a PFO initiative. HRSA considers PFO initiatives to be an innovative approach to funding home visiting service delivery, which may result in social benefit, as well as cost savings or cost avoidance to the public sector.

In response to the forthcoming SIR, MIECHV awardees planning to use MIECHV grant funds for outcomes or success payments related to a PFO initiative will be required to submit a PFO SIR Response outlining how their plans will meet all of the applicable statutory requirements and identifying what specific MIECHV funds (*e.g.,* fiscal year 2021 formula funding) they propose to use to (1) develop and implement their PFO initiative; and (2) make PFO outcomes or success payments based on the planned PFO initiative.

Regarding a PFO initiative, the MIECHV authorizing statute requires the following:

(1) A PFO initiative may not result in a reduction of funding for services delivered by the entity under a childhood home visitation program under this section while the eligible entity develops or operates such an initiative (section 711(c)(3)); and

(2) The PFO initiative for which outcome or success payments may be made must include:

(a) A feasibility study that describes how the proposed intervention is based on evidence of effectiveness;

(b) A rigorous, third-party evaluation that uses experimental or quasi-experimental design or other research methodologies that allow for the strongest possible causal inferences to determine whether the initiative has met its proposed outcomes as a result of implementation;

(c) An annual, publicly available report on the progress of the initiative; and

(d) A requirement that payments are made to the recipient of the grant, contract, or cooperative agreement only when agreed upon outcomes are achieved, excluding payments made to a third party conducting the evaluation. *See* 42 U.S.C. 711(k)(4).

The forthcoming SIR will provide further instructions to awardees in proposing a PFO initiative and submitting the required information to HRSA. Awardees are not required to propose or implement a PFO initiative, but if they wish to do so, they must submit a PFO SIR Response describing how their PFO initiative will meet all of the applicable statutory requirements. HRSA will use the information collected through the PFO SIR Response to ensure that MIECHV awardees proposals to use grant funds for PFO initiatives meet statutory requirements and to provide technical assistance to awardees. The implementation of a PFO initiative is not intended to disrupt current services or negatively impact communities that have benefited from home visiting programs and must not result in a reduction of funding for home visiting services.

Likely Respondents: MIECHV Program awardees that are states, territories, and, where applicable, nonprofit organizations providing home visiting services within states.

Burden Statement: Burden in this context means the time expended by persons to generate, maintain, retain, disclose, or provide the information requested. This includes the time needed to review instructions and supporting materials; to collect and analyze data and information to develop the PFO SIR Response; engage with stakeholders and coordinate with state level partners; and to draft and submit the PFO SIR Response. The table below summarizes the total annual burden hours estimated for this SIR.

TOTAL ESTIMATED ANNUALIZED BURDEN HOURS

Instrument	Number of respondents	Number of responses per respondent	Total responses	Average burden hours per response	Total burden hours
MIECHV PAY FOR OUTCOMES SIR	15	1	15	92	1,380
Total	15		15		1,380

HRSA specifically requests comments on (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.

Maria G. Button,

Director, Executive Secretariat.

[FR Doc. 2020–14658 Filed 7–7–20; 8:45 am]

BILLING CODE 4165–15–P

DEPARTMENT OF HEALTH AND HUMAN SERVICES

[Document Identifier OS–0990–0379]

Agency Information Collection Request: 30-Day Public Comment Request

AGENCY: Office of the Secretary, HHS.

ACTION: Notice.

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 7, 2020.

ADDRESSES: Written comments and recommendations for the proposed information collection should be sent within 30 days of publication of this notice to *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.

FOR FURTHER INFORMATION CONTACT: Sherrette Funn, *Sherrette.Funn@hhs.gov* or (202) 795–7714. When submitting comments or requesting information, please include the document identifier 0990-New-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: Father Generic ICR.

OMB No. 0990–0379—Office within OS—Specific program collecting the data (is applicable)

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.) (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

Terry Clark,

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

[FR Doc. 2020–14595 Filed 7–7–20; 8:45 am]

BILLING CODE 4150–25–P

DEPARTMENT OF HEALTH AND HUMAN SERVICES

National Institutes of Health

National Institute of Diabetes and Digestive and Kidney Diseases; Notice of Closed Meeting

Pursuant to section 10(d) of the Federal Advisory Committee Act, as amended, notice is hereby given of the following meeting.

