

**III. Comments**

Interested persons may submit either electronic comments regarding this document to <http://www.regulations.gov> or written comments to the Division of Dockets Management (see **ADDRESSES**). It is only necessary to send one set of comments. Identify comments with the docket number found in brackets in the heading of this document. Received comments may be seen in the Division of Dockets Management between 9 a.m. and 4 p.m., Monday through Friday, and will be posted to the docket at <http://www.regulations.gov>.

Dated: October 2, 2014.

**Leslie Kux,**

*Assistant Commissioner for Policy.*

[FR Doc. 2014–23965 Filed 10–7–14; 8:45 am]

**BILLING CODE 4164–01–P**

## DEPARTMENT OF HEALTH AND HUMAN SERVICES

### National Institutes of Health

#### Submission for OMB Review; 30-day Comment Request; a Generic Submission for Theory Development and Validation (NCI)

**SUMMARY:** Under the provisions of Section 3507(a)(1)(D) of the Paperwork Reduction Act of 1995, the National Institutes of Health (NIH), has submitted to the Office of Management and Budget (OMB) a request for review and approval of the information collection listed below. This proposed information collection was previously published in the **Federal Register** on July 14, 2014,

Vol. 79, page 40763 and allowed 60-days for public comment. No public comments were received. The purpose of this notice is to allow an additional 30 days for public comment. The National Cancer Institute (NCI), National Institutes of Health, may not conduct or sponsor, and the respondent is not required to respond to, an information collection that has been extended, revised, or implemented on or after October 1, 1995, unless it displays a currently valid OMB control number.

**Direct Comments to OMB:** Written comments and/or suggestions regarding the item(s) contained in this notice, especially regarding the estimated public burden and associated response time, should be directed to the: Office of Management and Budget, Office of Regulatory Affairs, [OIRA\\_submission@omb.eop.gov](mailto:OIRA_submission@omb.eop.gov) or by fax to 202–395–6974, Attention: NIH Desk Officer.

**Comment Due Date:** Comments regarding this information collection are best assured of having their full effect if received within 30-days of the date of this publication.

**FOR FURTHER INFORMATION CONTACT:** To obtain a copy of the data collection plans and instruments or request more information on the proposed project contact: Rebecca A. Ferrer, Division of Cancer Control and Population Sciences, 9609 Medical Center Dr., Room 3E114, Bethesda, MD 20892 or call non-toll-free number 240–276–6914 or Email your request, including your address to: [ferrerra@mail.nih.gov](mailto:ferrerra@mail.nih.gov). Formal requests for additional plans and instruments must be requested in writing.

**Proposed Collection:** A Generic Submission for Theory Development and Validation (NCI), Revision, National Cancer Institute (NCI), National Institutes of Health (NIH).

**Need and Use of Information Collection:** The National Cancer Institute is requesting approval for this revised generic clearance to conduct formative research related to behavioral science theory development and validation for the next three years. Formative research in the area of theory development and validation would provide the basis for developing effective cancer prevention and control strategies, allow for a better understanding of theoretical constructs that influence decisions and actions related to cancer, and ultimately contribute to reducing the U.S. cancer burden. Sub-studies proposed under this generic clearance would involve methodological testing and a standard set of research approaches, including surveys (internet, phone, and paper-and-pencil) and focus groups. Respondents would include individuals in the general public, recruited through established online panels or internet/newspaper advertisements. Development of each study or survey would involve consulting with NCI scientists as well as experts from the behavioral science research community.

OMB approval is requested for 3 years. There are no costs to respondents other than their time. The total estimated burden is 2,000 hours per year.

Estimated Burden Hours for Three Years

TABLE A.12–1—ESTIMATES OF ANNUAL BURDEN HOURS

Type of respondents	Number of respondents	Frequency of response	Average time per response (minutes/hour)	Total burden hours
General Public .....	667	1	15/60	167
Physicians .....	2,000	1	30/60	1,000
Health Professionals .....	333	1	1	333
Researchers .....	333	1	90/60	500

Dated: October 1, 2014.

**Karla Bailey,**

*NCI Project Clearance Liaison, National Institutes of Health.*

[FR Doc. 2014–23999 Filed 10–7–14; 8:45 am]

**BILLING CODE 4140–01–P**

## DEPARTMENT OF HEALTH AND HUMAN SERVICES

### National Institutes of Health

#### Proposed Collection; Comment Request; Application Forms for Research Development and Training Grants

**SUMMARY:** In compliance with the requirement of Section 3506(c)(2)(A) of the Paperwork Reduction Act of 1995, for opportunity for public comment on proposed data collections via

application forms, the National Institute on Drug Abuse (NIDA), the National Institutes of Health (NIH) will publish periodic summaries of proposed collections to be submitted to the Office of Management and Budget (OMB) for review and approval.

Written comments and/or suggestions from the public and affected agencies are invited on one or more of the following points: (1) Whether the proposed collection of information is necessary for the proper performance of the function of the agency, including