Effectiveness Trials for Treatment, Preventive and Services Interventions (R34)

Date: June 24, 2016.

Time: 11:30 a.m. to 3:00 p.m.

Agenda: To review and evaluate grant applications.

Place: National Institutes of Health, Neuroscience Center, 6001 Executive Boulevard, Rockville, MD 20852, (Telephone Conference Call).

Contact Person: Karen Gavin-Evans, Ph.D., Scientific Review Officer, Division of Extramural Activities, National Institute of Mental Health, NIH, Neuroscience Center, 6001 Executive Boulevard, Room 6153, MSC 9606, Bethesda, MD 20892, 301–451–2356, *gavinevanskm@mail.nih.gov.*

Name of Committee: National Institute of Mental Health Special Emphasis Panel, Fellowship and Dissertation Grants Review Meeting.

Date: June 30, 2016.

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

Agenda: To review and evaluate grant applications.

Place: National Institutes of Health, Neuroscience Center, 6001 Executive Boulevard, Rockville, MD 20852, (Telephone Conference Call).

Contact Person: Marcy Ellen Burstein, Ph.D., Scientific Review Officer, Division of Extramural Activities, National Institute of Mental Health, NIH Neuroscience Center, 6001 Executive Blvd., Room 6143, MSC 9606, Bethesda, MD 20892–9606, 301–443–9699, *bursteinme@mail.nih.gov.*

Name of Committee: National Institute of Mental Health Special Emphasis Panel, Mental Health Services Conflicts (Teleconference).

Date: June 30, 2016.

Time: 12:30 p.m. to 1:30 p.m.

Agenda: To review and evaluate grant applications.

Place: National Institutes of Health, Neuroscience Center, 6001 Executive Boulevard, Rockville, MD 20852, (Telephone Conference Call).

Contact Person: Karen Gavin-Evans, Ph.D., Scientific Review Officer, Division of Extramural Activities, National Institute of Mental Health, NIH Neuroscience Center, 6001 Executive Boulevard, Room 6153, MSC 9606, Bethesda, MD 20892, 301–451–2356, *gavinevanskm@mail.nih.gov.*

(Catalogue of Federal Domestic Assistance Program No. 93.242, Mental Health Research Grants, National Institutes of Health, HHS)

Dated: June 1, 2016.

Carolyn A. Baum,

Program Analyst, Office of Federal Advisory Committee Policy.

[FR Doc. 2016–13310 Filed 6–6–16; 8:45 am]

BILLING CODE 4140–01–P

DEPARTMENT OF HEALTH AND HUMAN SERVICES

National Institutes of Health

Submission for OMB Review; 30-day Comment Request, U.S. Nuclear Medicine Technologists Study (NCI)

SUMMARY: Under the provisions of Section 3507(a)(1)(D) of the Paperwork Reduction Act of 1995, the National Cancer Institute, the National Institutes of Health, has submitted to the Office of Management and Budget (OMB) a request for review and approval of the information collection listed below. This proposed information collection was previously published in the **Federal Register** on March 28, 2016 and allowed 60-days for public comment. No public comments were received. The purpose of this notice is to allow an additional 30 days for public comment. The National Cancer Institute, National Institutes of Health, may not conduct or sponsor, and the respondent is not required to respond to, an information collection that has been extended, revised, or implemented on or after October 1, 1995, unless it displays a currently valid OMB control number.

Direct Comments to OMB: Written comments and/or suggestions regarding the item(s) contained in this notice, especially regarding the estimated public burden and associated response time, should be directed to the: Office of Management and Budget, Office of Regulatory Affairs, *OIRA_submission@omb.eop.gov* or by fax to 202–395–6974, Attention: NIH Desk Officer.

Comment Due Date: Comments regarding this information collection are best assured of having their full effect if received within 30 days of the date of this publication.

FOR FURTHER INFORMATION CONTACT: To obtain a copy of the data collection plans and instruments, or request more information on the proposed project, contact*: Michele M. Doody, Radiation Epidemiology Branch, National Cancer Institute, 9609 Medical Center Drive, Room 7E566, Rockville, MD 20850, or call non-toll-free number 301–414–0308. Or Email your request, including your address to: *doodym@mail.nih.gov.* Formal requests for additional plans and instruments must be requested in writing.

Proposed Collection: U.S. Nuclear Medicine Technologists Study, 0925–0656, Expiration Date 04/30/2015—REINSTATEMENT WITH CHANGE, National Cancer Institute (NCI), National Institutes of Health (NIH).

