

and improved testing approaches applicable to the needs of U.S. federal agencies. NICEATM and ICCVAM welcome the public nomination of new, revised, and alternative test methods and strategies for validation studies and technical evaluations. Additional information about NICEATM can be found at <http://ntp.niehs.nih.gov/go/niceatm>.

SACATM, established by the ICCVAM Authorization Act [Section 285l–3(d)], provides advice on priorities and activities related to the development, validation, scientific review, regulatory acceptance, implementation, and national and international harmonization of new, revised, and alternative toxicological test methods to ICCVAM, NICEATM, and Director of NIEHS and NTP. Additional information about SACATM, including the charter, roster, and records of past meetings, can be found at <http://ntp.niehs.nih.gov/go/167>.

Brian R. Berridge,

Associate Director, National Toxicology Program.

[FR Doc. 2019–17269 Filed 8–12–19; 8:45 am]

BILLING CODE 4140–01–P

DEPARTMENT OF HEALTH AND HUMAN SERVICES

National Institutes of Health

National Center for Advancing Translational Sciences; Amended Notice of Meeting

Notice is hereby given of a change in the meeting of the National Center for Advancing Translational Sciences Advisory Council, which was published in the **Federal Register** on June 27, 2019, 84 FR 30740, Pg. 30744.

Amendment to change start time of the closed session from 9 a.m. to 8:30 a.m. and to change the start time of the open session from 11 a.m. to 10 a.m. The meeting is partially Closed to the public.

Dated: August 7, 2019.

Melanie J. Pantoja,

Program Analyst, Office of Federal Advisory Committee Policy.

[FR Doc. 2019–17245 Filed 8–12–19; 8:45 am]

BILLING CODE 4140–01–P

DEPARTMENT OF HEALTH AND HUMAN SERVICES

National Institutes of Health

National Institute of Allergy and Infectious 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 Allergy and Infectious Diseases Special Emphasis Panel; NIH Support for Conferences and Scientific Meetings (Parent R13 Clinical Trial Not Allowed).

Date: August 28, 2019.

Time: 1:00 p.m. to 3:00 p.m.

Agenda: To review and evaluate grant applications.

Place: National Institutes of Health, 5601 Fishers Lane, Rockville, MD 20892 (Telephone Conference Call).

Contact Person: Steven F. Santos, Ph.D., Scientific Review Officer, SRP, DEA, NIAID–NIH, DHHS, 5601 Fishers Lane, 3G33, Rockville, MD 20852, Phone: 301–761–7049, Cell: 202–306–4207, steven.santos@nih.gov. (Catalogue of Federal Domestic Assistance Program Nos. 93.855, Allergy, Immunology, and Transplantation Research; 93.856, Microbiology and Infectious Diseases Research, National Institutes of Health, HHS)

Dated: August 7, 2019.

Sylvia L. Neal,

Program Analyst, Office of Federal Advisory Committee Policy.

[FR Doc. 2019–17247 Filed 8–12–19; 8:45 am]

BILLING CODE 4140–01–P

DEPARTMENT OF HEALTH AND HUMAN SERVICES

Substance Abuse and Mental Health Services Administration

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

Periodically, the Substance Abuse and Mental Health Services Administration (SAMHSA) will publish a summary of information collection requests under OMB review, in compliance with the

Paperwork Reduction Act (44 U.S.C. Chapter 35). To request a copy of these documents, call the SAMHSA Reports Clearance Officer on (240) 276–1243.

Project: Protection and Advocacy for Individuals With Mental Illness (PAIMI) Final Rule, 42 CFR Part 51 (OMB No. 0930–0172)—Extension

These regulations meet the directive under 42 U.S.C. 10826(b) requiring the Secretary to promulgate final regulations to carry out the PAIMI Act (42 U.S.C. 10801 *et seq.*). The regulations contain information collection requirements associated with the rule. The Act authorizes funds to support activities on behalf of individuals with significant (severe) mental illness (adults) or significant (severe) emotional impairment (children/youth) as defined by the Act at 42 U.S.C. 10802(4) and 10804(d). Only entities designated by the governor of each state, including the American Samoa, Guam, Commonwealth of the Northern Mariana Islands, Commonwealth of Puerto Rico, U.S. Virgin Islands, District of Columbia (Mayor), and the tribal councils of the American Indian Consortium (the Hopi Tribe and the Navajo Nation located in the Four Corners region of the Southwest), to protect and advocate the rights of persons with developmental disabilities are eligible to receive PAIMI Program grants [*ibid* at 42 U.S.C. at 10802(2)]. These grants are based on a formula prescribed by the Secretary [*ibid* at 42 U.S.C. at 10822(a)(1)(A)].

