Landsdown St., North Bethesda, MD 20852, 301–796–8867, *PRAStaff@fda.hhs.gov.*

SUPPLEMENTARY INFORMATION: Under the PRA (44 U.S.C. 3501–3521), Federal Agencies must obtain approval from the Office of Management and Budget (OMB) for each collection of information they conduct or sponsor. ''Collection of information'' is defined in 44 U.S.C. 3502(3) and 5 CFR 1320.3(c) and includes Agency requests or requirements that members of the public submit reports, keep records, or provide information to a third party. Section 3506(c)(2)(A) of the PRA (44 U.S.C. 3506(c)(2)(A)) requires Federal Agencies to provide a 60-day notice in the **Federal Register** concerning each proposed collection of information, including each proposed extension of an existing collection of information, before submitting the collection to OMB for approval. To comply with this requirement, FDA is publishing notice of the proposed collection of information set forth in this document.

With respect to the following collection of information, FDA invites comments on these topics: (1) whether the proposed collection of information is necessary for the proper performance of FDA's functions, including whether the information will have practical utility; (2) the accuracy of FDA's estimate of the burden of the proposed collection of information, including the validity of the methodology and assumptions used; (3) ways to enhance the quality, utility, and clarity of the information to be collected; and (4) ways to minimize the burden of the collection of information on respondents, including through the use of automated collection techniques, when appropriate, and other forms of information technology.

Temporary Marketing Permit Applications—21 CFR 130.17(c) and (i)

OMB Control Number 0910–0133—Extension

This information collection request supports FDA regulations found in 21 CFR 130.17 (section 130.17). Section 401 of the Federal Food, Drug, and Cosmetic Act (the FD&C Act) (21 U.S.C. 341) directs FDA to issue regulations establishing definitions and standards of identity (SOIs) for food. Under section 403(g) of the FD&C Act (21 U.S.C. 343(g)), a food that is subject to a definition and SOI prescribed by regulation is misbranded if it does not conform to such definition and SOI. Section 130.17 provides for the issuance by FDA of temporary marketing permits (TMPs) that enable the food industry to test consumer acceptance and measure the technological and commercial feasibility in interstate commerce of experimental packs of food that deviate from applicable definitions and SOIs. Section 130.17(c) enables the Agency to monitor the manufacture, labeling, and distribution of experimental packs of food that deviate from applicable definitions and SOIs. The information so obtained can be used in support of a petition to establish or amend the applicable definition or SOI to provide for the variations. Section 130.17(i) specifies the information that a firm must submit to FDA to obtain an extension of a TMP. To assist respondents with the TMP process, we have developed guidance entitled ''Temporary Permits for Interstate Shipment of Experimental Packs of Food Varying from the Requirements of Definitions and Standards of Identity: Guidance for Industry'' (November 2021). This resource can be found on our website *https://www.fda.gov/regulatory-information/search-fda-guidance-documents/guidance-industry-temporary-permits-interstate-shipment-experimental-packs-food-varying-requirements.*

Description of Respondents: Respondents to this collection of information include private sector businesses including institutional and/or industrial customers and food industry members such as manufacturers, packers, or distributors desiring to apply for a TMP or TMP extension.

We estimate the burden of this collection of information as follows:

TABLE 1—ESTIMATED ANNUAL REPORTING BURDEN [1]

21 CFR section; activity	Number of respondents	Number of responses per respondent	Total annual responses	Average burden per response	Total hours
130.17(c); Request for TMP	13	2	26	25	650
130.17(i); Request for TMP extension	1	2	2	2	4
Total					654

[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: July 12, 2023.

Lauren K. Roth,

Associate Commissioner for Policy.

[FR Doc. 2023–15045 Filed 7–14–23; 8:45 am]

BILLING CODE 4164–01–P

DEPARTMENT OF HEALTH AND HUMAN SERVICES

[Document Identifier: OS–0990–0379]

Agency Father Generic Information Collection Request: 30-Day Public Comment Request

AGENCY: Office of the Secretary, Health and Human Service, HHS.

ACTION: Notice and request for comments. Office of the Assistant Secretary for Public Affairs is requesting OMB approval for a new father Generic Clearance.

SUMMARY: In compliance with the requirement of the Paperwork Reduction Act of 1995, the Office of the Secretary (OS), Department of Health and Human Services, is publishing the following summary of a proposed collection for public comment.

DATES: Comments on the ICR must be received on or before August 16, 2023.

ADDRESSES: Submit your comments to *OIRA_submission@omb.eop.gov* or via facsimile to (202) 395–5806.

