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

Contact Person: Rajiv Kumar, PhD, Scientific Review Officer, Center for Scientific Review, National Institutes of Health, 6701 Rockledge Drive, Room 4122, MSC 7802, Bethesda, MD 20892, 301–435–1212. *kumarra@csr.nih.gov*.

Name of Committee: Digestive, Kidney and Urological Systems Integrated Review Group. Clinical, Integrative and Molecular Gastroenterology Study Section.

Date: January 31–February 1, 2011.

Time: 8 a.m. to 5 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: Mushtaq A. Khan, DVM, PhD, Chief, Digestive, Kidney and Urological Systems, Center for Scientific Review, National Institutes of Health, 6701 Rockledge Drive, Room 2176, MSC 7818, Bethesda, MD 20892. 301–435–1778. *khanm@csr.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: December 10, 2010.

Jennifer S. Spaeth,

Director, Office of Federal Advisory Committee Policy.

[FR Doc. 2010–31601 Filed 12–15–10; 8:45 am]

BILLING CODE 4140–01–P

DEPARTMENT OF HEALTH AND HUMAN SERVICES

National Institutes of Health

National Library of Medicine; Notice of Meeting

Pursuant to section 10(a) of the Federal Advisory Committee Act, as amended (5 U.S.C. App), notice is hereby given of a meeting of the Board of Scientific Counselors, National Center for Biotechnology Information.

The meeting will be open to the public as indicated below, with attendance limited to space available. Individuals who plan to attend and need special assistance, such as sign language interpretation or other reasonable accommodations, should notify the Contact Person listed below in advance of the meeting.

The meeting will be closed to the public as indicated below in accordance with the provisions set forth in section 552b(c)(6), Title 5 U.S.C., as amended for review, discussion, and evaluation of individual intramural programs and projects conducted by the National Library of Medicine, including consideration of personnel qualifications and performance, and the competence of individual investigators, the disclosure of which would constitute a clearly unwarranted invasion of personal privacy.

Name of Committee: Board of Scientific Counselors, National Center for Biotechnology Information.

Date: April 12, 2011.

Open: 8:30 a.m. to 12 p.m.

Agenda: Program Discussion.

Place: National Library of Medicine, Building 38, 2nd Floor, Board Room, 8600 Rockville Pike, Bethesda, MD 20892.

Closed: 12 p.m. to 2 p.m.

Agenda: To review and evaluate personal qualifications and performance, and competence of individual investigators.

Place: National Library of Medicine, Building 38, 2nd Floor, Board Room, 8600 Rockville Pike, Bethesda, MD 20892.

Open: 2 p.m. to 3 p.m

Agenda: Program Discussion.

Place: National Library of Medicine, Building 38, 2nd Floor, Board Room, 8600 Rockville Pike, Bethesda, MD 20892.

Contact Person: David J. Lipman, MD, Director, National Center of Biotechnology Information, National Library of Medicine, Department of Health and Human Services, Building 38A, Room 8N805, Bethesda, MD 20892. 301–435–5985, *dlipman@mail.nih.gov*.

Any interested person may file written comments with the committee by forwarding the statement to the Contact Person listed on this notice. The statement should include the name, address, telephone number and when applicable, the business or professional affiliation of the interested person.

In the interest of security, NIH has instituted stringent procedures for entrance onto the NIH campus. All visitor vehicles, including taxicabs, hotel, and airport shuttles will be inspected before being allowed on campus. Visitors will be asked to show one form of identification (for example, a government-issued photo ID, driver's license, or passport) and to state the purpose of their visit.

(Catalogue of Federal Domestic Assistance Program No. 93.879, Medical Library Assistance, National Institutes of Health, HHS).

Dated: December 10, 2010.

Jennifer Spaeth,

Director, Office of Federal Advisory Committee Policy.

[FR Doc. 2010–31623 Filed 12–15–10; 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.

Proposed Project: 2011–2014 National Survey on Drug Use and Health: Methodological Field Tests (OMB No. 0930–0290)–Revision

The National Survey on Drug Use and Health (NSDUH) is a survey of the civilian, non-institutionalized population of the United States 12 years old and older. The data are used to determine the prevalence of use of tobacco products, alcohol, illicit substances, and illicit use of prescription drugs. The results are used by SAMHSA, ONDCP, Federal Government agencies, and other organizations and researchers to establish policy, direct program activities, and better allocate resources.

