Date: March 7, 2022.

Time: 3:00 p.m. to 6:00 p.m.

Agenda: To review and evaluate grant applications.

Place: Eunice Kennedy Shriver National Institute of Child Health and Human Development, National Institutes of Health, 6710B Rockledge Drive, Room 2121B, Bethesda, MD 20892 (Video Assisted Meeting).

Contact Person: Christiane M. Robbins, Scientific Review Officer, Scientific Review Branch, Eunice Kennedy Shriver National Institute of Child Health and Human Development, National Institute of Health, 6710B Rockledge Drive, Room 2121B, Bethesda, MD 20892, (301) 451–4989, *crobbins@mail.nih.gov.*

Name of Committee: National Institute of Child Health and Human Development Special Emphasis Panel; Pelvic Floor Disorders Network (UG1 Clinical Research).

Date: March 17, 2022.

Time: 10:00 a.m. to 6:00 p.m.

Agenda: To review and evaluate grant applications.

Place: Eunice Kennedy Shriver National Institute of Child Health and Human Development, National Institutes of Health, 6710B Rockledge Drive, Room 2131B, Bethesda, MD 20892 (Video Assisted Meeting).

Contact Person: Luis E. Dettin, Ph.D., Scientific Review Officer, Scientific Review Branch, Eunice Kennedy Shriver National Institute of Child Health and Human Development, National Institute of Health, 6710B Rockledge Drive, Room 2131B, Bethesda, MD 20892, (301) 827–8231, *luis_dettin@nih.gov.*

Name of Committee: National Institute of Child Health and Human Development Special Emphasis Panel; Pelvic Floor Disorders Network (U24 Resource-Related Research Projects).

Date: March 18, 2022.

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

Agenda: To review and evaluate grant applications.

Place: Eunice Kennedy Shriver National Institute of Child Health and Human Development, National Institutes of Health, 6710B Rockledge Drive, Room 2131B, Bethesda, MD 20892 (Video Assisted Meeting).

Contact Person: Luis E. Dettin, Ph.D., Scientific Review Officer, Scientific Review Branch, Eunice Kennedy Shriver National Institute of Child Health and Human Development, National Institutes of Health, 6710B Rockledge Drive, Room 2131B, Bethesda, MD 20892, (301) 827–8231, *luis_dettin@nih.gov.*

(Catalogue of Federal Domestic Assistance Program Nos. 93.864, Population Research; 93.865, Research for Mothers and Children; 93.929, Center for Medical Rehabilitation Research; 93.209, Contraception and Infertility Loan Repayment Program, National Institutes of Health, HHS)

Dated: January 13, 2022.

Melanie J. Pantoja,

Program Analyst, Office of Federal Advisory Committee Policy.

[FR Doc. 2022–00963 Filed 1–18–22; 8:45 am]

BILLING CODE 4140–01–P

DEPARTMENT OF HEALTH AND HUMAN SERVICES

National Institutes of Health

National Human Genome Research Institute; 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 Human Genome Research Institute Special Emphasis Panel; Knock Out Mouse Phenotyping Program (KOMP2).

Date: March 10, 2022.

Time: 12:00 p.m. to 4:30 p.m.

Agenda: To review and evaluate grant applications.

Place: National Human Genome Research Institute, National Institutes of Health, 6700B Rockledge Drive, Suite 3100, Bethesda, MD 20892 (Virtual Meeting).

Contact Person: Barbara J. Thomas, Ph.D., Scientific Review Officer, Scientific Review Branch, National Human Genome Research Institute, National Institutes of Health, 6700B Rockledge Drive, Suite 3100, Bethesda, MD 20892, 301–402–8837, *barbara.thomas@nih.gov.*

(Catalogue of Federal Domestic Assistance Program Nos. 93.172, Human Genome Research, National Institutes of Health, HHS)

Dated: January 12, 2022.

David W. Freeman,

Program Analyst, Office of Federal Advisory Committee Policy.

[FR Doc. 2022–00900 Filed 1–18–22; 8:45 am]

BILLING CODE 4140–01–P

DEPARTMENT OF HEALTH AND HUMAN SERVICES

National Institutes of Health

National Heart, Lung, and Blood Institute; 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: Heart, Lung, and Blood Initial Review Group; NHLBI Mentored Patient-Oriented Research Study Section.

Date: March 3–4, 2022.

Time: 10:00 a.m. to 6:00 p.m.

Agenda: To review and evaluate grant applications.

Place: National Institutes of Health, 6705 Rockledge Drive, Bethesda, MD 20817 (Virtual Meeting).

Contact Person: Stephanie Johnson Webb, Ph.D., Scientific Review Officer, Office of Scientific Review/DERA, National Heart, Lung, and Blood Institute, National Institutes of Health, 6705 Rockledge Drive, Room 208–V, Bethesda, MD 20892, (301) 827–7992, *stephanie.webb@nih.gov.*

(Catalogue of Federal Domestic Assistance Program Nos. 93.233, National Center for Sleep Disorders Research; 93.837, Heart and Vascular Diseases Research; 93.838, Lung Diseases Research; 93.839, Blood Diseases and Resources Research, National Institutes of Health, HHS)

Dated: January 12, 2022.

David W. Freeman,

Program Analyst, Office of Federal Advisory Committee Policy.

[FR Doc. 2022–00899 Filed 1–18–22; 8:45 am]

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DEPARTMENT OF HEALTH AND HUMAN SERVICES

National Institutes of Health

National Heart, Lung, and Blood Institute; 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: Heart, Lung, and Blood Initial Review Group; Clinical Trials Review Study Section.

