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**Donata Green,**

*Acting Associate Director, Office of Policy, Planning and Partnerships, Agency for Toxic Substances and Disease Registry.*

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**DEPARTMENT OF HEALTH AND HUMAN SERVICES**

**Food and Drug Administration**

[Docket No. FDA-2023-N-1053]

**Agency Information Collection Activities; Submission for Office of Management and Budget Review; Comment Request; Customer/Partner Service Satisfaction Surveys**

**AGENCY:** Food and Drug Administration, HHS.

**ACTION:** Notice.

**SUMMARY:** The Food and Drug Administration (FDA or Agency) 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 29, 2023.

**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 0910-0360. Also include the FDA docket number found in brackets in the heading of this document.

**FOR FURTHER INFORMATION CONTACT:**

JonnaLynn Capezzuto, Office of Operations, Food and Drug Administration, Three White Flint North, 10A-12M, 11601 Landsdown St., North Bethesda, MD 20852, 301-796-3794, [PRAStaff@fda.hhs.gov](mailto:PRAStaff@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.

**Customer/Partner Service Satisfaction Surveys**

*OMB Control Number 0910-0360—Extension*

Under section 1003 of the Federal Food, Drug, and Cosmetic Act (21 U.S.C. 393), FDA is authorized to conduct research and public information programs about regulated products and responsibilities of the Agency. Executive Order 12862, entitled “Setting Customer Service Standard,” directs Federal Agencies that “provide significant services directly to the public” to “survey customers to determine the kind and quality of services they want and their level of satisfaction with existing services.” FDA is seeking to extend OMB approval to conduct customer service satisfaction surveys to implement Executive Order 12862. Participation in the surveys is

voluntary. This request covers customer/partner (including State and local governments) service satisfaction surveys of regulated entities, such as food processors; cosmetic, drug, biologic, and medical device manufacturers; animal drugs, animal food and feed; tobacco products; and consumers and health professionals.

FDA will use the information from these surveys to identify strengths and weaknesses in service to customers/partners and to make improvements. The surveys will measure timeliness, appropriateness, clarity, and accuracy of information, courtesy, and problem resolution in the context of individual programs.

FDA estimates conducting approximately 20 customer/partner service satisfaction surveys per year, each requiring an average of 25 minutes for review and completion. We estimate respondents to these surveys to be between 100 and 20,000 customers/partners. Some of these surveys will be repeats of earlier surveys for purposes of monitoring customer/partner service and developing long-term data. Respondents to this collection of information cover a broad range of stakeholders who have experience with certain products regulated by or services provided by FDA.

In the **Federal Register** of April 25, 2023 (88 FR 24992), FDA published a 60-day notice requesting public comment on the proposed collection of information. One comment was received in support of this information collection.

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

TABLE 1—ESTIMATED ANNUAL REPORTING BURDEN <sup>1</sup>

Activity	Number of respondents	Number of responses per respondent	Total annual responses	Average burden per response	Total hours
Mail, telephone, web-based survey .....	85,000	1	85,000	0.42 (25 minutes) .....	35,700

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

Since the last OMB approval of this information collection request, FDA submitted three requests to increase the total burden hours. Therefore, this request for extension of OMB approval adjusts the number of respondents by an increase of 30,000 and the total burden hours by an increase of 21,950.

Dated: August 24, 2023.

**Lauren K. Roth,**

*Associate Commissioner for Policy.*

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**DEPARTMENT OF HEALTH AND HUMAN SERVICES**

**Health Resources and Services Administration**

**Meeting of the National Advisory Council on Migrant Health**

**AGENCY:** Health Resources and Services Administration (HRSA), Department of Health and Human Services.

**ACTION:** Notice.

**SUMMARY:** In accordance with the Federal Advisory Committee Act, this notice announces that the National Advisory Council on Migrant Health (NACMH) has scheduled a public meeting. Information about NACMH and the agenda for this meeting can be found on NACMH’s website at <https://www.hrsa.gov/advisory-committees/migrant-health>.

**DATES:** November 1–2, 2023, 9:00 a.m.–5:00 p.m. Eastern Standard Time.

**ADDRESSES:** This meeting will be held in-person at 5600 Fishers Lane,