Group; Clinical Neuroimmunology and Brain Tumors Study Section.

Date: October 17–18, 2013.

Time: 8:00 a.m. to 5:00 p.m.

Agenda: To review and evaluate grant applications.

Place: Melrose Hotel, 2430 Pennsylvania Avenue NW., Washington, DC 20037.

Contact Person: Jay Joshi, Ph.D., Scientific Review Officer, Center for Scientific Review, National Institutes of Health, 6701 Rockledge Drive, Room 5196, MSC 7846, Bethesda, MD 20892, (301) 408–9135, *joshij@csr.nih.gov.*

Name of Committee: Musculoskeletal, Oral and Skin Sciences Integrated Review Group; Musculoskeletal Tissue Engineering Study Section.

Date: October 17–18, 2013.

Time: 8:00 a.m. to 5:30 p.m.

Agenda: To review and evaluate grant applications.

Place: Hyatt Regency Bethesda, One Bethesda Metro Center, 7400 Wisconsin Avenue, Bethesda, MD 20814.

Contact Person: Baljit S. Moonga, Ph.D., Scientific Review Officer, Center for Scientific Review, National Institutes of Health, 6701 Rockledge Drive, Room 4214, MSC 7806, Bethesda, MD 20892, 301–435–1777, *moongabs@mail.nih.gov*.

Name of Committee: Oncology 1-Basic Translational Integrated Review Group; Molecular Oncogenesis Study Section.

Date: October 17–18, 2013.

Time: 8:00 a.m. to 6:00 p.m.

Agenda: To review and evaluate grant applications.

Place: Embassy Suites Alexandria, 1900 Diagonal Road, Alexandria, VA.

Contact Person: Nywana Sizemore, Ph.D., Scientific Review Officer, Center for Scientific Review, National Institutes of Health, 6701 Rockledge Drive, Room 6204, MSC 7804, Bethesda, MD 20892, 301–435–1718, *sizemoren@csr.nih.gov*.

Name of Committee: Cardiovascular and Respiratory Sciences Integrated Review Group; Cardiac Contractility, Hypertrophy, and Failure Study Section.

Date: October 17, 2013.

Time: 8:00 a.m. to 6:00 p.m.

Agenda: To review and evaluate grant applications.

Place: Renaissance Mayflower Hotel, 1127 Connecticut Avenue NW., Washington, DC 20036.

Contact Person: Olga A. Tjurmina, Ph.D., Scientific Review Officer, Center for Scientific Review, National Institutes of Health, 6701 Rockledge Drive, Room 4030B, MSC 7814, Bethesda, MD 20892, (301) 451–1375, *ot3d@nih.gov.*

Name of Committee: Risk, Prevention and Health Behavior Integrated Review Group; Social Psychology, Personality and Interpersonal Processes Study Section.

Date: October 17, 2013.

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

Agenda: To review and evaluate grant applications.

Place: Renaissance Mayflower Hotel, 1127 Connecticut Avenue NW., Washington, DC 20036.

Contact Person: Monica Basco, Ph.D., Scientific Review Officer, Center for Scientific Review, National Institutes of Health, 6701 Rockledge Drive, Room 3220, MSC 7808, Bethesda, MD 20892, 301–496–7010, *bascoma@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: September 17, 2013.

Melanie J. Gray,

Program Analyst, Office of Federal Advisory Committee Policy.

[FR Doc. 2013–22983 Filed 9–20–13; 8:45 am]

BILLING CODE 4140–01–P

DEPARTMENT OF HEALTH AND HUMAN SERVICES

National Institutes of Health

National Institute of Diabetes and Digestive and Kidney Diseases; Notice of Closed Meetings

Pursuant to section 10(d) of the Federal Advisory Committee Act, as amended (5 U.S.C. App.), notice is hereby given of the following meetings.

The meetings 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 Diabetes and Digestive and Kidney Diseases Special Emphasis Panel, Bariatric Surgery—Related Ancillary Studies (R01s).

Date: October 29, 2013.

Time: 1:00 p.m. to 2:30 p.m.

Agenda: To review and evaluate grant applications.

Place: National Institutes of Health, Two Democracy Plaza, 6707 Democracy Boulevard, Bethesda, MD 20892, (Telephone Conference Call).

Contact Person: Paul A. Rushing, Ph.D., Scientific Review Officer, Review Branch, DEA, NIDDK, National Institutes of Health, Room 747, 6707 Democracy Boulevard, Bethesda, MD 20892–5452, (301) 594–8895, *rushingp@extra.niddk.nih.gov.*

Name of Committee: National Institute of Diabetes and Digestive and Kidney Diseases Special Emphasis Panel, Regulatory Mechanisms in Intestinal Motility (P01).

Date: November 8, 2013.

Time: 2:00 p.m. to 5:00 p.m.

Agenda: To review and evaluate grant applications.

Place: National Institutes of Health, Two Democracy Plaza, 6707 Democracy Boulevard, Bethesda, MD 20892, (Telephone Conference Call).

