

plans and instruments, contact: Nina Goodman, Public Health Advisor, Office of Communications and Education (OCE), NCI, NIH, 6116 Executive Blvd., Suite 400, Rockville, MD 20892, call non-toll-free number (301) 435-7789 or email your request, including your address to: goodmann@mail.nih.gov. Formal requests for additional plans and instruments must be requested in writing.

Proposed Collection: A Generic Submission For Formative Research, Pretesting, and Customer Satisfaction of NCI's Communication and Education Resources, 0925-0046, Expiration Date 2/28/2013, Reinstatement without Change, National Cancer Institute (NCI), National Institutes of Health (NIH).

Need and Use of Information Collection: In order to carry out NCI's legislative mandate to educate and disseminate information about cancer prevention, detection, diagnosis, and treatment to a wide variety of audiences and organizations, it is beneficial for NCI through its Office of Communications and Education (OCE), to pretest NCI communications

strategies, concepts, and messages while they are under development. This pretesting, or formative evaluation, helps ensure that the messages, communication materials, and information services created by NCI have the greatest capacity of being received, understood, and accepted by their target audiences. Since NCI's OCE is also responsible for the design, implementation, and evaluation of education programs over the entire cancer continuum, and management of NCI initiatives that address specific challenges in cancer research and treatment, it is also necessary to ensure that customers are satisfied with programs. This customer satisfaction research helps ensure the relevance, utility, and appropriateness of the many educational programs and products that OCE and NCI produce. OCE will use a variety of qualitative (focus groups, interviews) and quantitative (paper, phone, in-person, and web surveys) methodologies to conduct this formative and customer satisfaction research, allowing NCI to: (1) Understand

characteristics (attitudes, beliefs, and behaviors) of the intended target audience and use this information in the development of effective communication tools and strategies; (2) use a feedback loop to help refine, revise, and enhance messages, materials, products, and programs—ensuring that they have the greatest relevance, utility, appropriateness, and impact for/to target audiences; and (3) expend limited program resource dollars wisely and effectively. The participants may include, but are not limited to, cancer patients, their families, the general public, health providers, the media, voluntary groups, scientific and medical organizations (affected public could include individuals or households; businesses or other for profit; not-for-profit institutions; and Federal Government; State, Local, or Tribal Government).

OMB approval is requested for 3 years. There are no costs to respondents other than their time. The total estimated burden, over three years for this generic request are 6,600 hours.

3-YEAR ESTIMATED BURDEN HOURS (GENERIC REQUEST)

Category of respondents	Form name	Number of respondents	Frequency of response per respondent	Time per response (in hours)	Burden hours
Individuals, Households, Local, State, and Federal Governments, and Private Sector.	Focus Groups, Individual In-Depth Interviews, Brief Interviews, Surveys, Website Usability Testing.	33,000	1	12/60	6,600

Dated: February 27, 2013.

Vivian Horovitch-Kelley,

NCI Project Clearance Liaison, NCI, NIH.

[FR Doc. 2013-05164 Filed 3-5-13; 8:45 am]

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DEPARTMENT OF HEALTH AND HUMAN SERVICES

National Institutes of Health

Eunice Kennedy Shriver National Institute of Child Health & Human Development; 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 Child Health and Human Development Special Emphasis Panel; Multigenic Disease Models.

Date: March 20, 2013.

Time: 9:00 a.m. to 11:00 a.m.

Agenda: To review and evaluate grant applications.

Place: National Institutes of Health, 6100 Executive Boulevard, Rockville, MD 20852, (Telephone Conference Call).

Contact Person: Cathy J. Wedeen, Ph.D., Scientific Review Officer, Division of Scientific Review, OD, Eunice Kennedy Shriver National Institute of Child Health and Human Development, NIH, 6100 Executive Blvd., Room 5B01-G, Bethesda, MD 20892, 301-435-6878, wedeenc@mail.nih.gov.

(Catalogue of Federal Domestic Assistance Program Nos. 93.864, Population Research; 93.865, Research for Mothers and Children; 93.929, Center for Medical Rehabilitation Research; 93.209, Contraception and

Infertility Loan Repayment Program, National Institutes of Health, HHS)

Dated: February 28, 2013.

Michelle Trout,

Program Analyst, Office of Federal Advisory Committee Policy.

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