currently approved through May 31, 2023. An Agency may not conduct or sponsor and a person is not required to respond to a collection of information unless it displays a currently valid OMB control number.

DATES: Comments must be submitted on or before January 30, 2023.

ADDRESSES: Submit your comments, referencing Docket ID No. EPA–HQ–OECA–2013–0547, online using *www.regulations.gov* (our preferred method), by email to *docket.oeca@epa.gov,* or by mail to: EPA Docket Center, Environmental Protection Agency, Mail Code 28221T, 1200 Pennsylvania Ave. NW, Washington, DC 20460.

EPA's policy is that all comments received will be included in the public docket without change including any personal information provided, unless the comment includes profanity, threats, information claimed to be Confidential Business Information (CBI) or other information whose disclosure is restricted by statute.

FOR FURTHER INFORMATION CONTACT: Gregory Savitske, Monitoring, Assistance, and Media Programs Division, Office of Compliance, (2227A), Environmental Protection Agency, 1200 Pennsylvania Ave. NW, Washington, DC 20460; telephone number: (202) 564–2601; fax number: (202) 564–0050; email address: *Savitske.Gregory@epa.gov.*

SUPPLEMENTARY INFORMATION: Supporting documents which explain in detail the information that the EPA will be collecting are available in the public docket for this ICR. The docket can be viewed online at *www.regulations.gov* or in person at the EPA Docket Center, WJC West, Room 3334, 1301 Constitution Ave. NW, Washington, DC. The telephone number for the Docket Center is 202–566–1744. For additional information about EPA's public docket, visit *https://www.epa.gov/dockets.*

Pursuant to section 3506(c)(2)(A) of the Paperwork Reduction Act, EPA is soliciting comments and information to enable it to: (i) evaluate whether the proposed collection of information is necessary for the proper performance of the functions of the Agency, including whether the information will have practical utility; (ii) evaluate the accuracy of the Agency's estimate of the burden of the proposed collection of information, including the validity of the methodology and assumptions used; (iii) enhance the quality, utility, and clarity of the information to be collected; and (iv) minimize the burden of the collection of information on those who are to respond, including through the use of appropriate automated electronic, mechanical, or other technological collection techniques or other forms of information technology, *e.g.,* permitting electronic submission of responses. EPA will consider the comments received and amend the ICR as appropriate. The final ICR package will then be submitted to OMB for review and approval. At that time, EPA will issue another **Federal Register** notice to announce the submission of the ICR to OMB and the opportunity to submit additional comments to OMB.

Abstract: Discharge Monitoring Report-Quality Assurance (DMR–QA) study program participation is mandatory for Major and selected Minor National Pollutant Discharge Elimination System (NPDES) permit holders in accordance with Clean Water Act Section 308. The DMR–QA study program is designed to evaluate the analytic ability of laboratories that perform chemical, microbiological and whole effluent toxicity (WET) analyses required in NPDES permits for reporting results in the Discharge Monitoring Reports (DMR). Under DMR–QA, the permit holder is responsible for having their in-house and/or contract laboratories analyze proficiency test samples and submit results to proficiency testing (PT) providers for grading. Graded results are transmitted by either the permit holder or PT provider to the appropriate federal or state NPDES permitting authority. Permit holders are responsible for submitting corrective action reports to the appropriate permitting authority.

Form Numbers: 6400–01.

Respondents/affected entities: Major and selected Minor permit holders under the Clean Water Act's National Pollutant Discharge Elimination System (NPDES).

Respondent's obligation to respond: Major permit holders must participate annually. Minor permit holders must participate if selected by the state or EPA DMR–QA coordinator.

Estimated number of respondents: 5,500 (total).

Frequency of response: Major permit holders must participate annually. Minor permit holders must participate if selected by the state or EPA DMR–QA coordinator.

Total estimated burden: 36,300 hours (per year). Burden is defined at 5 CFR 1320.03(b).

Total estimated cost: $5,240,070 (per year), includes $3,243,350 annualized capital or operation & maintenance costs.

Changes in estimates: The total estimated respondent burden is projected to remain the same as the ICR currently approved by OMB; this is attributed to the estimated number of respondents receiving this ICR remaining stable over the past three years. Labor costs will likely increase to account for changes in employee benefit and compensation costs as well as inflation. Non-labor costs for obtaining proficiency test samples will also likely increase.

Dated: November 23, 2022.

Elizabeth Vizard,

Acting Director, Monitoring, Assistance, and Media Programs Division, Office of Compliance.

[FR Doc. 2022–26087 Filed 11–29–22; 8:45 am]

BILLING CODE 6560–50–P

DEPARTMENT OF HEALTH AND HUMAN SERVICES

Centers for Disease Control and Prevention

[Docket No. CDC–2022–0136]

Advisory Committee on Immunization Practices; Cancellation of Meeting

AGENCY: Centers for Disease Control and Prevention (CDC), Department of Health and Human Services (HHS).

