Agenda: To review and evaluate grant applications.

Place: National Institutes of Health, 6701 Rockledge Drive, Bethesda, MD 20892, (Telephone Conference Call).

Contact Person: Barna Dey, Ph.D., Scientific Review Officer, Center for Scientific Review, National Institutes of Health, 6701 Rockledge Drive, Room 3184, Bethesda, MD 20892, 301–451–2796, *bdey@mail.nih.gov*.

Name of Committee: Center for Scientific Review Special Emphasis Panel Member Conflict: Research Grant Program.

Date: July 26, 2017.

Time: 11:00 a.m. to 9:00 p.m.

Agenda: To review and evaluate grant applications.

Place: National Institutes of Health, 6701 Rockledge Drive, Bethesda, MD 20892, (Virtual Meeting).

Contact Person: Kenneth A. Roebuck, Ph.D., Scientific Review Officer, Center for Scientific Review, National Institutes of Health, 6701 Rockledge Drive, Room 5106, MSC 7852, Bethesda, MD 20892, (301) 435–1166, *roebuckk@csr.nih.gov.*

Name of Committee: Center for Scientific Review Special Emphasis Panel Member conflict: Exploratory/Developmental Research.

Date: July 26, 2017.

Time: 11:00 a.m. to 2:00 p.m.

Agenda: To review and evaluate grant applications.

Place: National Institutes of Health, 6701 Rockledge Drive, Bethesda, MD 20892, (Telephone Conference Call).

Contact Person: Mark P. Rubert, Ph.D., Scientific Review Officer, Center for Scientific Review, National Institutes of Health, 6701 Rockledge Drive, Room 5218, MSC 7852, Bethesda, MD 20892, 301–435–1775, *rubertm@csr.nih.gov.*

Name of Committee: Center for Scientific Review Special Emphasis Panel Member conflict: AIDS and AIDS Related Research.

Date: July 27–28, 2017.

Time: 10:30 a.m. to 5:00 p.m.

Agenda: To review and evaluate grant applications.

Place: National Institutes of Health, 6701 Rockledge Drive, Bethesda, MD 20892, (Virtual Meeting).

Contact Person: Jingsheng Tuo, Ph.D., Scientific Review Officer, Center for Scientific Review, National Institutes of Health, 6701 Rockledge Drive, Room 5207, Bethesda, MD 20892, 301–451–8754, *tuoj@nei.nih.gov.*

Name of Committee: Center for Scientific Review Special Emphasis Panel Member Conflict: AIDS and Related Research.

Date: August 2, 2017.

Time: 9:00 a.m. to 6:00 p.m.

Agenda: To review and evaluate grant applications.

Place: National Institutes of Health, 6701 Rockledge Drive, Bethesda, MD 20892, (Virtual Meeting).

Contact Person: Barna Dey, Ph.D., Scientific Review Officer, Center for Scientific Review, National Institutes of Health, 6701 Rockledge Drive, Room 3184, Bethesda, MD 20892, 301–451–2796, *bdey@mail.nih.gov.*

(Catalogue of Federal Domestic Assistance Program Nos. 93.306, Comparative Medicine; 93.333, Clinical Research, 93.306, 93.333, 93.337, 93.393–93.396, 93.837–93.844, 93.846–93.878, 93.892, 93.893, National Institutes of Health, HHS)

Dated: June 29, 2017.

David Clary,

Program Analyst, Office of Federal Advisory Committee Policy.

[FR Doc. 2017–14126 Filed 7–5–17; 8:45 am]

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DEPARTMENT OF HEALTH AND HUMAN SERVICES

National Institutes of Health

Proposed Collection; 60-Day Comment Request; Identifying Experts in Prevention Science Methods To Include on NIH Review Panels, Office of Disease Prevention (NIH ODP)

AGENCY: National Institutes of Health, Department of Health and Human Services.

ACTION: Notice.

SUMMARY: In compliance with the requirement of Section 3506(c)(2)(A) of the Paperwork Reduction Act of 1995, to provide opportunity for public comment on proposed data collection projects, the National Institutes of Health (NIH), Office of Disease Prevention (ODP) will publish periodic summaries of proposed projects to be submitted to the Office of Management and Budget (OMB) for review and approval.

DATES: Comments regarding this information collection are best assured of having their full effect if received within 60 days of the date of this publication.

FOR FURTHER INFORMATION CONTACT: To obtain a copy of the data collection plans and instruments, submit comments in writing, or request more information on the proposed project, contact: Dr. Ranell Myles, Public Health Analyst, NIH Office of Disease Prevention, 6100 Executive Blvd., Room 2B03, Bethesda, MD 20892 or call (301) 827–5579 or email your request, including your address to *prevention@mail.nih.gov.* Formal requests for additional plans and instruments must be requested in writing.

SUPPLEMENTARY INFORMATION: Section 3506(c)(2)(A) of the Paperwork Reduction Act of 1995 requires: Written comments and/or suggestions from the public and affected agencies are invited to address one or more of the following points: (1) Whether the proposed collection of information is necessary for the proper performance of the function of the agency, including whether the information will have practical utility; (2) 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; (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 those who are to respond, including the use of appropriate automated, electronic, mechanical, or other technological collection techniques or other forms of information technology.

