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

Contact Person: Abubakar A. Shaikh, PhD, DVM, Scientific Review Administrator, Center for Scientific Review, National Institutes of Health, 6701 Rockledge Drive, Room 6168, MSC 7892, Bethesda, MD 20892, (301) 435–1042, *shaikha@csr.nih.gov.*

Name of Committee: Center for Scientific Review Special Emphasis Panel, SEP to Review AOIC Member Conflicts.

Date: November 24, 2004.

Time: 11 a.m. to 3 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: Ranga V. Srinivas, PhD, Scientific Review Administrator, Center for Scientific Review, National Institutes of Health, 6701 Rockledge Drive, Room 5222, MSC 7852, Bethesda, MD 20892, (301) 435–1167, *srinivar@csr.nih.gov.*

Name of Committee: Center for Scientific Review Special Emphasis Panel High Resolution Electron Microscopy.

Date: November 29, 2004.

Time: 1 p.m. to 4 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: Richard D. Rodewald, PhD, Scientific Review Administrator, Center for Scientific Review, National Institutes of Health, 6701 Rockledge Drive, Room 5142, MSC 7840, Bethesda, MD 20892, (301) 435–1024, *rodewalr@csr.nih.gov.*

Name of Committee: Center for Scientific Review Special Emphasis Panel Neural Engineering.

Date: December 1, 2004.

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

Agenda: To review and evaluate grant applications.

Place: Sofitel Lafayette Square Washington, 806 15th Street, NW., Washington, DC 20005.

Contact Person: Mary Custer, PhD, Scientific Review Administrator, Center for Scientific Review, National Institutes of Health, 6701 Rockledge Drive, Room 4136, MSC 7850, Bethesda, MD 20892, (301) 435–1164, *custerm@csr.nih.gov.*

Name of Committee: Center for Scientific Review Special Emphasis Panel Pelvic Floor Physiology.

Date: December 1, 2004.

Time: 12 p.m. to 1 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: M. Chris Langub, PhD, Scientific Review Administrator, Center for Scientific Review, National Institutes of Health, 6701 Rockledge Drive, Room 4112, MSC 7814, Bethesda, MD 20892, (301) 496–8551, *langubm@csr.nih.gov.*

Name of Committee: Center for Scientific Review Special Emphasis Panel PAR–03–137 Topical Microbicides.

Date: December 8–10, 2004.

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

Agenda: To review and evaluate grant applications.

Place: One Washington Circle Hotel, One Washington Circle, Washington, DC 20037.

Contact Person: Ranga V. Srinivas, PhD, Scientific Review Administrator, Center for Scientific Review, National Institutes of Health, 6701 Rockledge Drive, Room 5222, MSC 7852, Bethesda, MD 20892, (301) 435–1167, *srinivar@csr.nih.gov.*

Name of Committee: Center for Scientific Review Special Emphasis Panel ZRG1 ONC F (02) Gene Expression.

Date: December 13, 2004.

Time: 1:30 p.m. to 2:30 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: Marcia Litwack, PhD, Scientific Review Administrator, Center for Scientific Review, National Institutes of Health, 6701 Rockledge Drive, Room 6206, MSC 7804, Bethesda, MD 20892, (301) 435–1719, *litwackm@csr.nih.gov.*

Name of Committee: Center for Scientific Review Special Emphasis Panel, Member Conflict—Neurogenesis.

Date: December 14, 2004.

Time: 1 p.m. to 3 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: Carole L. Jelsema, PhD, Chief and Scientific Review Administrator, MDCN Scientific Review Group, Center for Scientific Review, National Institutes of Health, 6701 Rockledge Drive, Room 4146, MSC 7850, Bethesda, MD 20892, (301) 435–1248, *jelsemac@csr.nih.gov.*

Name of Committee: Center for Scientific Review Special Emphasis Panel, Pharmacogenetics and Bioinformatics.

Date: December 14–15, 2004.

Time: 5 p.m. to 4 p.m.

Agenda: To review and evaluate grant applications.

Place: Four Points by Sheraton Bethesda, 8400 Wisconsin Avenue, Bethesda, MD 20814.

Contact Person: Barbara Whitmarsh, PhD, Scientific Review Administrator, Center for Scientific Review, National Institutes of Health, 6701 Rockledge Drive, Room 2206, MSC 7890, Bethesda, MD 20892, (301) 435–4511, *whitmarshb@csr.nih.gov.*

Name of Committee: Center for Scientific Review Special Emphasis Panel, Angiotensin Receptors.

Date: December 17, 2004.

Time: 11 a.m. to 12 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: Joyce C. Gibson, DSC, Scientific Review Administrator, Center for Scientific Review, National Institutes of Health, 6701 Rockledge Drive, Room 4130, MSC 7814, Bethesda, MD 20892, (301) 435–4522, *gibsonj@csr.nih.gov.*

Name of Committee: Center for Scientific Review Special Emphasis Panel, ZRG1 RUS–E (02) Renal Urology Special Member Conflicts Meeting.

Date: December 20, 2004.

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

Agenda: To review and evaluate grant applications.

Place: Bethesda Marriott Suites, 6711 Democracy Boulevard, Bethesda, MD 20817.

Contact Person: Aftab A. Ansari, PhD, Health Scientist Administrator, National Institutes of Health, Center for Scientific Review, 6701 Rockledge Drive, Room 4108, MSC 7814, Bethesda, MD 20892, (301) 435–1173, *ansaria@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: November 8, 2004.

LaVerne Y. Stringfield,

Director, Office of Federal Advisory Committee Policy.