The meeting will be closed to the public in accordance with the provisions set forth in sections 552b(c)(4) and 552b(c)(6), Title 5 U.S.C., as amended. The grant applications and the discussions could disclose confidential trade secrets or commercial property such as patentable material, and personal information concerning individuals associated with the grant applications, the disclosure of which would constitute a clearly unwarranted invasion of personal privacy.

Name of Committee: National Institute of Diabetes and Digestive and Kidney Diseases Special Emphasis Panel PAR19–202: High impact, Interdisciplinary Science in NIDDK Research Areas (RC2 Clinical Trial Optional)—Kidney and Urological Diseases.

Date: September 1, 2020.

Time: 11:00 a.m. to 2:30 p.m.

Agenda: To review and evaluate grant applications.

Place: National Institutes of Health, Two Democracy Plaza, 6707 Democracy Boulevard, Bethesda, MD 20892, (Video Meeting).

Contact Person: Najma S. Begum, Ph.D., Scientific Review Officer, Review Branch, DEA, NIDDK, National Institutes of Health, Room 7349, 6707 Democracy Boulevard, Bethesda, MD 20892–5452, (301) 594–8894, *begumn@niddk.nih.gov*.

(Catalogue of Federal Domestic Assistance Program Nos. 93.847, Diabetes, Endocrinology and Metabolic Research; 93.848, Digestive Diseases and Nutrition Research; 93.849, Kidney Diseases, Urology and Hematology Research, National Institutes of Health, HHS)

Dated: July 2, 2020.

Miguelina Perez,

Program Analyst, Office of Federal Advisory Committee Policy.

[FR Doc. 2020–14708 Filed 7–7–20; 8:45 am]

BILLING CODE 4140–01–P

DEPARTMENT OF HEALTH AND HUMAN SERVICES

National Institutes of Health

Center for Scientific Review; Notice of Closed Meeting

Pursuant to section 10(d) of the Federal Advisory Committee Act, as amended, notice is hereby given of the following meeting.

The meeting will be closed to the public in accordance with the provisions set forth in sections 552b(c)(4) and 552b(c)(6), Title 5 U.S.C., as amended. The grant applications and the discussions could disclose confidential trade secrets or commercial property such as patentable material, and personal information concerning individuals associated with the grant applications, the disclosure of which would constitute a clearly unwarranted invasion of personal privacy.

Name of Committee: Center for Scientific Review Special Emphasis Panel Small Business: Microbial (non-HIV) Diagnostics and Detection of Infectious Agents, Food and Waterborne Pathogens, and Methods in Microbial Sterilization, Disinfection and Bioremediation.

Date: July 15, 2020.

Time: 12:00 p.m. to 5:30 p.m.

Agenda: To review and evaluate grant applications.

Place: National Institutes of Health, Rockledge II, 6701 Rockledge Drive, Bethesda, MD 20892, (Virtual Meeting).

Contact Person: Gagan Pandya, Ph.D., Scientific Review Officer, National Institutes of Health, Center for Scientific Review, 6701 Rockledge Drive, RM 3200, MSC 7808, Bethesda, MD 20892, 301–435–1167, *pandyaga@mail.nih.gov*.

This notice is being published less than 15 days prior to the meeting due to the timing limitations imposed by the review and funding cycle.

(Catalogue of Federal Domestic Assistance Program Nos. 93.306, Comparative Medicine; 93.333, Clinical Research, 93.306, 93.333, 93.337, 93.393–93.396, 93.837–93.844, 93.846–93.878, 93.892, 93.893, National Institutes of Health, HHS)

Dated: July 2, 2020.

Melanie J. Pantoja,

Program Analyst, Office of Federal Advisory Committee Policy.

[FR Doc. 2020–14709 Filed 7–7–20; 8:45 am]

BILLING CODE 4140–01–P