following points: (1) Whether the proposed collection of information is necessary for the proper performance of the function of the agency, including whether the information will have practical utility; (2) The accuracy of the agency's estimate of the burden of the proposed collection of information, including the validity of the methodology and assumptions used; (3) Ways to enhance the quality, utility, and clarity of the information to be collected; and (4) Ways to minimize the burden of the collection of information on those who are to respond, including the use of appropriate automated, electronic, mechanical, or other technological collection techniques or other forms of information technology.

To Submit Comments and For Further Information: To obtain a copy of the data collection plans and instruments, submit comments in writing, 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.* Formal requests for additional plans and instruments must be requested in writing.

DATES: *Comment Due Date:* Comments regarding this information collection are best assured of having their full effect if received within 60 days of the date of this publication.

Proposed Collection: A Generic Submission for Theory Development and Validation (NCI), Revision, 0925–0645, National Cancer Institute (NCI), National Institutes of Health (NIH).

Need and Use of Information Collection: The National Cancer Institute is requesting terms of clearance and 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.

There are no costs to respondents other than their time. The total estimated burden is 6,500 hours.

ESTIMATED BURDEN HOURS FOR THREE YEARS

Type of respondents	Number of respondents	Number of responses per respondent	Average burden per response (in hours)	Total burden hours
General Public	2,000	1	15/60	500
Physicians	6,000	1	30/60	3,000
Health Professionals	1,000	1	1	1,000
And Researchers	1,000	1	2	2,000

Dated: July 8, 2014.

Karla Bailey,

NCI Project Clearance Liaison, National Institutes of Health.

[FR Doc. 2014–16447 Filed 7–11–14; 8:45 am]

BILLING CODE 4140–01–P

DEPARTMENT OF HEALTH AND HUMAN SERVICES

National Institutes of Health

Draft Report on Carcinogens Monograph on Trichloroethylene; Amended Notice

SUMMARY: The notice amends the **Federal Register** notice, 79 FR 33203, published June 10, 2014, announcing availability of documents, request for comments, and notice of meeting to peer review the Draft Report on Carcinogens (RoC) Monograph on Trichloroethylene (TCE). The deadline for written public comment submissions has been extended to August 4, 2014. All other information in the original notice has not changed. Information about the meeting and registration is available at *http://ntp.niehs.nih.gov/go/38853*.

DATES: *Written Public Comments Submissions:* Deadline is August 4, 2014.

Dated: July 7, 2014.

John R. Bucher,

Associate Director, National Toxicology Program.

[FR Doc. 2014–16449 Filed 7–11–14; 8:45 am]

BILLING CODE 4140–01–P

DEPARTMENT OF HEALTH AND HUMAN SERVICES

National Institutes of Health

Scientific Advisory Committee on Alternative Toxicological Methods; Announcement of Meeting; Request for Comments

SUMMARY: This notice announces a meeting of the Scientific Advisory Committee on Alternative Toxicological Methods (SACATM). SACATM advises the Interagency Coordinating Committee on the Validation of Alternative Methods (ICCVAM), the National Toxicology Program (NTP) Interagency Center for the Evaluation of Alternative Toxicological Methods (NICEATM), and the Director of the National Institute of Environmental Health Sciences (NIEHS) and NTP regarding statutorily mandated duties of ICCVAM and activities of NICEATM. The meeting is open to the public. Registration is requested for both public attendance and oral comment and required to access the webcast. Information about the meeting and registration is available at *http://ntp.niehs.nih.gov/go/32822.*

DATES: *Meeting:* September 16, 2014, beginning at 8:30 a.m. Eastern Daylight Time and continuing until adjournment at approximately 5:00 p.m.

Written Public Comments Submissions: Deadline is September 2, 2014. Registration for Meeting and Oral Comments: Requested by September 9, 2014. Registration to View Webcast: Deadline is September 16, 2014. Registration to view the meeting via the webcast is required.

ADDRESSES: *Meeting Location:* Rodbell Auditorium, Rall Building, NIEHS, 111