[FR Doc. 04–25346 Filed 11–15–04; 8:45 am]

BILLING CODE 4140–01–M

DEPARTMENT OF HEALTH AND HUMAN SERVICES

Substance Abuse and Mental Health Services Administration

Agency Information Collection Activities: Proposed Collection; Comment Request

In compliance with Section 3506(c)(2)(A) of the Paperwork Reduction Act of 1995 concerning opportunity for public comment on proposed collections of information, the Substance Abuse and Mental Health Services Administration will publish periodic summaries of proposed projects. To request more information on the proposed projects or to obtain a copy of the information collection plans, call the SAMHSA Reports Clearance Officer on (240) 276–1243.

Comments are invited on: (a) Whether the proposed collections of information are necessary for the proper performance of the functions of the agency, including whether the information shall have practical utility; (b) the accuracy of the agency's estimate of the burden of the proposed collection of information; (c) ways to enhance the quality, utility, and clarity of the information to be collected; and (d) ways to minimize the burden of the collection of information on respondents, including through the use of automated collection techniques or other forms of information technology.

National Survey on Drug Use and Health: Clinical Validation Study of the Substance Dependence and Abuse Measures—(OMB No. 0930–0231)—Revision

The Substance Abuse and Mental Health Services Administration's (SAMHSA) National Survey on Drug Use and Health (NSDUH), formerly the National Household Survey on Drug Abuse, is a survey of the civilian, noninstitutionalized 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, the Office of National Drug Control Policy, other Federal government agencies, and other organizations and researchers to establish policy, direct program activities, and better allocate resources.

From 2001–2004, the NSDUH conducted the first phase (phase 1) of a two-phase Clinical Validation Study of the Substance Dependence and Abuse Measures. From 2005–2007 the NSDUH plans to conduct the second phase (Phase 2) of this study. Specific aims of the two-phase study are to achieve the best overarching format, and the best wording and ordering for the assessment questions. The goal is improved validity and reduced respondent burden.

In phase 1 a field test was conducted. Half of all subjects in this field test were between 12 and 17, and half 18 years of age or older; subjects were recruited from the Research Triangle and the Triad areas of North Carolina through fliers and newspaper ads, and asked (1) demographic information and (2) questions from two self-administered sections of the NSDUH questionnaire: questions about the quantity and frequency of use of drugs and alcohol, and questions about symptoms of substance dependence and abuse. A semi-structured clinical interview was administered to these same subjects by a trained clinician to determine the presence or absence of substance dependence and abuse. The clinical instruments used to assess subjects were the substance abuse modules from the Structured Clinical Interview for DSM–IV (SCID) (for adults) and the Kiddie Schedule for Affective Disorders and Schizophrenia (K–SADS) (for those between 12 and 17 years of age). The correspondence of the diagnosis of substance dependence and abuse between the clinical and survey interview were compared.

The results of this comparison from the field test in phase 1 indicated a lack of sufficient correspondence between the clinical and survey interview. Overall there was a strong tendency toward over reporting in the survey interview. Recommendations were made to revise specific questions. Problems with specific questions were identified and reasons for lack of correspondence were examined. Modifications to the NSDUH questions on substance dependence and abuse to achieve better correspondence are being made for phase 2 of the study. In order to reduce a tendency to over report some questions have been revised to be more specific.

In Phase 2, a second clinical validation study will be conducted using the same procedures as Phase 1 but with the revised questions on dependence or abuse. This will allow a determination of the correspondence (kappa) between the revised diagnosis obtained from the NSDUH substance dependence and abuse module and the diagnosis from the structured clinical interviews. Final revisions to the survey instrument will be made based on findings from Phase 2. All decisions about final revisions to the module will balance the need for correspondence across different groups. The following table summarizes the burden associated with phase two of the project.

Phase II	Number of respondents	Responses per respondent	Hours per response	Total burden
Adults:				
Screening	400	1	.08	32
Screener and interview	200	1	1.5	300
Adolescents:				
Screening	200	1	1.5	300
Screener and interview	170	1	1.50	255
Total	370	.....	.....	887

Send comments to Summer King, SAMHSA Reports Clearance Officer, Room 7–1044, 1 Choke Cherry Road, Rockville, MD 20850. Written comments should be received by January 18, 2005.

Dated: November 9, 2004.

Anna Marsh,

Executive Officer, SAMHSA.

[FR Doc. 04–25365 Filed 11–15–04; 8:45 am]

BILLING CODE 4162–20–P

DEPARTMENT OF HOMELAND SECURITY

Coast Guard

[USCG–2004–19591]

Collection of Information Under Review by Office of Management and Budget (OMB): OMB Control Numbers: 1625–0005, 1625–0020, 1625–0029, 1625–0031, 1625–0085, and 1625–0096

AGENCY: Coast Guard, DHS.

ACTION: Request for comments.

SUMMARY: In compliance with the Paperwork Reduction Act of 1995, the Coast Guard intends to seek the approval of OMB for the renewal or revision of six Information Collection Requests (ICRs). The ICRs comprise (1) 1625–0005, Application and Permit to Handle Hazardous Materials; (2) 1625–0020, Security Zones, Regulated Navigation Areas, and Safety Zones; (3) 1625–0029, Self-propelled Liquefied Gas Vessels; (4) 1625–0031, Plan Approval and Records for Electrical Engineering Regulations—title 46 CFR subchapter J; (5) 1625–0085, Streamlined Inspection Program; and (6) 1625–0096, Report of Oil or Hazardous Substance Discharge, and Report of Suspicious Maritime Activity. Before submitting the ICRs to OMB, the Coast Guard is inviting comments on them as described below.

DATES: Comments must reach the Coast Guard on or before January 18, 2005.

ADDRESSES: To make sure that your comments and related material do not