FOR FURTHER INFORMATION CONTACT: Sherrette Funn, *Sherrette.Funn@hhs.gov* or (202) 264–0041. When requesting information, please include the document identifier 0990–0379–30D and project title for reference.

SUPPLEMENTARY INFORMATION: Interested persons are invited to send comments

regarding this burden estimate or any other aspect of this collection of information, including any of the following subjects: (1) The necessity and utility of the proposed information collection for the proper performance of the agency's functions; (2) the accuracy of the estimated burden; (3) ways to enhance the quality, utility, and clarity of the information to be collected; and (4) the use of automated collection techniques or other forms of information technology to minimize the information collection burden.

Title of the Collection: Generic Clearance for the Collection of Qualitative Feedback on Agency Service Delivery (Online Customer Surveys).

Type of Collection: Extension.

OMB No.: 0990–0379—OS/ASPA.

Abstract: This collection of information is necessary to enable the Agency to garner customer and stakeholder feedback in an efficient, timely manner, in accordance with our commitment to improving service delivery. The information collected from our customers and stakeholders will help ensure that users have an effective, efficient, and satisfying experience with the Agency's programs. This feedback will provide insights into customer or stakeholder perceptions, experiences and expectations, provide an early warning of issues with service, or focus attention on areas where communication, training or changes in operations might improve delivery of products or services. These collections will allow for ongoing, collaborative and actionable communications between the Agency and its customers and stakeholders.

Type of respondent; frequency (annual, quarterly, monthly, etc.); and the affected public (individuals, public or private businesses, state or local governments, etc.).

ANNUALIZED BURDEN HOUR TABLE

Forms (if necessary)	Number of respondents	Number of responses per respondents	Average burden per response	Total burden hours
Website Customer Satisfaction Survey	3,000,000	1	10/60	500,000

Sherrette A. Funn,

Office of the Secretary, Paperwork Reduction Act Reports Clearance Officer.

[FR Doc. 2023–15030 Filed 7–14–23; 8:45 am]

BILLING CODE 4150–25–P

DEPARTMENT OF HEALTH AND HUMAN SERVICES

[Document Identifier: OS–0990–0459]

Agency Father Generic Information Collection Request: 30-Day Public Comment Request

AGENCY: Office of the Secretary, Health and Human Service, HHS.

ACTION: Notice and request for comments. Office of the Assistant Secretary for Public Affairs is requesting OMB approval for a new father Generic Clearance.

SUMMARY: In compliance with the requirement of the Paperwork Reduction Act of 1995, the Office of the Secretary (OS), Department of Health and Human Services, is publishing the following summary of a proposed collection for public comment.

DATES: Comments on the ICR must be received on or before August 16, 2023.

ADDRESSES: Submit your comments to *OIRA_submission@omb.eop.gov* or via facsimile to (202) 395–5806.

FOR FURTHER INFORMATION CONTACT: Sherrette Funn, *Sherrette.Funn@hhs.gov* or (202) 264–0041. When requesting information, please include the document identifier 0990–0459 30D and project title for reference.

SUPPLEMENTARY INFORMATION: Interested persons are invited to send comments regarding this burden estimate or any other aspect of this collection of information, including any of the following subjects: (1) The necessity and utility of the proposed information collection for the proper performance of the agency's functions; (2) the accuracy of the estimated burden; (3) ways to enhance the quality, utility, and clarity of the information to be collected; and (4) the use of automated collection techniques or other forms of information technology to minimize the information collection burden.

Title of the Collection: Fast-Track Generic Clearance for the Collection of Routine Customer Feedback on HHS Communications.

Type of Collection: Father Generic ICR.

OMB No.: 0990–0459—ASPA.

Abstract: This collection of information is necessary to enable HHS to garner customer and stakeholder feedback. Information will be collected from our customers and stakeholders from the concept phase to the end of the product life cycle. This will help ensure that users have an effective, efficient, and satisfying experience with HHS communications products. If this information is not collected, vital feedback on HHS communications will be unavailable, preventing programs from developing communications products that meets the needs of the audience and demonstrating impact of the communications products developed.

Type of respondent; frequency (annual, quarterly, monthly, etc); and the affected public (individuals, public or private businesses, state or local governments, etc.).

ANNUALIZED BURDEN HOUR TABLE

Forms (If necessary)	Number of respondents	Number of responses per respondents	Average burden per response	Total burden hours
HHS communications products	1,000,000	1	30/60	500,000