Contact Person: Paul A. Rushing, Ph.D., Scientific Review Officer, Review Branch, DEA, NIDDK, National Institutes of Health, Room 747, 6707 Democracy Boulevard, Bethesda, MD 20892–5452, (301) 594–8895, *rushingp@extra.niddk.nih.gov.*

(Catalogue of Federal Domestic Assistance Program Nos. 93.847, Diabetes, Endocrinology and Metabolic Research; 93.848, Digestive Diseases and Nutrition Research; 93.849, Kidney Diseases, Urology and Hematology Research, National Institutes of Health, HHS)

Dated: September 17, 2013.

David Clary,

Program Analyst, Office of Federal Advisory Committee Policy.

[FR Doc. 2013–22987 Filed 9–20–13; 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: SAMHSA Tobacco Prevention, Cessation, and Behavioral Health Message Testing—NEW

The Substance Abuse and Mental Health Services Administration (SAMHSA) is conducting message testing to inform the development and implementation of a tobacco use prevention and cessation campaign aimed at youth with substance use and/or mental health conditions.

The purpose of the project is to inform messaging efforts, through focus groups with youth and in-depth interviews with health care providers, to improve tobacco use prevention and cessation efforts in populations with mental health and substance use concerns, particularly youth and vulnerable populations. The focus groups and interviews are an integral part of the process to test messages and, ultimately, develop effective campaign materials and efficient implementation plans.

SAMHSA will screen parents (because focus group participants are

under the age of consent) and youth, conduct focus groups with youth with substance use and/or mental health conditions, and interview health care professionals who treat youth with these conditions. The screen will be administered by telephone to parents first and, as eligible, to youth and will take 10 minutes to complete for parents and for youth. Questions will include a mix of open-ended and closed-ended responses and are intended to gather information on previous diagnosis and symptomology of mental health conditions and availability to participate in the focus group. The focus groups with youth will be conducted in person and will take up to 90 minutes. Questions are primarily open-ended and intended to gather information on the reasons youth with substance use and/or mental health conditions use tobacco, the barriers and facilitators to tobacco use prevention and cessation, the appeal of various tobacco use prevention and cessation messages, and the best dissemination strategies and communication channels for a future campaign aimed at this specialized group. The interviews with health care professionals who treat youth with mental health and/or substance use conditions will be conducted in person, as feasible, or by telephone and will take up to 45 minutes. Questions are primarily open-ended and intended to gather information to better understand how various health care professionals screen for and address tobacco use in youth receiving care in their practice, identify messages and materials aimed at health care professionals to address tobacco use prevention and cessation in youth with substance use and/or mental health conditions, determine the most efficient communication strategies and channels to disseminate this information. All data collections are voluntary.

Below is the table of the estimated total burden hours:

Respondent	Number of respondents	Responses per respondent	Average burden hour	Total hour burden
Screener (Parent)	576	1	.15	86.4
Screener (Youth)	144	1	.15	21.6
Youth Focus Group	*108	1	1.50	162
Provider Interview	42	1	.75	31.5
Total	762			301.5

*The 108 respondents identified for the youth focus groups are included in the 144 respondents for the youth screener.

Written comments and recommendations concerning the proposed information collection should be sent by October 23, 2013 to the SAMHSA Desk Officer at the Office of Information and Regulatory Affairs, Office of Management and Budget (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. 2013–23053 Filed 9–20–13; 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: Fetal Alcohol Spectrum Disorders (FASD) Center for Excellence (CFE) Screening and Brief Intervention (SBI) Project and Project CHOICES Evaluation (OMB No. 0930–0302)—Reinstatement

Since 2001, SAMHSA's Center for Substance Abuse Prevention has been operating the SAMHSA Fetal Alcohol Spectrum Disorders (FASD) Center for Excellence (CFE). The purpose of the FASD Center for Excellence is to prevent alcohol-exposed pregnancies among women of childbearing age and pregnant women and to improve the quality of life for individuals affected by FASD. Data will be collected from women served across approximately 10 sites in local/community-based agencies. Women will be screened for alcohol use, and provided appropriate interventions based on their pregnancy status.

The FASD CFE will be integrating Screening and Brief Intervention (SBI) for pregnant women and Project CHOICES for non-pregnant women through service delivery organizations and will monitor the results. Approximately 10 sites will implement the SBI program and/or Project CHOICES.

At baseline, an assessment form will be administered by the counselor to screen women at the participating sites or health care delivery programs. Basic demographic data will be collected for all women screened (age, race/ethnicity, education, and marital status) at baseline by participating sites but no personal identification information will be transmitted to SAMHSA. Both quantity and frequency of drinking will be assessed for all women. Pregnant women will be assessed for risk of alcohol use using the TWEAK screening instrument, which has been used successfully with pregnant women. Non-pregnant women will be assessed for ability to conceive and use of effective birth control.

SBI focuses on 10- to 15-minute counseling sessions, conducted by a counselor who will use a scripted manual to guide the program. Participants in SBI will be assessed throughout their pregnancy to monitor