ACTION: Notice.

SUPPLEMENTARY INFORMATION: Notice is hereby given of a change in the meeting of the Advisory Committee on Immunization Practices (ACIP); December 9, 2022, from 10 a.m. to 5 p.m., EST. The virtual meeting was published in the **Federal Register** on November 23, 2022, Volume 87, Number 225, pages 71641–71642. This meeting is being canceled in its entirety.

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, Mailstop H24–8, Atlanta, Georgia 30329–4027; Telephone: 404–639–8836; Email: *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. 2022–26084 Filed 11–29–22; 8:45 am]

BILLING CODE 4163–18–P

DEPARTMENT OF HEALTH AND HUMAN SERVICES

Administration for Community Living

[OMB Control No. 0985–0029]

Agency Information Collection Activities: Proposed Collection; Public Comment Request of the State Councils on Developmental Disabilities (Councils) State Plan

AGENCY: Administration for Community Living, Department of Health and Human Services.

ACTION: Notice.

SUMMARY: The Administration for Community Living (ACL) is announcing an opportunity for the public to comment on the proposed collection of information listed above. Under the Paperwork Reduction Act of 1995 (PRA), Federal agencies are required to publish a 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 Information Collection (IC) Revision solicits comments on the information collection requirements relating to the Developmental Disabilities State Plan OMB control number 0985–0029.

DATES: Comments on the collection of information must be submitted electronically by 11:59 p.m. (EST) or postmarked by January 30, 2023.

ADDRESSES: Submit comments on the collection of information via email to *Sara.Newell-Perez@acl.hhs.gov* or to Administration for Community Living, 330 C Street SW, Washington, DC 20201, Attention: Sara Newell-Perez.

FOR FURTHER INFORMATION CONTACT: Sara Newell-Perez, 202–795–7413 or *Sara.Newell-Perez@acl.hhs.gov.*

SUPPLEMENTARY INFORMATION: Under the 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. ''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. The PRA 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, ACL is publishing a notice of the proposed collection of information set forth in this document.

With respect to the following collection of information, ACL invites comments on our burden estimates or any other aspect of this collection of information, including:

(1) whether the proposed collection of information is necessary for the proper performance of ACL's functions, including whether the information will have practical utility;

(2) the accuracy of ACL's estimate of the burden of the proposed collection of information, including the validity of the methodology and assumptions used to determine burden estimates;

(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.

The State Councils on Developmental Disabilities (Councils) are authorized in Subtitle B, of the Developmental Disabilities Assistance and Bill of Rights Act of 2000 (DD Act), as amended, [42 U.S.C. 15001 *et seq.*] (The DD Act). The DD Act requires Councils to submit a five-year State plan. Section 124(a) [42 U.S.C. 15024(a)], states that: *Any State desiring to receive assistance under this subtitle shall submit to the Secretary, and obtain approval of, a 5-year strategic State plan under this section.* The DD Act regulations outlines additional guiding requirements in 45 CFR part 1326.30(a), which states that: *In order to receive Federal financial assistance under this subpart, each State Developmental Disabilities Council must prepare and submit to the Secretary, and have in effect, a State plan which meets the requirements of sections 122 and 124 of the Act (42 U.S.C. 6022 and 6024) and these regulations.*

The Council is responsible for the development, and submission of the State plan as well as implementation of the activities described in the plan. The Council updates the Sate plan annually during the five years. The State plan provides information on individuals with developmental disabilities in the State, and a description of the services available to them and their families. The State plan sets forth the goals and specific objectives to be achieved by the State Council in pursuing systems change and capacity building that result in empowering people with developmental disabilities to lead independent lives within the community. It describes State priorities, strategies, and actions, and the allocation of funds to meet these goals and objectives. Additionally, the data collected in the State plan and submitted to ACL is also used to comply with the GPRA Modernization Act of 2010 (GPRAMA).

The State Plan is used in three ways. First, it provides a framework for citizens, State governments, and other key stakeholder to provide input and comments to help shape the goals and objectives during the development stage. Secondly, it is used by each Council as a planning document to operationalize its goals and strategies. Finally, it provides information the Department needs for monitoring and providing technical assistance to ensure the Council is compliant.

This is a revision of a currently approved information collect that expires March 30, 2023. To ensure the DD Council State plan is consistent with the Executive Order on Advancing Racial Equity and Support for Underserved Communities Through the Federal Government and the Executive Order on Advancing Equality for Lesbian, Gay, Bisexual, Transgender, Queer, and Intersex Individuals, ACL intends to determine whether sexual orientation and gender identity (SOGI) data elements need to be adapted prior to adding them to ensure accessibility of the questions for individuals with intellectual and developmental disabilities.

The proposed data collection tool may be found on the ACL website for review at: *https://www.acl.gov/about-acl/public-input.*

Estimated Program Burden: ACL estimates the burden of this collection of information as follows: