• Is the Solution easy to manage (ease of use, ease of data movement, user friendly)?

Viability

• Does the Solution present a deep understanding of the market for the Solution?
• Is there a clear advantage that differentiates this Solution from others?
• Is the Solution a model for real world implementation practical?
• Is the Solution economically viable and scalable/replicable?
• Are consumers and/or providers already participating (*e.g.* have signed up to test the Solution)?

Impact

• Does the participant present a theory or explanation of how the proposed Solution would improve the future of consumer-mediated health information sharing?
• Is there clear evidence of a health care need based on research for a specific consumer population, and is there evidence that Solution impacts this population?
• Could the Solution improve the experience of information sharing between consumers and their health care providers?
• Is the Solution's design human-centered so that it enables the consumer to understand and manage their health?

Phase 3

Impact

• Do the results indicate how the Solution will enable consumers to share data in a ''real-life'' setting?
• Does the Solution improve the experience of information sharing between consumers and their health care providers?

Deployability

• Is the Solution readily available to consumers to be used on existing mobile platforms or a public facing Web site?
• Is the Solution designed for ease of learning and ease of use by the target user population?

Scalability

• How scalable is the Solution in a real-world setting? How likely are cost efficiencies for delivery at greater scale?
• Is the user experience optimized for the greater population of consumers and/or providers?
• Is there a plan for getting consumers and/or providers to adopt and use the Solution?

Additional Information

General Conditions: ONC reserves the right to cancel, suspend, and/or modify the Challenge, or any part of it, for any reason, at ONC's sole discretion.

Intellectual Property: Each participant retains title and full ownership in and to their Submission. Participants expressly reserve all intellectual property rights not expressly granted under the challenge agreement. By participating in the Challenge, each entrant hereby irrevocably grants to the Government a limited, non-exclusive, royalty-free, perpetual, worldwide license and right to reproduce, publically perform, publically display, and use the Submission to the extent necessary to administer the challenge, and to publically perform and publically display the Submission, including, without limitation, for advertising and promotional purposes relating to the Challenge. This may also include displaying the results of the Challenge on a public Web site or during a public presentation.

Representation, Warranties and Indemnification

By entering the Challenge, each applicant represents, warrants and covenants as follows:

(a) Participant is the sole author, creator, and owner of the Submission;

(b) The Submission is not the subject of any actual or threatened litigation or claim;

(c) The Submission does not and will not violate or infringe upon the intellectual property rights, privacy rights, publicity rights, or other legal rights of any third party;

(d) The Submission does not and will not contain any harmful computer code (sometimes referred to as ''malware,'' ''viruses'' or ''worms''); and

(e) The Submission, and participants' use of the Submission, does not and will not violate any applicable laws or regulations, including, without limitation, HIPAA, applicable export control laws and regulations of the U.S. and other jurisdictions.

If the Submission includes any third party works (such as third party content or open source code), participant must be able to provide, upon request, documentation of all appropriate licenses and releases for such third party works. If participant cannot provide documentation of all required licenses and releases, the Federal Agency sponsor reserves the right, at their sole discretion, to disqualify the applicable Submission. Conversely, they may seek to secure the licenses and releases and allow the applicable Submission to remain in the Challenge, while reserving all rights with respect to such licenses and releases.

Participants must indemnify, defend, and hold harmless the Federal Government from and against all third party claims, actions, or proceedings of any kind and from any and all damages, liabilities, costs, and expenses relating to or arising from participant's Submission or any breach or alleged breach of any of the representations, warranties, and covenants of participant hereunder. The Federal Agency sponsors reserve the right to disqualify any Submission that, in their discretion, deems to violate these Official Rules, Terms & Conditions.

Authority: 15 U.S.C. 3719.

Karen DeSalvo,

National Coordinator for Health Information Technology.

[FR Doc. 2016–11102 Filed 5–9–16; 8:45 am]

BILLING CODE 4150–45–P

DEPARTMENT OF HEALTH AND HUMAN SERVICES

Office of the Secretary

[Document Identifier: HHS–OS–0990–new–60D]

Agency Information Collection Activities; Proposed Collection; Public Comment Request

AGENCY: Office of the Assistant Secretary for Health, Office of Adolescent Health, HHS.

ACTION: Notice.

