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

T.W. Alexander Drive, Research Triangle Park, NC 27709.

Meeting Web page: The preliminary agenda, registration, and other meeting materials are at *http://ntp.niehs.nih.gov/go/32822.*

Webcast: The meeting will be webcast; the URL will be provided to those who register for viewing.

FOR FURTHER INFORMATION CONTACT: Dr. Lori White, Designated Federal Officer for SACATM, Office of Liaison, Policy and Review, Division of NTP, NIEHS, P.O. Box 12233, K2–03, Research Triangle Park, NC 27709. Phone: 919–541–9834, fax: (301) 480–3272, email: *whiteld@niehs.nih.gov.* Hand Deliver/Courier address: 530 Davis Drive, Room K2136, Morrisville, NC 27560.

SUPPLEMENTARY INFORMATION:

Preliminary Agenda and Other Meeting Information: A preliminary agenda, roster of SACATM members, background materials, public comments, and any additional information, when available, will be posted on the SACATM meeting Web site (*http://ntp.niehs.nih.gov/go/32822*) or is available upon request from the Designated Federal Officer. Following the meeting, summary minutes will be prepared and available on the SACATM Web site or upon request.

Meeting and Registration: This meeting is open to the public with time scheduled for oral public comments. The public may attend the meeting at NIEHS, where attendance is limited only by the space available, or view the webcast. Registration is required to view the webcast; the URL for the webcast will be provided in the email confirming registration. Individuals who plan to attend and/or provide comments are encouraged to register at *http://ntp.niehs.nih.gov/go/32822* by September 9, 2014, to facilitate planning for the meeting. Individuals interested in the meeting are encouraged to access this Web site to stay abreast of the most current information regarding the meeting. Visitor and security information for those attending in person is available at niehs.nih.gov/about/visiting/index.cfm. Individuals with disabilities who need accommodation to participate in this event should contact Ms. Robbin Guy at phone: (919) 541–4363 or email: *guyr2@niehs.nih.gov.* TTY users should contact the Federal TTY Relay Service at 800–877–8339. Requests should be made at least five business days in advance of the event.

Request for Comments: Both written and oral public input on the agenda topics is invited. Written comments received in response to this notice will be posted on the meeting Web site and persons submitting them will be identified by their name and affiliation and/or sponsoring organization, if applicable. Persons submitting written comments should include their name, affiliation (if applicable), and sponsoring organization (if any) with the document. Time is allotted during the meeting for presentation of oral comments and each organization (sponsoring organization or affiliation) is allowed one time slot per topic. At least 7 minutes will be allotted for each speaker, and if time permits, may be extended up to 10 minutes at the discretion of the chair. Registration for oral comments will also be available on-site, although time allowed for presentation by on-site registrants may be less than for registered speakers and will be determined by the number of persons who register at the meeting. In addition to in-person oral comments at the meeting, public comments can be presented by teleconference line. There will be 50 lines for this call; availability will be on a first-come, first-served basis. The lines will be open from 8:30 a.m. until approximately 5:00 p.m., although public comments will be received only during the formal public comment periods, which will be indicated on the preliminary agenda. The access number for the teleconference line will be provided to registrants by email prior to the meeting.

Persons wishing to present oral comments are encouraged to register using the SACATM meeting registration form (*http://ntp.niehs.nih.gov/go/32822*), indicate the topic(s) on which they plan to comment, and, if possible, send a copy of their statement to *whiteld@niehs.nih.gov* by September 9, to enable review by SACATM, NICEATM, ICCVAM, and NIEHS/NTP staff prior to the meeting. Written statements can supplement and may expand the oral presentation. If registering on-site and reading from written text, please bring 30 copies of the statement for distribution and to supplement the record.

Background Information on ICCVAM, NICEATM, and SACATM: ICCVAM is an interagency committee composed of representatives from 15 Federal regulatory and research agencies that require, use, generate, or disseminate toxicological and safety testing information. ICCVAM conducts technical evaluations of new, revised, and alternative safety testing methods with regulatory applicability and promotes the scientific validation and regulatory acceptance of toxicological and safety-testing methods that more accurately assess the safety and hazards of chemicals and products and that reduce, refine (decrease or eliminate pain and distress), or replace animal use. The ICCVAM Authorization Act of 2000 (42 U.S.C. 285*l*–3) established ICCVAM as a permanent interagency committee of the NIEHS under NICEATM.

NICEATM administers ICCVAM, provides scientific and operational support for ICCVAM-related activities, and conducts independent validation studies to assess the usefulness and limitations of new, revised, and alternative test methods and strategies. NICEATM and ICCVAM work collaboratively to evaluate new and improved test methods and strategies applicable to the needs of U.S. Federal agencies. NICEATM and ICCVAM welcome the public nomination of new, revised, and alternative test methods and strategies for validation studies and technical evaluations. Additional information about ICCVAM and NICEATM can be found at *http://ntp.niehs.nih.gov/go/iccvam* and *http://ntp.niehs.nih.gov/go/niceatm.*

SACATM was established in response to the ICCVAM Authorization Act [Section 285*l*–3(d)] and is composed of scientists from the public and private sectors. SACATM advises ICCVAM, NICEATM, and the Director of the NIEHS and NTP regarding statutorily mandated duties of ICCVAM and activities of NICEATM. SACATM provides advice on priorities and activities related to the development, validation, scientific review, regulatory acceptance, implementation, and national and international harmonization of new, revised, and alternative toxicological test methods. Additional information about SACATM, including the charter, roster, and records of past meetings, can be found at *http://ntp.niehs.nih.gov/go/167.*

Dated: July 7, 2014.

John R. Bucher,

Associate Director, National Toxicology Program.

[FR Doc. 2014–16452 Filed 7–11–14; 8:45 am]

BILLING CODE 4140–01–P

DEPARTMENT OF HEALTH AND HUMAN SERVICES

Substance Abuse and Mental Health Services Administration

Agency Information Collection Activities: Proposed Collection; Comment Request

In compliance with Section 3506(c)(2)(A) of the Paperwork Reduction Act of 1995 concerning