Need and Use of Information Collection: We propose to collect from U.S. nuclear medicine technologists (USNMT) certified after 1980 historical information about nuclear medicine procedures performed, radioisotopes used, related work and safety practices, and places of employment. The primary objectives of the current feasibility effort are: (a) To identify a cohort of nuclear medicine technologists certified after 1980 by the American Registry of Radiologic Technologists (ARRT) and/or the Nuclear Medicine Technologist Certification Board (NMTCB); and (b) to characterize individual organ-specific occupational radiation doses from radioisotope procedures. More recently certified technologists, who specialized in nuclear medicine, are expected to have greater exposures to radioisotopes than the general radiologic technologists in the U.S. Radiologic Technologist (USRT) cohort owing to performing such procedures with greater frequency. The proposed USNMT study would be a direct follow-on to the USRT Study to assess health risks associated with occupational exposure to these much higher-energy radiopharmaceuticals.

OMB approval is requested for 3 years. There are no costs to respondents other than their time. The total estimated annualized burden hours are 125.

ESTIMATED ANNUALIZED BURDEN HOURS

Type of respondent	Instrument	Number of respondents	Frequency of response	Average time per response (hours)	Annual hour burden
Nuclear Medicine Technologists	Nuclear Medicine Questionnaire	250	1	30/60	125
Total		250	250		125

Dated: May 31, 2016.

Karla Bailey,

Project Clearance Liaison, National Cancer Institute, NIH.

[FR Doc. 2016–13308 Filed 6–6–16; 8:45 am]

BILLING CODE 4140–01–P

DEPARTMENT OF HEALTH AND HUMAN SERVICES

National Institutes of Health

National Cancer Institute Amended; Notice of Meeting

Notice is hereby given of a change in the meeting of the National Cancer Institute Special Emphasis Panel, June 23, 2016, 08:00 a.m. to June 24, 2016, 06:00 p.m., Doubletree Hotel Bethesda, 8120 Wisconsin Avenue, Bethesda, MD, 20814 which was published in the **Federal Register** on May 16, 2016, 81 FR 30318.

The meeting notice is amended to change the Committee name from National Cancer Institute, Special Emphasis Panel; NCI Omnibus R03 SEP–1 to National Cancer Institute, Special Emphasis Panel; NCI R03 SEP–2. The meeting is closed to the public.

Dated: June 1, 2016.

Melanie J. Gray,

Program Analyst, Office of Federal Advisory Committee Policy.

[FR Doc. 2016–13309 Filed 6–6–16; 8:45 am]

BILLING CODE 4140–01–P

DEPARTMENT OF HEALTH AND HUMAN SERVICES

Substance Abuse and Mental Health Services Administration

Notice of Meeting

Pursuant to Public Law 92–463, notice is hereby given that the Substance Abuse and Mental Health Services Administration's (SAMHSA) Center for Substance Abuse Prevention (CSAP) National Advisory Council will meet on June 13, 2016, 1:30 p.m.–2:30 p.m., via teleconference.

The meeting will include the review, discussion, and evaluation of grant applications reviewed by the Initial Review Group, and involve an examination of confidential financial and business information as well as personal information concerning the applicants. Therefore, these meetings will be closed to the public as determined by the SAMHSA Administrator, in accordance with Title 5 U.S.C. 552b(c)(4) and (c)(6); and 5 U.S.C. App. 2, Section 10(d).

Committee Name: Substance Abuse and Mental Health Services, Administration Center for Substance Abuse Prevention National Advisory Council.

Date/Time/Type: June 13, 2016, 1:30 p.m.–2:30 p.m. (CLOSED).

Place: SAMHSA Building, 5600 Fishers Lane, Rockville, MD 20857.

Contact: Matthew J. Aumen, Designated Federal Officer, SAMHSA/CSAP National Advisory Council, 5600 Fishers Lane, Rockville, MD 20857, Email: *Matthew.Aumen@samhsa.hhs.gov*.

Summer King,

Statistician, SAMHSA.

[FR Doc. 2016–13369 Filed 6–6–16; 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

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. The regulations contain information collection requirements. 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 42 U.S.C. 10802(4) and 10804(d). Only entities designated by the governor of each State, including American Samoa, Guam, the Commonwealth of the Northern Mariana Islands, the Commonwealth of Puerto Rico, the U.S. Virgin Islands, the Mayor of the District of Columbia, and the tribal councils for the American Indian Consortium (the Hopi and Navajo Nations 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 [the Act at 42 U.S.C. at 10802(2)]. These grants are based on a formula prescribed by the Secretary [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 a report that describes its 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. The PAIMI Act at 42 U.S.C. 10824(a) requires that each P&A system transmit a copy of its annual report to the Secretary (via SAMHSA/CMHS) and to the State Mental Health Agency where the system is located. These annual PAIMI Program Performance Reports (PPR) to the Secretary must include 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; and,
- 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;
- 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.

The PAIMI Rules [42 CFR part 51] mandate that each State P&A system may place restrictions on either its case or client acceptance criteria developed as part of its annual PAIMI priorities. Each P&A system is required to inform