

information on the families being served under the Tribal MIECHV Program;

- Collect the number of newly enrolled and continuing families being served;
- Collect the number of home visits;
- Track and improve the quality of benchmark measures data submitted by the tribal grantees;
- Improve program monitoring and oversight;

• Improve rigorous data analyses that help to assess the effectiveness of the programs and enable ACF to better monitor projects;

• Ensure adequate and timely reporting of program data to relevant federal agencies and stakeholders, including Congress and members of the public; and

• Collect data on caseload capacity and the retention and attrition of

enrolled families and the retention and attrition of program staff on a quarterly basis.

Overall, this information collection will provide valuable information to HHS that will guide understanding of the Tribal MIECHV Program and the provision of technical assistance to Tribal MIECHV Program grantees.

*Respondents:* Tribal MIECHV Grantees.

#### ANNUAL BURDEN ESTIMATES

Instrument	Total number of respondents	Annual number of responses per respondent	Average burden hours per response	Annual burden hours
Tribal MIECHV Demographic and Service Utilization Data Form .....	55	1	317	17,435
Tribal MIECHV Demographic & Service Utilization Data Report (Families) ...	1,668	1	233	389
Tribal MIECHV Performance Measures Form .....	55	1	288	15,840
Tribal MIECHV Quarterly Performance Report .....	55	4	2.5	550

#### *Estimated Total Annual Burden*

Hours: 33,825.

*Authority:* Section 511 of title V of the Social Security Act.

**Mary B. Jones,**

*ACF/OPRE Certifying Officer.*

[FR Doc. 2023–13065 Filed 6–16–23; 8:45 am]

**BILLING CODE 4184–77–P**

#### DEPARTMENT OF HEALTH AND HUMAN SERVICES

[Document Identifier: OS–0990–0473–30D]

#### Agency Information Collection Request. 30-Day Public Comment Request

**AGENCY:** Office of the Secretary, HHS.

**ACTION:** Notice.

**SUMMARY:** In compliance with the requirement of the Paperwork Reduction Act of 1995, the Office of the Secretary (OS), Department of Health and Human Services, is publishing the following summary of a proposed collection for public comment.

**DATES:** Comments on the ICR must be received on or before July 20, 2023.

**ADDRESSES:** Written comments and recommendations for the proposed information collection should be sent within 30 days of publication of this notice to [www.reginfo.gov/public/do/PRAMain](http://www.reginfo.gov/public/do/PRAMain). Find this particular information collection by selecting “Currently under 30-day Review—Open for Public Comments” or by using the search function.

#### **FOR FURTHER INFORMATION CONTACT:**

Sherrette Funn, [Sherrette.Funn@hhs.gov](mailto:Sherrette.Funn@hhs.gov) or (202) 264–0041, or [PRA@HHS.GOV](mailto:PRA@HHS.GOV). When submitting comments or requesting information, please include the document identifier 0990–0473–30D and project title for reference.

**SUPPLEMENTARY INFORMATION:** Interested persons are invited to send comments regarding this burden estimate or any other aspect of this collection of information, including any of the following subjects: (1) The necessity and utility of the proposed information collection for the proper performance of the agency’s functions; (2) the accuracy of the estimated burden; (3) ways to enhance the quality, utility, and clarity of the information to be collected; and (4) the use of automated collection

techniques or other forms of information technology to minimize the information collection burden.

*Title of the Collection:* HHS Subpart C Certification Form.

*Type of Collection:* Revision.

*Abstract:* The Office for Human Research Protections (OHRP) is requesting a three-year extension of OMB No. 0990–0473, the HHS Subpart C Certification Form. The purpose of this form is to provide a simplified, standardized procedure for institutions to submit subpart C research certifications to OHRP in order to obtain authorization to include prisoners in HHS-conducted or supported human subjects research. The form also simplifies the internal process used by OHRP to review and record such certifications, resulting in faster processing while reducing unnecessary and burdensome staff time.

