TABLE 3—ESTIMATED ANNUAL THIRD-PARTY DISCLOSURE BURDEN [1]—Continued

21 CFR section; activity	Number of respondents	Number of disclosures per respondent	Total annual disclosures	Average burden per disclosure	Total hours
Total					1,030,270

[1] There are no capital costs or operating and maintenance costs associated with this collection of information.

These estimates reflect our continued experience with the information collection. We have made nominal adjustments to reflect the addition of burden associated with gluten and certain bottled or otherwise packaged beer; petition submissions received since our last evaluation of the information collection; and informal communications with industry regarding food product labeling.

Dated: April 7, 2023.

Lauren K. Roth,

Associate Commissioner for Policy.

[FR Doc. 2023–07683 Filed 4–11–23; 8:45 am]

BILLING CODE 4164–01–P

DEPARTMENT OF HEALTH AND HUMAN SERVICES

National Institutes of Health

National Center for Advancing Translational Sciences; Notice of Closed Meeting

Pursuant to section 10(d) of the Federal Advisory Committee Act, as amended, 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 Center for Advancing Translational Sciences Special Emphasis Panel; NCATS CTSA Training Grants Review Meeting.

Date: May 17, 2023.

Time: 9:30 a.m. to 6:30 p.m.

Agenda: To review and evaluate grant applications.

Place: Office of Grants Management and Scientific Review National Center for Advancing Translational Sciences, National Institutes of Health, 6701 Democracy Boulevard, Suite 1001 Bethesda, MD 20892 (Virtual Meeting).

Contact Person: Nakia C. Brown, Ph.D., Scientific Review Officer, Office of Grants Management and Scientific Review, National Center for Advancing Translational Sciences, National Institutes of Health, 6701 Democracy Blvd., Suite 101, Bethesda, MD 20892, 301–827–4905, *brownnac@mail.nih.gov*.

(Catalogue of Federal Domestic Assistance Program Nos. 93.859, Pharmacology, Physiology, and Biological Chemistry Research; 93.350, B—Cooperative Agreements; 93.859, Biomedical Research and Research Training, National Institutes of Health, HHS)

Dated: April 7, 2023.

Melanie J. Pantoja,

Program Analyst, Office of Federal Advisory Committee Policy.

[FR Doc. 2023–07656 Filed 4–11–23; 8:45 am]

BILLING CODE 4140–01–P

DEPARTMENT OF HEALTH AND HUMAN SERVICES

National Institutes of Health

Proposed Collection; 60-Day Comment Request; Specimen Resource Locator (National Cancer Institute)

AGENCY: National Institutes of Health, HHS.

ACTION: Notice.

SUMMARY: In compliance with the requirement of the Paperwork Reduction Act of 1995 to provide an opportunity for public comment on proposed data collection projects, the National Institutes of Health, National Cancer Institute (NCI) will publish periodic summaries of proposed projects to be submitted to the Office of Management and Budget (OMB) for review and approval.

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

FOR FURTHER INFORMATION CONTACT: To obtain a copy of the data collection plans and instruments, submit comments in writing, or request more information on the proposed project, contact: Joanne Demchok, Program Director, Cancer Diagnosis Program, Division of Cancer Treatment and Diagnosis, 9609 Medical Center Drive, Rockville, Md. 20892 or call non-toll-free number 240–276–5959 or Email your request, including your address to: *peterjo@mail.nih.gov.* Formal requests for additional plans and instruments must be requested in writing.

SUPPLEMENTARY INFORMATION: Section 3506(c)(2)(A) of the Paperwork Reduction Act of 1995 requires: written comments and/or suggestions from the public, and affected agencies are invited to address 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 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 minimizes 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.

Proposed Collection Title: Specimen Resource Locator (NCI), 0925–0703: Expiration Date 11/30/2023, EXTENSION, National Cancer Institute (NCI), National Institutes of Health (NIH).

Need and Use of Information Collection: The availability of specimens and associated data is critical to increase our knowledge of cancer biology and to translate important research discoveries into clinical applications. The development of molecular technologies in cancer patients with defined molecular abnormalities advances the identification and development of clinically useful biomarkers and diagnostic assays that guide treatment.

The discovery and validation of cancer prevention markers require researchers' access to quality clinical biospecimens. In response to this need, NCI's Cancer Diagnosis Program developed, and is expanding, a searchable database: Specimen Resource Locator (SRL) *https://specimens.cancer.gov/tissue/default.htm.* The SRL allows scientists in the research community and the NCI

to locate specimens needed for their research. The SRL lists all NCI-supported and non-NCI-supported biospecimens repositories and their links. It is not NCI's intent to collect the biospecimens; instead, the collections are descriptions of the available data that can act as a resource and be shared with interested researchers and scientists. This submission does not involve any analysis.

