

human-drugs/evaluating-immunosuppressive-effects-utero-exposure-drug-and-biologic-products-07112024. The transcripts and recordings will also be accessible at <https://www.regulations.gov> and may be viewed at the Dockets Management Staff (HFA-305), Food and Drug Administration, 5630 Fishers Lane, