Guatemala. This award will build upon previously funded Global Health Security projects, and specifically surveillance activities by CDC to collaboratively work with host countries to create sustainable systems that can inform local public health policy.

DATES: The period for this award will be September 30, 2024, through September 29, 2029.

FOR FURTHER INFORMATION CONTACT: Shana Eatman, Centers for Disease Control and Prevention, 1600 Clifton Rd., Atlanta, GA, Telephone: 770–488–3933, Email: *DGHPNOFOs@cdc.gov.*

SUPPLEMENTARY INFORMATION: The single source award will build on existing public health infrastructure and CDC-supported surveillance and workforce development platform in the region. This award will enable CDC to work collaboratively with recipients to sustain its Global Health Security Agenda (GHSA) efforts to strengthen health protection, detection, and response capacity in Guatemala/Central America. Universidad del Valle de Guatemala is in a unique position to conduct this work, as it will focus on multi-pathogen key public health impact disease surveillance of respiratory pathogen surveillance, support for entomological activities and public health workforce development.

Summary of the Award

Recipient: Universidad del Valle de Guatemala.

Purpose of the Award: The purpose of this award is to enhance global health security in Guatemala/CA through epidemiologic, clinical, and laboratory-based surveillance. This award should contribute to strengthening national notifiable disease surveillance and translating data into public health action.

Amount of Award: $5,000,000 in Federal Fiscal Year (FFY) 2024 funds, with a total estimated $25,000,000 for the 5-year period of performance, subject to availability of funds. Please note, Year 1 funding is as follows: $5,000,000 for Core Component 1, $10,000,000 for Approved but Unfunded (ABU) Component 2, $15,000,000 Component 3 (ABU), and $15,000,000 for (ABU) Component 4.

Period of Performance: September 30, 2024, through September 29, 2029.

Authority: This program is authorized under section 307 of the Public Health Service Act [42 U.S.C. 242*l*] and section 317(k)(2) of the Public Health Service Act [42 U.S.C. 247(b)(k2].

Dated: May 21, 2024.

Jamie Legier,

Acting Director, Office of Grants Services, Centers for Disease Control and Prevention.

[FR Doc. 2024–11725 Filed 5–28–24; 8:45 am]

BILLING CODE 4163–18–P

DEPARTMENT OF HEALTH AND HUMAN SERVICES

Centers for Medicare & Medicaid Services

[Document Identifiers: CMS–2540–24]

Agency Information Collection Activities: Submission for OMB Review; Comment Request

AGENCY: Centers for Medicare & Medicaid Services, Health and Human Services (HHS).

ACTION: Notice.

SUMMARY: The Centers for Medicare & Medicaid Services (CMS) is announcing an opportunity for the public to comment on CMS' intention to collect information from the public. Under the Paperwork Reduction Act of 1995 (PRA), federal agencies are required to publish notice in the **Federal Register** concerning each proposed collection of information, including each proposed extension or reinstatement of an existing collection of information, and to allow a second opportunity for public comment on the notice. Interested persons are invited to send comments regarding the burden estimate or any other aspect of this collection of information, including the necessity and utility of the proposed information collection for the proper performance of the agency's functions, the accuracy of the estimated burden, ways to enhance the quality, utility, and clarity of the information to be collected, and the use of automated collection techniques or other forms of information technology to minimize the information collection burden.

DATES: Comments on the collection(s) of information must be received by the OMB desk officer by June 28, 2024.

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.

To obtain copies of a supporting statement and any related forms for the proposed collection(s) summarized in this notice, please access the CMS PRA website by copying and pasting the following web address into your web browser: *https://www.cms.gov/Regulations-and-Guidance/Legislation/PaperworkReductionActof1995/PRA-Listing.*

FOR FURTHER INFORMATION CONTACT: William Parham at (410) 786–4669.

SUPPLEMENTARY INFORMATION: Under the Paperwork Reduction Act of 1995 (PRA) (44 U.S.C. 3501–3520), federal agencies must obtain approval from the Office of Management and Budget (OMB) for each collection of information they conduct or sponsor. The term ''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 publish a 30-day notice in the **Federal Register** concerning each proposed collection of information, including each proposed extension or reinstatement of an existing collection of information, before submitting the collection to OMB for approval. To comply with this requirement, CMS is publishing this notice that summarizes the following proposed collection(s) of information for public comment:

1. *Type of Information Collection Request:* Reinstatement with change of a previously approved collection; *Title of Information Collection:* Skilled Nursing Facility and Skilled Nursing Facility Complex Cost Report; *Use:* The primary function of the cost report is to implement the principles of cost reimbursement that require that SNFs maintain sufficient financial records and statistical data for proper determination of costs payable under the program. Specifically, CMS–2540–24 collects discrete data, previously reported in summary form, used in determining the cost weights for the SNF market basket and for payment adequacy analyses. SNFs and SNF health care complexes participating in the Medicare program submit these cost reports annually to report cost and statistical data used by CMS to determine reasonable costs. Essentially the methods of determining costs payable under Medicare involve making use of data available from the provider's accounting records, as usually maintained, to arrive at equitable and proper payment for services to beneficiaries.

We originally published the 30-day notice on April 17, 2024 (89 FR 27413); however, there was a significant delay

between the publication of the notice and the associated files being publicly available. For this reason, we are publishing the 30-day notice again. *Form Number:* CMS–2540–24 (OMB control number: 0938–0463); *Frequency:* Annually; *Affected Public:* Private Sector—Business or other for-profits, Not-for-profit institutions; *Number of Respondents:* 14,189; *Total Annual Responses:* 14,189; *Total Annual Hours:* 2,866,178. (For policy questions regarding this collection contact Luann Piccione at 410–786–5423.)