OMB approval is requested for 3 years. There are no costs to respondents other than their time. The total estimated annualized burden hours are 105.

ESTIMATED ANNUALIZED BURDEN HOURS

Type of respondent	Form name	Number of respondents	Number of responses per respondent	Average burden per response (in hours)	Total burden hour
Private Sector	Initial Request	70	1	30/60	35
State Government		70	1	30/60	35
Federal Government		60	1	30/60	30
Private Sector	Annual Update	20	1	5/60	2
State Government		20	1	5/60	2
Federal Government		10	1	5/60	1
Total			250		105

Dated: April 7, 2023.

Diane Kreinbrink,

Project Clearance Liaison, National Cancer Institute, National Institutes of Health.

[FR Doc. 2023–07684 Filed 4–11–23; 8:45 am]

BILLING CODE 4140–01–P

DEPARTMENT OF HEALTH AND HUMAN SERVICES

National Institutes of Health

Interagency Coordinating Committee on the Validation of Alternative Methods; Notice of Public Meeting; Request for Public Input

AGENCY: National Institutes of Health, HHS.

ACTION: Notice.

SUMMARY: The Interagency Coordinating Committee on the Validation of Alternative Methods (ICCVAM) will hold a public forum to share information and facilitate direct communication of ideas and suggestions from stakeholders. Interested persons may attend in person or view the meeting remotely by webcast. Time will be set aside for questions and public statements on the topics discussed. Registration is requested for attending in person and required for viewing the webcast and presenting oral statements. Information about the meeting and registration are available at *https://ntp.niehs.nih.gov/go/iccvamforum-2023.*

DATES: Meeting: May 18 and 19, 2023, 9 a.m. to approximately 3 p.m. EDT both days.

Registration for Onsite Meeting: Deadline is May 17, 2023. Registration for Webcast: Deadline is May 19, 2023. Registration for Oral Statements: Deadline is May 9, 2023.

Registration to attend in person is requested; registration to view the webcast and present oral public statements is required.

ADDRESSES: Meeting Location: William H. Natcher Conference Center, National Institutes of Health (NIH), Bethesda, MD 20892. Meeting web page: Registration and other meeting materials are at *https://ntp.niehs.nih.gov/go/iccvamforum-2023.* A preliminary agenda will be posted on this page by May 2, 2023.

FOR FURTHER INFORMATION CONTACT: Dr. Nicole Kleinstreuer, Director, National Toxicology Program Interagency Center for the Evaluation of Alternative Toxicological Methods (NICEATM), email: *nicole.kleinstreuer@nih.gov,* telephone: 984–287–3150.

SUPPLEMENTARY INFORMATION:

Background: ICCVAM, a congressionally mandated committee, coordinates the development and validation of alternative testing strategies that protect human health and the environment while replacing, reducing, or refining animal use.

ICCVAM's goals include promotion of national and international partnerships between governmental and nongovernmental groups, including academia, industry, advocacy groups, and other key stakeholders. To foster these partnerships ICCVAM initiated annual public forums in 2014 to share information and facilitate direct communication of ideas and suggestions from stakeholders (79 FR 25136).

This year's meeting will be held on May 18 and 19, 2023. NICEATM and ICCVAM members will give presentations on current activities related to the development and validation of alternative test methods and approaches.

There will be opportunities for participants to ask clarifying or follow-up questions of the ICCVAM members about their presentations. Instructions for submitting these questions will be provided to webcast viewers prior to the event. The agenda will also include time for public oral statements relevant to the ICCVAM mission and current activities from participants who have registered to do so in advance.

Preliminary Agenda and Other Meeting Information: A preliminary agenda will be posted by May 2 at *https://ntp.niehs.nih.gov/go/iccvamforum-2023.* Interested individuals are encouraged to visit this web page to stay abreast of the most current meeting information.

Meeting and Registration: This meeting is open to the public. The public may attend the meeting at NIH, where attendance is limited only by the space available, or view remotely by webcast. Those planning to attend the meeting in person are encouraged to register at *https://ntp.niehs.nih.gov/go/iccvamforum-2023* by May 17, 2023, to facilitate planning for appropriate meeting space. Registration for the webcast is required and is open through 3 p.m. EDT on May 19, 2023, at *https://ntp.niehs.nih.gov/go/iccvamforum-2023.* Registrants will receive instructions on how to access and participate in the webcast in the email confirming their registration.

NIH visitor and security information is available at *http://www.nih.gov/about/visitor/index.htm.* Individuals with disabilities who need accommodation to participate in this event should contact Milene Brownlow at phone: 984–287–3364 or email: