

proposed project or to obtain a copy of the data collection plans and draft instruments, email [paperwork@hrsa.gov](mailto:paperwork@hrsa.gov) or call Lisa Wright-Solomon, the HRSA Information Collection Clearance Officer at (301) 443-1984.

**SUPPLEMENTARY INFORMATION:** When submitting comments or requesting information, please include the information request collection title for reference.

*Information Collection Request Title:* Radiation Exposure Screening and Education Program, OMB No. 0906-0012—Revision.

*Abstract:* The Radiation Exposure Screening and Education Program (RESEP) is authorized by section 417C of the Public Health Service Act (42 U.S.C. 285a-9). The purpose of RESEP is to assist individuals who live (or lived) in areas where U.S. nuclear weapons testing occurred and who are diagnosed with cancer and other radiogenic diseases caused by exposure to nuclear fallout or nuclear materials such as uranium. RESEP funds support eligible health care organizations in implementing cancer screening programs; developing education programs; disseminating information on

radiogenic diseases and the importance of early detection; screening eligible individuals for cancer and other radiogenic diseases; providing appropriate referrals for medical treatment; and facilitating documentation of radiation exposure.

*Need and Proposed Use of the Information:* For this program, performance measures were drafted to provide data useful to the program and to enable HRSA to provide aggregate program data required by Congress under the Government Performance and Results Act of 1993 (Pub. L. 103-62). These measures cover the principal topic areas of interest to the Federal Office of Rural Health Policy (FORHP), including demographics for the RESEP program user population, medical screening activities for cancers and other radiogenic diseases, exposure and presentation types for eligible radiogenic malignant and nonmalignant diseases, referrals for appropriate medical treatment, eligibility counseling and referral assistance for the Radiation Exposure Compensation Act, and program outreach and education activities. These measures speak to FORHP's progress toward meeting the

established goals. In order to reduce the reporting burden by the award recipients, a number of questions have been removed with the new set of measures reflecting an effort to streamline data collection and collect consistent and uniform measures across FORHP's grant programs.

*Likely Respondents:* Radiation Exposure Screening and Education Program award recipients.

*Burden Statement:* Burden in this context means the time expended by persons to generate, maintain, retain, disclose, or provide the information requested. This includes the time needed to: (1) Review instructions; (2) develop, acquire, install, and utilize technology and systems for the purpose of collecting, validating and verifying information, processing and maintaining information, and disclosing and providing information; (3) train personnel and to be able to respond to a collection of information; (4) search data sources; to complete and review the collection of information; and (5) transmit or otherwise disclose the information. The total annual burden hours estimated for this ICR are summarized in the table below.

TOTAL ESTIMATED ANNUALIZED BURDEN HOURS

Form name	Number of respondents	Number of responses per respondent	Total responses	Average burden per response (in hours)	Total burden hours
Radiation Exposure Screening and Education Program .....	8	1	8	12	96
	8	.....	8	.....	96

HRSA specifically requests comments on: (1) The necessity and utility of the proposed information collection for the proper performance of the agency's functions; (2) the accuracy of the estimated burden; (3) ways to enhance the quality, utility, and clarity of the information to be collected; and (4) the use of automated collection techniques or other forms of information technology to minimize the information collection burden.

**Amy McNulty,**

*Acting Director, Division of the Executive Secretariat.*

[FR Doc. 2018-00145 Filed 1-8-18; 8:45 am]

**BILLING CODE 4165-15-P**

**DEPARTMENT OF HEALTH AND HUMAN SERVICES**

**National Institutes of Health**

**National Institute of Dental & Craniofacial Research; Notice of Meeting**

Pursuant to section 10(d) of the Federal Advisory Committee Act, as amended, notice is hereby given of a meeting of the National Advisory Dental and Craniofacial Research Council.

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 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 Advisory Dental and Craniofacial Research Council.

*Date:* January 31, 2018.

*Open:* 8:30 a.m. to 12:00 p.m.

*Agenda:* Report to the Director, NIDCR.

*Place:* National Institutes of Health, Building 31, 6th Floor, Conference Room 6, 31 Center Drive, Bethesda, MD 20892.

*Closed:* 1:30 p.m. to 3:00 p.m.

*Agenda:* To review and evaluate grant applications.

*Place:* National Institutes of Health, Building 31, 6th Floor, Conference Room 6, 31 Center Drive, Bethesda, MD 20892.

*Contact Person:* Alicia J. Dombroski, Ph.D., Director, Division of Extramural Activities,

National Institute of Dental and Craniofacial Research, National Institutes of Health, Bethesda, MD 20892, 301-594-4805, [adombroski@nidcr.nih.gov](mailto:adombroski@nidcr.nih.gov).

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.

Information is also available on the Institute's/Center's home page: <http://www.nidcr.nih.gov/about>, where an agenda and any additional information for the meeting will be posted when available.

(Catalogue of Federal Domestic Assistance Program Nos. 93.121, Oral Diseases and Disorders Research, National Institutes of Health, HHS)

Dated: January 2, 2018.