*Type of Respondent:* Institutions or Organizations operating Institutional Review Boards (IRBs) that have enrolled or are planning to enroll prisoners in human subjects research conducted or supported by HHS.

#### ESTIMATED ANNUALIZED BURDEN TABLE

Type of respondent	Number of respondents	Number responses per respondent	Average burden per response (in hours)	Total burden hours
Institutions or Organizations operating IRBs .....	25	2	1.0	50
Institutions or Organizations operating IRBs .....	5	3	1.0	15
Total .....				65

**Sherrette A. Funn,**

*Paperwork Reduction Act Reports Clearance Officer, Office of the Secretary.*

[FR Doc. 2023–13047 Filed 6–16–23; 8:45 am]

**BILLING CODE 4150–28–P**

## DEPARTMENT OF HEALTH AND HUMAN SERVICES

### National Institutes of Health

#### National Institute on Drug Abuse; Notice of Closed Meeting

Pursuant to section 1009 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 Institute on Drug Abuse Special Emphasis Panel; Cutting-Edge Basic Research Awards (CEBRA) Review Panel.

*Date:* July 18, 2023.

*Time:* 10 a.m. to 5:30 p.m.

*Agenda:* To review and evaluate grant applications.

*Place:* National Institute of Health, National Institute on Drug Abuse, 301 North Stonestreet Avenue, Bethesda, MD 20892 (Virtual Meeting).

*Contact Person:* Sheila Pirooznia, Ph.D., Scientific Review Officer, Division of Extramural Review, Scientific Review Branch, National Institute on Drug Abuse, NIH, 301 North Stonestreet Avenue, MSC 6021, Bethesda, MD 20892, (301) 496–9350, [sheila.pirooznia@nih.gov](mailto:sheila.pirooznia@nih.gov).

(Catalogue of Federal Domestic Assistance Program Nos. 93.277, Drug Abuse Scientist Development Award for Clinicians, Scientist Development Awards, and Research Scientist Awards; 93.278, Drug Abuse National Research Service Awards for Research Training; 93.279, Drug Abuse and Addiction Research Programs, National Institutes of Health, HHS)

Dated: June 14, 2023.

**Tyeshia M. Roberson-Curtis,**

*Program Analyst, Office of Federal Advisory Committee Policy.*

[FR Doc. 2023–13057 Filed 6–16–23; 8:45 am]

**BILLING CODE 4140–01–P**

## DEPARTMENT OF HEALTH AND HUMAN SERVICES

### National Institutes of Health

#### Submission for OMB Review; 30-Day Comment Request; Specimen Resource Locator (National Cancer Institute)

**AGENCY:** National Institutes of Health, HHS.

**ACTION:** Notice.

**SUMMARY:** In compliance with 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.

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

**ADDRESSES:** Written comments and recommendations for the proposed information collection should be sent within 30 days of publication of this notice to [www.reginfo.gov/public/do/PRAMain](http://www.reginfo.gov/public/do/PRAMain). Find this particular information collection by selecting “Currently under 30-day Review—Open for Public Comments” or by using the search function.

#### FOR FURTHER INFORMATION 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](mailto:peterjo@mail.nih.gov).

**SUPPLEMENTARY INFORMATION:** This proposed information collection was previously published in the **Federal Register** on April 12, 2023, page 22049 (Vol. 88, No. 70 FR 22049) 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.

In compliance with 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.

*Proposed Collection:* Specimen Resource Locator, OMB #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 increasing our knowledge of cancer biology and translating important research discoveries to clinical applications. The discovery and validation of cancer prevention markers require access, by researchers, to quality clinical biospecimens. In response to this need, the National Cancer Institute’s (NCI) Cancer Diagnosis Program has developed and is expanding a searchable database: Specimen Resource Locator (SRL). The SRL allows scientists in the research community and the NCI to locate specimens needed for their research. The SRL will list all NCI-supported repositories and their links. This administrative submission is an online form that will collect information to manage and improve a program and its resources for the use of all 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