On January 1, each eligible state protection and advocacy (P&A) system is required to prepare an annual PAIMI Program Performance Report (PPR). Each annual PPR describes a P&A system's activities, accomplishments and expenditures to protect the rights of individuals with mental illness supported with payments from PAIMI program allotments during the most recently completed fiscal year. Each P&A system transmit a copy of its annual report to the Secretary (via SAMHSA) and to the State Mental Health Agency where the system is located per the PAIMI Act at 42 U.S.C. 10824(a). Each annual PPR must provide the Secretary with the following information:

- The number of (PAIMI-eligible) individuals with mental illness served;
- A description of the types of activities undertaken;
- A description of the types of facilities providing care or treatment to which such activities are undertaken;
- A description of the manner in which the activities are initiated;

- A description of the accomplishments resulting from such activities;
- A description of systems to protect and advocate the rights of individuals with mental illness supported with payments from PAIMI Program allotments;
- A description of activities conducted by States to protect and advocate such rights;
- A description of mechanisms established by residential facilities for individuals with mental illness to protect such rights;
- A description of the coordination among such systems, activities and mechanisms;
- Specification of the number of public and nonprofit P&A systems established with PAIMI Program allotments; and
- Recommendations for activities and services to improve the protection and advocacy of the rights of individuals with mental illness and a description of the need for such activities and services that were not met by the state P&A systems established under the PAIMI

Act due to resource or annual program priority limitations.

Each PAIMI grantee's annual PPR must include a separate section, prepared by its PAIMI Advisory Council (PAC), that describes the council's activities and its assessment of the state P&A system's operations per the PAIMI Act at 42 U.S.C. 10805(7).

In 2017, SAMHSA included the annual PAIMI PPR in the Web-based Block Grant Application System (WebBGAS). WebBGAS, SAMHSA's electronic data system, is used to collect grantee information for the following reasons:

(1) To meet the OMB requirements for data collection for mandatory (formula) grant programs;

(2) To comply with the annual program reporting requirements of the PAIMI Act 42 U.S.C. 10801 *et seq.* and the PAIMI Rules 42 CFR, Part 51;

(3) To simplify the submission of PAIMI program data by the state P&A systems;

(4) To meet the Government Performance Results Act (GPRA) requirements;

(5) To comply with the Government Accountability Office (GAO) evaluation

recommendations that SAMHSA obtain information that closely measures the actual outcomes of the programs it funds;

(6) To reduce the grantee data collection burden by removing information that did not facilitate evaluation of a PAIMI grantee's programmatic and financial management systems;

(7) To provide immediate access to the PAIMI program data used to prepare a section of the Secretary's biennial report to the President, Congress, and National Council on Disability in accordance with the *Developmental Disabilities Assistance Act of 2000* at 42 U.S.C. 15005. Reports of the Secretary;

(8) To improve SAMHSA's ability to create reports, analyze trends and provide timely feedback to the P&A grantees when PPR revisions are needed.

On July 17, 2017, OMB approved SAMHSA's PPR and Advisory Council Report (Control No. 0930-0169, Expiration Date July 31, 2020). The burden estimate for the annual State P&A system reporting requirements for these regulations is as follows:

42 CFR citation	Number of respondents	Responses per respondent	Burden/response (hrs.)	Total hour burden
51.8(a)(2) Program Performance Report ¹				
51.8(b)(8) Advisory Council Report*				
51.10 Remedial Actions: Corrective Action Plans & Implementation Status Reports.	5	2	8	80
51.23(c) Reports, materials and fiscal data provided to the Advisory Council	5	3	2	30
51.25(b)(3) Grievance Procedure	57	1	1	57
51.43 Written denial of access by P&A system**	57	1	0.5	28.5
Total	57		11.5	195.5

Note: Burden for the annual application [42 CFR 51.5(b–d)] is approved at a standard level per application under OMB control number 0920–0428.

¹ Responses and burden hours associated with these reports are approved under OMB No. 0930–0169.

^{**} There is no burden estimate associated with this program provision. State P&A systems report that when a facility denies a P&A system access to the facility, a client, or records, the P&A attempts to resolve the dispute through negotiation, conciliation, mediation, and other non-adversarial techniques. Only after exhausting the non-legal remedies provided under state and federal laws will a P&A system file a formal complaint in the appropriate federal district court. See also, the PAIMI Act at 42 U.S.C. 10807(a)—Legal Actions and the PAIMI Final Rule at 42 CFR 51.32—Resolving Disputes.

Written comments and recommendations concerning the proposed information collection should be sent by September 12, 2019 to the SAMHSA Desk Officer at the Office of Information and Regulatory Affairs, OMB. To ensure timely receipt of comments, and to avoid potential delays in OMB's receipt and processing of mail sent through the U.S. Postal Service, commenters are encouraged to submit their comments to OMB via email to: OIRA_Submission@omb.eop.gov. Although commenters are encouraged to send their comments via email, commenters may also fax their

comments to: (202) 395–7285. Commenters may also mail them to: Office of Management and Budget, Office of Information and Regulatory Affairs, New Executive Office Building, Room 10102, Washington, DC 20503.

Summer King,
Statistician.

[FR Doc. 2019–17263 Filed 8–12–19; 8:45 am]

BILLING CODE 4162–20–P

DEPARTMENT OF HEALTH AND HUMAN SERVICES

Substance Abuse and Mental Health Services Administration

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

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