**Natasha M. Copeland,**

*Program Analyst, Office of Federal Advisory Committee Policy.*

[FR Doc. 2018-00155 Filed 1-8-18; 8:45 am]

**BILLING CODE 4140-01-P**

## DEPARTMENT OF HEALTH AND HUMAN SERVICES

### National Institutes of Health

#### National Institute of Biomedical Imaging and Bioengineering; Notice of Closed Meeting

Pursuant to section 10(d) of the Federal Advisory Committee Act, as amended, notice is hereby given of the meeting of the National Institute of Biomedical Imaging and Bioengineering Special Emphasis Panel.

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 Biomedical Imaging and Bioengineering Special Emphasis Panel; MSM Program Review (2018/05).

*Date:* February 12, 2018.

*Time:* 5:00 p.m. to 12:00 p.m.

*Agenda:* To review and evaluate grant applications.

*Place:* National Institutes of Health, Two Democracy Plaza, 6707 Democracy Plaza, Suite 920, Bethesda, MD 20892 (Virtual Meeting).

*Contact Person:* Manana Sukhareva, Ph.D., Scientific Review Officer, National Institutes of Biomedical Imaging and Bioengineering, National Institutes of Health, 6707 Democracy Boulevard, Suite 959, Bethesda, MD 20892, (301) 451-3397, [sukharem@mail.nih.gov](mailto:sukharem@mail.nih.gov).

Dated: January 2, 2018.

**Melanie J. Pantoja,**

*Program Analyst, Office of Federal Advisory Committee Policy.*

[FR Doc. 2018-00154 Filed 1-8-18; 8:45 am]

**BILLING CODE 4140-01-P**

## DEPARTMENT OF HEALTH AND HUMAN SERVICES

### Substance Abuse and Mental Health Services Administration

#### Confidentiality of Substance Use Disorder Patient Records

**AGENCY:** Substance Abuse and Mental Health Services Administration, HHS.

**ACTION:** Notice of public meeting.

**SUMMARY:** The Substance Abuse and Mental Health Services Administration (SAMHSA) announces that it will hold a public listening session on Wednesday, January 31, 2018, to solicit information concerning the Confidentiality of Substance Use Disorder Patient Records regulations as required by Section 11002 of the 21st Century Cures Act. The listening session will provide an opportunity for the public to provide input to SAMHSA concerning the effect of part 2 on "patient care, health outcomes, and patient privacy" as well as potential regulatory changes and future subregulatory guidance.

**DATES:** The listening session will be held on Wednesday, January 31, 2018, from 8:30 a.m. (Eastern) to 1:00 p.m. (Eastern).

**FOR FURTHER INFORMATION CONTACT:** For information concerning the listening session, please contact Rachel Karton, Senior Legislative and Regulatory Analyst, SAMHSA, 5600 Fishers Lane, Rockville, MD 20857, (240) 276-0416 or email [PrivacyRegulations@SAMHSA.hhs.gov](mailto:PrivacyRegulations@SAMHSA.hhs.gov).

#### SUPPLEMENTARY INFORMATION:

*Participation:* The Listening Session proceeding will be recorded, and subsequently archived and posted on the SAMHSA website. The public may attend the listening session:

- *Via Teleconference/Webcast:* The entire proceeding will be streamed live over the internet (requires prior registration). Audio and streaming information will be sent to those who register prior to the meeting. Capacity

for the Teleconference/Webcast participation is limited so early registration is recommended.

- *In Person:* The address for this meeting is 5600 Fishers Lane, 5th (Main) Floor Pavilion rooms, Rockville, Maryland 20852. The building is a federal facility; prior registration, a security screening and a federally-approved identification (e.g., driver's license) are required to attend in-person. Capacity for in-person attendance is limited so early registration is recommended.

*Registration:* Registration is required for participation in the listening session in person or via Teleconference/Webcast. Registration is now open. Registration for the in-person session will close on 01/22/2018 at 12:00 p.m. Eastern Time (ET). Registration for the Teleconference/Webcast will close on 01/31/2018 at 8:30 a.m. ET. Persons registering should indicate if they wish to make a public comment. SAMHSA recommends that when commenters suggest changes or revisions to current regulations that they indicate specifically, when feasible, how such regulation text should be revised. Only one representative of an organization may be allowed to present oral comments. Presentations will be limited to three minutes per speaker. SAMHSA will try to accommodate all speakers who wish to present based on the time allotted for this meeting. Persons making oral presentations are encouraged to also submit written comments as discussed below.

To register, go to: <https://42-cfr-part2-listening-session.eventbrite.com>.

*Special Assistance:* Individuals who plan to attend and need special assistance, such as sign language interpretation or other reasonable accommodations, should notify Rachel Karton (contact information provided below) at least 10 days prior to the meeting.

*Public Comments:* In addition to attending the session in person or joining via Teleconference/Webcast, the Agency offers several ways to provide comments. SAMHSA recommends that when commenters suggest changes or revisions to current regulations that they indicate specifically, when feasible, how such regulation text should be revised. You may provide comments through the following means:

- *Electronically:* [PrivacyRegulations@SAMHSA.hhs.gov](mailto:PrivacyRegulations@SAMHSA.hhs.gov) (preferred).
- *Regular, Express or Overnight Mail, or Hand Delivery or Courier:* Written comments must be sent to the following address ONLY: Substance Abuse and Mental Health Services Administration (SAMHSA), Department of Health and