). ACCESS is available to registered users at https://access. trade.gov. In addition, a complete version of the Preliminary Decision Memorandum can be accessed directly at https://access.trade.gov/public/ FRNoticesListLayout.aspx.

Scope of the Investigation

The products covered by this investigation are capsules from Brazil. For a complete description of the scope of this investigation, *see* Appendix I.

Scope Comments

In accordance with the Preamble to Commerce's regulations,4 the Initiation $Notice \ {\rm set} \ {\rm aside} \ {\rm a} \ {\rm period} \ {\rm of} \ {\rm time} \ {\rm for}$ parties to raise issues regarding product coverage (i.e., scope). 5 Certain interested parties commented on the scope of the investigation as it appeared in the Initiation Notice. For a summary of the scope comments and rebuttal responses submitted for this preliminary determination, and Commerce's accompanying preliminary analysis of all comments timely received, see the Preliminary Scope Decision Memorandum.⁶ Commerce is not preliminarily modifying the scope

¹ See Hard Empty Capsules from Brazil, the People's Republic of China, India, and the Socialist Republic of Vietnam: Initiation of Countervailing Duty Investigations, 89 FR 91680 (November 20, 2024) (Initiation Notice).

² See Hard Empty Capsules from Brazil, the People's Republic of China, India, and the Socialist Republic of Vietnam: Postponement of Preliminary Determinations in the Countervailing Duty Investigations, 90 FR 3788 (January 15, 2025).

³ See Memorandum, "Decision Memorandum for the Preliminary Determination of the Countervailing Duty Investigation of Hard Empty Capsules from Brazil," dated concurrently with, and hereby adopted by, this notice (Preliminary Decision Memorandum).

⁴ See Antidumping Duties; Countervailing Duties, Final Rule, 62 FR 27296, 27323 (May 19, 1997) (Preamble).

⁵ See Initiation Notice.

⁶ See Memorandum, "Less-Than-Fair-Value and Countervailing Duty Investigations of Hard Empty Capsules from Brazil, the People's Republic of China, India, and the Socialist Republic of Vietnam: Scope Comments Decision Memorandum for the Preliminary Determination," dated concurrently with, and hereby adopted by, this notice (Preliminary Scope Decision Memorandum).