William N. Parham, III,

Director, Division of Information Collections and Regulatory Impacts, Office of Strategic Operations and Regulatory Affairs.

[FR Doc. 2024–11741 Filed 5–28–24; 8:45 am]

BILLING CODE 4120–01–P

DEPARTMENT OF HEALTH AND HUMAN SERVICES

Administration for Children and Families

Tribal Consultation Meetings

AGENCY: Office of Early Childhood Development (ECD), Office of Head Start (OHS), and Office of Child Care (OCC), Administration for Children and Families (ACF), U.S. Department of Health and Human Services (HHS).

ACTION: Notice of meetings.

SUMMARY: Pursuant to the Head Start Act, notice is hereby given of one joint Tribal consultation session to be held between HHS/ACF ECD, OHS, and OCC leadership and the leadership of Tribal governments operating Tribal Maternal, Infant, and Early Childhood Home Visiting; Tribal Child Care and Development Fund; and Head Start and Early Head Start programs. The purpose of this consultation session is to discuss ways to better meet the needs of Tribal children and their families and issues affecting the delivery of early childhood services in their geographic locations. The consultation will also provide an opportunity for discussion on the review and promulgation of Head Start Program Performance Standards, as required under the Head Start Act. To meet this legislative requirement, one Tribal consultation will be held as part of HHS/ACF or ACF Tribal consultation sessions.

DATES: The meetings will take place July 9 and 10, 2024, at the following times:

- July 9 from 9 a.m.–5 p.m. Mountain Standard Time; and
- July 10 from 9 a.m.–3 p.m. Mountain Standard Time

ADDRESSES: The meetings are in-person in Fort McDowell, Arizona, at:

We-Ko-Pa Casino Resort, 10438 WeKoPa Way, Fort McDowell, AZ 85264

FOR FURTHER INFORMATION CONTACT: Todd Lertjuntharangool, Regional Program Manager, Region 11, Office of Head Start, email *Todd.Lertjuntharangool@acf.hhs.gov,* or phone (866) 763–6481. Additional information and online meeting registration will be forthcoming.

SUPPLEMENTARY INFORMATION: In accordance with section 640(l)(4) of the Head Start Act, 42 U.S.C. 9835(1)(4), ACF announces a joint Tribal consultation session to be held between HHS/ACF ECD, OHS, and OCC leadership and the leadership of Tribal governments operating Tribal Maternal, Infant, and Early Childhood Home Visiting; Tribal Child Care and Development Fund; and Head Start and Early Head Start programs.

The agenda for the scheduled joint consultation reflects the statutory purposes of Head Start Tribal consultations related to meeting the needs of American Indian and Alaska Native (AIAN) children and families and will include the opportunity to discuss topics such as the Tribal Request for Information and recent changes to AIAN Head Start eligibility requirements. The consultation will also provide an opportunity for discussion on the review and promulgation of Head Start Program Performance Standards under section 641A(a)(2)(D) of the Head Start Act. OHS will also highlight the progress made in addressing issues and concerns raised in the previous OHS Tribal consultations.

The consultation session includes elected or appointed leaders of Tribal governments and their designated representatives. Designees must have a letter from the Tribal government authorizing them to represent the Tribe. Tribal governments must submit the designee letter at least 3 days before the consultation session to *AIANheadstart@acf.hhs.gov*. Other representatives of Tribal organizations and Native non-profit organizations are welcome to attend as observers.

Within 45 days of the consultation session, a detailed report of each consultation session will be available for all Tribal governments receiving funds for Head Start and Early Head Start programs. Tribes can submit written testimony for the report to *AIANheadstart@acf.hhs.gov* prior to each consultation session or within 30 days of each meeting. OHS will summarize oral testimony and comments from the consultation sessions in each report without attribution, along with topics of concern and recommendations.

Khari M. Garvin,

Director, Office of Head Start.

Megan E. Steel,

ACF Certifying Officer.

[FR Doc. 2024–11783 Filed 5–24–24; 11:15 am]

BILLING CODE 4184–40–P

DEPARTMENT OF HEALTH AND HUMAN SERVICES

Food and Drug Administration

[Docket No. FDA–2024–N–2149]

Agency Information Collection Activities; Proposed Collection; Comment Request; De Novo Classification Process (Evaluation of Automatic Class III Designation)

AGENCY: Food and Drug Administration, HHS.

ACTION: Notice.

SUMMARY: The Food and Drug Administration (FDA or Agency) is announcing an opportunity for public comment on the proposed collection of certain information by the Agency. Under the Paperwork Reduction Act of 1995 (PRA), Federal Agencies are required to publish notice in the **Federal Register** concerning each proposed collection of information, including each proposed extension of an existing collection of information, and to allow 60 days for public comment in response to the notice. This notice solicits comments on the information collection provisions related to the De Novo Classification Process.

DATES: Either electronic or written comments on the collection of information must be submitted by July 29, 2024.

ADDRESSES: You may submit comments as follows. Please note that late, untimely filed comments will not be considered. The *https://www.regulations.gov* electronic filing system will accept comments until 11:59 p.m. Eastern Time at the end of July 29, 2024. Comments received by mail/hand delivery/courier (for written/paper submissions) will be considered timely if they are received on or before that date.

Electronic Submissions

Submit electronic comments in the following way:

- *Federal eRulemaking Portal: https://www.regulations.gov.* Follow the instructions for submitting comments. Comments submitted electronically,