

**DATES:** The meeting will be held on October 20–21, 2021, from 10:00 a.m. to 5:00 p.m., EDT (times subject to change). The docket is currently open to receive written comments. Written comments must be received on or before October 21, 2021.

**SUPPLEMENTARY INFORMATION:** Notice is hereby given of a change in the meeting of the Advisory Committee on Immunization Practices (ACIP); October 20, 2021, 10:00 a.m.–5:00 p.m., EDT, and October 21, 2021, 10:00 a.m.–5:00 p.m., EDT (times subject to change), in the original FRN.

The virtual meeting was published in the **Federal Register** on Wednesday, September 22, 2021, Volume 86, Number 181, pages 52683–52684.

The virtual meeting is being amended to change the date the docket was opened to receive written public comments, and updates to the Matters To Be Considered and Written Public Comment sections of the notice and should read as follows:

**DATES:** The meeting will be held on October 20–21, 2021, from 10:00 a.m. to 5:00 p.m., EDT (times subject to change). The public may submit written comments from October 7, 2021 through October 21, 2021.

*Matters To Be Considered:* The agenda will include discussions on herpes zoster vaccines, influenza vaccines, pneumococcal vaccine, and COVID–19 vaccines. Recommendation votes on herpes zoster vaccine, pneumococcal vaccine, and COVID–19 vaccines are scheduled. No Vaccines for Children (VFC) votes are scheduled. Agenda items are subject to change as priorities dictate. For more information on ACIP please visit the ACIP website: <http://www.cdc.gov/vaccines/acip/index.html>.

*Written Public Comment:* The docket will be opened to receive written comments on October 7, 2021. Written comments must be received on or before October 21, 2021.

The virtual meeting is open to the public.

**FOR FURTHER INFORMATION CONTACT:** Stephanie Thomas, ACIP Committee Management Specialist, Centers for Disease Control and Prevention, National Center for Immunization and Respiratory Diseases, 1600 Clifton Road NE, MS–H24–8, Atlanta, GA 30329–4027; Telephone: 404–639–8367; Email: [ACIP@cdc.gov](mailto:ACIP@cdc.gov).

The Director, Strategic Business Initiatives Unit, Office of the Chief Operating Officer, Centers for Disease Control and Prevention, has been delegated the authority to sign **Federal Register** notices pertaining to announcements of meetings and other

committee management activities, for both the Centers for Disease Control and Prevention and the Agency for Toxic Substances and Disease Registry.

**Kalwant Smagh,**

*Director, Strategic Business Initiatives Unit, Office of the Chief Operating Officer, Centers for Disease Control and Prevention.*

[FR Doc. 2021–24320 Filed 11–5–21; 8:45 am]

**BILLING CODE 4163–18–P**

## DEPARTMENT OF HEALTH AND HUMAN SERVICES

### Centers for Disease Control and Prevention

#### 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, and the Determination of the Director, Strategic Business Initiatives Unit, Office of the Chief Operating Officer, CDC, pursuant to Public Law 92–463. 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:* Disease, Disability, and Injury Prevention and Control Special Emphasis Panel (SEP)—RFA–CK–22–003, Emerging Infections Sentinel Networks (EISN) Research.

*Date:* January 11, 2022.

*Time:* 10:00 a.m.–5:00 p.m., EST.

*Place:* Teleconference, Centers for Disease Control and Prevention, Room 1080, 8 Corporate Square Boulevard, Atlanta, Georgia 30329–4027.

*Agenda:* To review and evaluate grant applications.

*For Further Information Contact:* Gregory Anderson, M.S., M.P.H., Scientific Review Officer, National Center for HIV/AIDS, Viral Hepatitis, STD, and TB Prevention, CDC, 1600 Clifton Road NE, Mailstop US8–1, Atlanta, Georgia 30329–4027, Telephone: (404) 718–8833, Email: [GAnderson@cdc.gov](mailto:GAnderson@cdc.gov).

The Director, Strategic Business Initiatives Unit, Office of the Chief Operating Officer, Centers for Disease Control and Prevention, has been delegated the authority to sign **Federal Register** notices pertaining to announcements of meetings and other committee management activities, for both the Centers for Disease Control and

Prevention and the Agency for Toxic Substances and Disease Registry.

**Kalwant Smagh,**

*Director, Strategic Business Initiatives Unit, Office of the Chief Operating Officer, Centers for Disease Control and Prevention.*

[FR Doc. 2021–24322 Filed 11–5–21; 8:45 am]

**BILLING CODE 4163–18–P**

## DEPARTMENT OF HEALTH AND HUMAN SERVICES

### Centers for Medicare & Medicaid Services

[Document Identifiers: CMS–10280]

#### 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 December 8, 2021.

**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](http://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, you may make your request using one of following:

1. Access CMS' website address at website address at: <https://www.cms.gov/Regulations-and-Guidance/Legislation/PaperworkReductionActof1995/PRA-Listing.html>.