Proposed Collection Title: Identifying Experts in Prevention Science Methods to Include on NIH Review Panels,—REVISION, Office of Disease Prevention (ODP), National Institutes of Health (NIH).

Need and Use of Information Collection: The Office of Disease Prevention (ODP) is the lead Office at the National Institutes of Health (NIH) responsible for assessing, facilitating, and stimulating research in disease prevention and health promotion, and disseminating the results of this research to improve public health. Prevention is preferable to treatment, and research on disease prevention is an important part of the NIH's mission. The knowledge gained from this research leads to stronger clinical practice, health policy, and community health programs. ODP collaborates with the NIH, other Department of Health and Human Services (DHHS) agencies, and other public and private partners to achieve the Office's mission and goals. One of our priorities is to promote the use of the best available methods in prevention research and support the development of better methods. One of our strategies is to help the Center for Scientific Review (CSR) identify experts in prevention science methods to include on their review panels. This will strengthen the panels and improve the quality of the prevention research supported by the NIH. To identify experts in prevention science methods, we worked with our contractor, IQ Solutions, Inc., to develop online software which will allow us to collect scientists' names, contact information, and resumes, as well as to have those scientists identify their level of

expertise in a variety of prevention science methods and content areas. The a collected with this software was used to create a web-based tool that CSR staff can use to identify scientists with expertise in specific prevention science methods and content areas for invitation to serve on one of the CSR review panels. This system will also be shared with review staff in the other Institutes and Centers at the NIH, as well as other DHHS agencies, to use in the same way. Given our plans to create an automated system for reviewer information collection, we are now seeking OMB approval for a revision to our data collection plan.

This OMB revision request is for the collection of additional data not collected in the previously deployed online software and survey including additional study design topics, research methods, content topics, as well as the geographic region of research of the investigator/respondent and the income category of the region/country in which the investigator's/respondent's research is performed.

OMB approval is requested for 3 years. There are no costs to respondents other than their time. The total estimated annualized burden hours are 1,300.

ESTIMATED ANNUALIZED BURDEN HOURS

Type of respondents	Number of respondents	Number of responses per respondent	Average time per response (in hours)	Total annual burden hour
Investigators	3,120	1	25/60	1,300
Total		3,120		1,300

Dated: June 29, 2017.

Lawrence A. Tabak,

Principal Deputy Director, National Institutes of Health.

[FR Doc. 2017–14087 Filed 7–5–17; 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: State Targeted Response to the Opioid Crisis (Opioid STR) Evaluation—NEW

The Substance Abuse and Mental Health Services Administration (SAMHSA) recently awarded 57 grants to states and territories to help address the national opioid crisis by increasing access to treatment, reducing unmet treatment needs, and reducing opioid overdose related deaths through the provision of prevention, treatment, and recovery activities for opioid use disorder (OUD).

SAMHSA's Center for Behavioral Health Statistics and Quality (CBHSQ), will be conducting a cross-site evaluation of the Opioid STR grant program. The proposed data collection is necessary to evaluate how the Opioid STR state/territory grantees plan and implement prevention, treatment and recovery services. Additionally, a subset of communities/programs will be selected to participate in supplemental evaluation activities designed to provide detailed information related to the implementation of services at the program/community level, as well as the impacts of the program on client outcomes.

SAMHSA has developed a set of interview protocols and survey measures that will collect information from all state/territory grantees (57), and subset (up to 20) of programs/communities that provide services and activities funded by the grant. In addition, SAMHSA's Performance Accountability and Reporting System (SPARS) will be used to collect individual-level data using CSAT's Government Performance and Results Act (GPRA) for Discretionary Grant Programs Client Outcome Measure (OMB No. 0930–0208) from individuals receiving services from participating communities/programs.

Specific data collected as part of the Opioid STR evaluation include the following:

State Survey: The State Survey will be administered to State Project Directors/Program Managers to collect information about the state/territory's current, planned, and implemented activities to address opioid misuse using Opioid STR funding. State Surveys will be administered three (3) times.

State Interview: The State Interview Protocol will be used to collect information from the State Project Director/Program Manager during phone interviews at two (2) time points. Interviews will collect information about the state's substance abuse treatment systems prior to STR funding, the types of activities states plan to implement with STR funding, challenges and successes when implementing these activities, and plans for sustaining the activities.

Community/Program Survey: The Community/Program Survey will be administered to Community/Program Directors or Program Managers to collect information about the community/program's readiness to implement activities that address opioid misuse, their actual implementation of activities to address opioid misuse, and initial outcomes of their implemented activities. Community/Program Survey will be administered three (3) times.

Community/Program Director/Manager Interview Protocol: The Community/Program Director/Manager Interview Protocol will be used to collect information from Community/Program Directors or Program Managers during in-person site visits to each participating community/program. Interviews will collect in-depth information about the community's/program's implementation of activities to address opioid misuse using Opioid STR funding, and factors facilitating and impeding the implementation of STR-funded activities. Community/Program Director/Manager Interviews will be conducted two (2) times.

Community/Program Data Manager Interview Protocol: The Community/Program Data Manager Interview Protocol will be used to collect information from Data Managers during in-person site visits to each participating community/program. Interviews will collect in-depth information about how the program used community/program-level data to