Date: March 3–4, 2022.

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

Agenda: To review and evaluate grant applications.

Place: National Institutes of Health, 6705 Rockledge Drive, Bethesda, MD 20817 (Virtual Meeting).

Contact Person: Keary A. Cope, Ph.D., Scientific Review Officer, Office of Scientific Review/DERA, National Heart, Lung, and Blood Institute, National Institutes of Health, 6705 Rockledge Drive, Room 209–A, Bethesda, MD 20892–7924, (301) 827–7912, *copeka@mail.nih.gov*.

(Catalogue of Federal Domestic Assistance Program Nos. 93.233, National Center for Sleep Disorders Research; 93.837, Heart and Vascular Diseases Research; 93.838, Lung Diseases Research; 93.839, Blood Diseases and Resources Research, National Institutes of Health, HHS)

Dated: January 12, 2022.

David W. Freeman,

Program Analyst, Office of Federal Advisory Committee Policy.

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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. To request a copy of these documents, call the SAMHSA Reports Clearance Officer on (240) 276–0361.

Project: Revision of Mental Health Client/Participant Outcome Measures and Infrastructure, Prevention, and Mental Health Promotion Indicators (OMB No. 0930–0285)

SAMHSA is requesting approval from the Office of Management and Budget (OMB) for revisions to the previously approved instruments and data collection activities for the Government Performance and Results Act (GPRA) Center Mental Health Services (OMB No 0930–0285) that expires on February 28, 2022.

To be fully accountable for the spending of federal funds, SAMHSA requires all programs to collect and report data to ensure that program goals and objectives are met. Data is collected and used to monitor and improve performance of each program and ensure appropriate and thoughtful spending of federal funds.

SAMHSA requests the following revisions to the NOMS Mental Health Client/Participant Outcome measures: (1) Merge the CMHS NOMS Child Client-level Measures for Discretionary Programs data collection instrument with the current CMHS NOMS Adult Client-level Measures for Discretionary Programs data collection instrument; (2) delete questions for data not being utilized for program monitoring and quality improvement; (3) reduce grantee burden by shifting questions for a five-point psychometric response scale to ''Yes'', ''No'', and ''No response/Refused'' responses; (4) modify IDC–10 diagnoses to expand the F 40–48, F60–63, and F90–99 codes to allow for more specificity. Also, add ICD–10 ''Z'' codes to allow for a focus on social determinants of health that may affect the diagnosis, course, prognosis, or treatment of a client/consumer mental disorder; (6) shift reporting NOMS data to baseline assessment, 3-month or 6-month reassessment, and a final clinical discharge assessment; (7) reduce the number of physical health indictors and reporting frequency from quarterly to three points in time (baseline, 3- or 6-month reassessment, clinical discharge).

SAMHSA also requests the following revisions to the Infrastructure, Prevention, and Mental Health Promotion indicators:

(1) Delete four indicators not used by any SAMSHA programs: PD1: The number of policy changes completed as a result of the grant; WD4: The number of changes made to credentialing and licensing policies in order to incorporate expertise needed to improve mental health-related practices/activities; F1: The amount of additional funding obtained for specific mental health-related practices/activities that are consistent with the goals of the grant; and O2: The total number of contacts made through program outreach efforts).

(2) Revise two indicators to provide more clarity A3: The number of communities that enhance health information-sharing for provision of services between agencies and program; and A1: The number of grant project activities in which fidelity is monitored as a result of the grant); and

(3) Add eleven indicators to reflect program developments during the past three years: R2: The number of individuals referred to trauma-informed care services as a result of the grant; R3: The number of individuals referred to crisis or other mental health services for suicidality; S2: The number of individuals screened for trauma-related experiences as a result of the grant; S3: The number of individuals screened for suicidal ideation as a result of the grant; T5: The number of activities modified, adapted, or changed to reflect trauma-informed practices for the population(s) being served by the grant; T6: The number of activities modified, adapted, or changed to reflect culturally appropriate services for the population(s) being served by the grant; T7: As a result of the grant, reduce the percentage of individuals who died by suicide; and T8: As a result of the grant, reduce the number of individuals who attempted suicide).

These changes will lessen grantee burden with data collection and improve capacity to report qualitative performance and quantitative outcomes for all discretionary grant programs, including: Demographic characteristics of clients' served; clinical characteristics of clients' served before, during, and after receipt of services; numbers of clients served; and characteristics of services and activities provided to clients'.

Currently, the information collected from this instrument is entered and stored on SAMHSA's Performance Accountability and Reporting System (SPARS), a real-time, performance management system that captures information on mental health and substance abuse treatment services delivered in the United States. Continued approval of this information collection will allow SAMHSA to continue to meet Government Performance and Results Modernization Act of 2010 (GPRMA) reporting requirements that quantify the effects and accomplishments of its discretionary grant programs, which are consistent with OMB guidance.

SAMHSA and its Centers will use the data collected for annual reporting required by required by GPRMA, to describe and understand changes in outcomes from baseline to follow-up to discharge. SAMHSA and its Centers will use the data for annual reporting comparing baseline with discharge and follow-up data. SAMHSA's report for each fiscal year will include actual results of performance monitoring for the three preceding fiscal years. Information collected through this request will allow SAMHSA to report on the results of these performance outcomes as well as be consistent with SAMHSA-specific performance domains, and to assess the