**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:* Extension of a currently approved collection; *Title of the Information Collection:* Home Health Change of Care Notice; *Use:* The purpose of the Home Health Change of Care Notice (HHCCN) is to notify original Medicare beneficiaries receiving home health care benefits of plan of care changes. Home health agencies (HHAs) are required to provide written notice to Original Medicare beneficiaries under various circumstances involving the reduction or termination of items and/or services consistent with Home Health Agencies Conditions of Participation (COPs).

The home health COP requirements are set forth in § 1891 [42 U.S.C. 1395bbb] of the Social Security Act (the Act). The implementing regulations under 42 CFR 484.10(c) specify that Medicare patients receiving HHA

services have rights. The patient has the right to be informed, in advance about the care to be furnished, and of any changes in the care to be furnished. The HHA must advise the patient in advance of the disciplines that will furnish care, and the frequency of visits proposed to be furnished. The HHA must advise the patient in advance of any change in the plan of care before the change is made."

Notification is required for covered and non-covered services listed in the plan of care (POC). The beneficiary will use the information provided to decide whether or not to pursue alternative options to continue receiving the care noted on the HHCCN. *Form Number:* CMS-10280 (OMB control number: 0938-1196); *Frequency:* Yearly; *Affected Public:* Private Sector (Business or other for-profits, Not-for-Profit Institutions); *Number of Respondents:* 11,157; *Total Annual Responses:* 12,385,108; *Total Annual Hours:* 824,848. (For policy questions regarding this collection contact Jennifer McCormick at 410-786-2852.)

Dated: November 3, 2021.

**William N. Parham, III,**  
*Director, Paperwork Reduction Staff, Office of Strategic Operations and Regulatory Affairs.*

[FR Doc. 2021-24396 Filed 11-5-21; 8:45 am]

**BILLING CODE 4120-01-P**

## DEPARTMENT OF HEALTH AND HUMAN SERVICES

### Administration for Children and Families

#### Proposed Information Collection Activity; Notification of Obligation Target Status for CCDF American Rescue Plan (ARP) Act Stabilization Funds (0970-0510)

**AGENCY:** Office of Child Care (OCC), Administration for Children and Families, HHS.

**ACTION:** Request for public comment.

**SUMMARY:** The Administration for Children and Families (ACF), Office of Child Care (OCC) plans to submit a generic information collection (GenIC) request under the umbrella generic: Generic Clearance for Financial Reports used for ACF Mandatory Grant Programs (0970-0510). This request includes an information collection for

Child Care and Development Fund (CCDF) state and territory grant recipients to report obligation progress of the American Rescue Plan (ARP) Act Stabilization funds.

**DATES:** *Comments due within 14 days of publication.* In compliance with the requirements of the Paperwork Reduction Act of 1995, ACF is soliciting public comment on the specific aspects of the information collection described above and below.

**ADDRESSES:** Copies of the proposed collection of information can be obtained and comments may be submitted by emailing [infocollection@acf.hhs.gov](mailto:infocollection@acf.hhs.gov). All requests should be identified by the title of the information collection.

#### SUPPLEMENTARY INFORMATION:

*Description:* ACF programs require detailed financial information from their grantees that allows ACF to monitor various specialized cost categories within each program, to closely manage program activities, and to have sufficient financial information to enable periodic thorough and detailed audits. The Generic Clearance for Financial Reports used for ACF Mandatory Grant Programs allows ACF programs to efficiently develop and receive approval for financial reports that are tailored to specific funding recipients and the associated needs of the program. For more information about the umbrella generic, see: [https://www.reginfo.gov/public/do/PRAViewDocument?ref\\_nbr=202108-0970-002](https://www.reginfo.gov/public/do/PRAViewDocument?ref_nbr=202108-0970-002).

This specific GenIC is to meet the one-time statutory financial reporting requirement established by the ARP Act (Pub. L. 117-2, Sec. 2202). The ARP Act allocated \$24 billion for CCDF for lead agencies to award subgrants to child care providers in order to stabilize the child care market. The ARP Act requires lead agencies to notify the Secretary of HHS if they are unable to obligate at least 50 percent of the Stabilization funds that are available for subgrants within 9 months of enactment.

Generic clearance approval is requested to allow ACF's OCC to collect the necessary information from CCDF lead agencies by the statutory deadline of December 11, 2021.

*Respondents:* State and territory CCDF administrators.