In March 2008, SAMHSA received a 3-year renewal of its generic clearance for methodological field tests. This will be a request for another renewal of the generic approval to continue methodological tests over the next 3 years, with conditions similar to the previous clearance. These methodological tests will continue to be designed to examine the feasibility, quality, and efficiency of new procedures or revisions to existing survey protocol. Specifically, the tests will measure the reliability and validity of certain questionnaire sections and items through multiple measurements on a set of respondents; assess new methods for gaining cooperation and participation of respondents with the goal of increasing response and decreasing potential bias in the survey estimates; and assess the impact of new sampling techniques and technologies on respondent behavior and reporting. Research will involve focus groups, cognitive laboratory testing, field tests, and customer surveys.

The next wave of methodological tests will continue to examine ways to increase data quality, lower operating costs, and gain a better understanding of

sources and effects of nonsampling error on the NSDUH estimates. Particular attention will be given to minimizing the impact of design changes so that survey data continue to remain comparable over time. If these tests provide successful results, current procedures or data collection instruments may be revised.

The number of respondents to be included in each field test will vary, depending on the nature of the subject being tested and the target population. However, the total estimated response burden is 8,251 hours. The exact number of subjects and burden hours for each test are unknown at this time, but will be clearly outlined in each individual submission. The table below, however, describes the anticipated burden for each of the major testing activities for which generic approval is being tested.

ESTIMATED BURDEN FOR NSDUH METHODOLOGICAL FIELD TESTS

Activity	Number of respondents	Responses per respondent	Total number of responses	Average burden per response (hrs.)	Total burden (hrs.)
a. Focus Groups	270	1	270	2.0	540
b. Cognitive laboratory testing	200	1	200	1.0	200
c. Field Tests	6,600	1	6,600	1.0	6,600
d. Customer Satisfaction Surveys	300	1	300	0.25	75
Household screening for c	8,910	1	8,910	0.083	740
Screening Verification for c	445	1	445	0.067	30
Interview Verification for c	990	1	990	0.067	66
Total	17,715	.....	17,715	.....	8,251
Annual Average (Total divided by 3 years)	5,905	.....	5,905	.....	2,750

Written comments and recommendations concerning the proposed information collection should be sent by January 18, 2011 to: SAMHSA Desk Officer, Human Resources and Housing Branch, Office of Management and Budget, New Executive Office Building, Room 10235, Washington, DC 20503; due to potential delays in OMB's receipt and processing of mail sent through the U.S. Postal Service, respondents are encouraged to submit comments by fax to: 202–395–7285.

Dated: December 1, 2010.

Elaine Parry,

Director, Office of Management, Technology and Operations.

[FR Doc. 2010–31585 Filed 12–15–10; 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: Screening, Brief Intervention, Brief Treatment and Referral to Treatment (SBIRT) Cross-Site Evaluation—New

SAMHSA is conducting a cross-site external evaluation of the impact of programs of screening, brief intervention (BI), brief treatment (BT) and referral to treatment on patients presenting at various health care delivery units with a continuum of severity of substance use. SAMHSA's SBIRT program is a cooperative agreement grant program designed to help States and Tribal Councils expand the continuum of care available for substance misuse and use disorders. The program includes screening, brief intervention, brief treatment and referrals to treatment for persons at risk for dependence on alcohol or drugs. The cross-site evaluation will provide a comprehensive assessment of the effects of SBIRT on patient outcomes, performance site practices, and treatment systems. This information will allow SAMHSA to determine the extent to which SBIRT has met its objectives of implementing a comprehensive system of identification and care to meet the needs of individuals at all points along the substance use continuum.

A paper and pencil survey will be administered to practitioners in sites where SBIRT services are being delivered. The practitioner survey is designed to evaluate the implementation of proposed SBIRT models by measuring their penetration and practitioners' willingness to adopt. Furthermore, the survey will document moderating factors related to practitioner and health care delivery unit characteristics.

The 93 question practitioner survey includes collection of demographic information as well as questions that attempt to assess barriers to implementation encountered by the practitioners and to gauge the effectiveness of the training they received. These measures were developed and used by Babor et al. (2005) in their comparable study comparing different implementation strategies for primary care screening and brief intervention programs for hazardous and harmful drinkers. The practitioner survey also includes an instrument developed by Panzano and Roth (2006) to measures an organization's willingness to adopt new innovative practices.

TOTAL BURDEN HOURS FOR THE CROSS-SITE PATIENT SURVEY

Instrument/activity	Number of respondents	Responses per respondent	Hours per response	Total burden hours	Hourly wage	Total respondent cost [a]
Practitioner Survey	1,075	1	.30	322.5	$32	$10,320