SUMMARY: In compliance with section 3506(c)(2)(A) of the Paperwork Reduction Act of 1995, the Office of the Secretary (OS), Department of Health and Human Services, announces plans to submit a Information Collection Request (ICR), described below, to the Office of Management and Budget (OMB). Prior to submitting that ICR to OMB, OS seeks comments from the public regarding the burden estimate, below, or any other aspect of the ICR.

DATES: Comments on the ICR must be received on or before July 11, 2016.

ADDRESSES: Submit your comments to *Information.CollectionClearance@hhs.gov* or by calling (202) 690–6162.

FOR FURTHER INFORMATION CONTACT: Information Collection Clearance staff, *Information.CollectionClearance@hhs.gov* or (202) 690–6162.

SUPPLEMENTARY INFORMATION: When submitting comments or requesting information, please include the document identifier HHS–OS–0990–new–60D for reference.

Information Collection Request Title: Federal Evaluation of Making Proud Choices! (MPC!)

Abstract: The Office of Adolescent Health (OAH), U.S. Department of Health and Human Services (HHS) is requesting approval by OMB for a new data collection. The Federal Evaluation of Making Proud Choices (MPC!) will provide information about program design, implementation, and impacts through a rigorous assessment of program of a highly popular teen pregnancy prevention curriculum—MPC; it includes the baseline survey instrument related to the impact study and instruments for the implementation and fidelity assessment. The evaluation will be conducted in 39 schools nationwide. The data collected from these instruments will be used to describe the characteristics of the study sample of youth, being used in the models for estimating program impacts, and provide a detailed understanding of program implementation.

Need and Proposed Use of the Information: The baseline survey data will be used to describe the study sample and to assess whether random assignment successfully generated treatment and control groups balanced on important baseline characteristics. The findings from these analyses of program impacts and implementation will be of interest to the general public, to policymakers, and to schools and other organizations interested in supporting a comprehensive approach to teen pregnancy prevention.

Likely Respondents: The baseline data will be collected through a Web based survey with study participants in the participating evaluation schools. Study participants will primarily be in 8th or 9th grade at the time of the baseline survey, and will be enrolled in the schools' mandatory health class.

TOTAL ESTIMATED ANNUALIZED BURDEN—HOURS

Form name	Number of respondents	Number of responses per respondent	Average burden per response (in hours)	Total burden hours
Baseline survey of impact study participants	865	1	30/60	432.5
Master topic guide for staff interviews	39	1	1	39
Staff survey	26	1	30/60	13
Program attendance data and collection protocol	13	14	9/60	27.3
Program fidelity checklist	9	14	15/60	31.5
Youth focus group	87	1	1	87
Total				630.5

OS 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.

Terry S. Clark,

Asst Information Collection Clearance Officer.

[FR Doc. 2016–11069 Filed 5–10–16; 8:45 am]

BILLING CODE 4168–11–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 (5 U.S.C. App.), 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 Institute of Biomedical Imaging and Bioengineering Special Emphasis Panel, Health Disparity SBIR Review (2016/10).

Date: June 29, 2016.

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

Agenda: To review and evaluate grant applications.

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

Contact Person: Dennis Hlasta, Ph.D., Scientific Review Officer, National Institute of Biomedical Imaging and Bioengineering, National Institutes of Health, 6707 Democracy Blvd., Bethesda, MD 20892, (301) 451–4794, *dennis.hlasta@mail.nih.gov.*

Dated: May 5, 2016.

David Clary,

Program Analyst, Office of Federal Advisory Committee Policy.

[FR Doc. 2016–11023 Filed 5–10–16; 8:45 am]

BILLING CODE 4140–01–P

DEPARTMENT OF HEALTH AND HUMAN SERVICES

National Institutes of Health

National Institute of Mental Health; Notice of Closed Meeting

Pursuant to section 10(d) 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 Institute of Mental Health.

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 the review, discussion, and evaluation of individual intramural programs and projects conducted by the NATIONAL INSTITUTE OF MENTAL HEALTH, 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 Institute of Mental Health.

Date: June 6–8, 2016.

Time: June 06, 2016, 1:00 p.m. to 4:30 p.m.

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

Place: National Institutes Of Health, Porter Neuroscience Research Center GE 620/630/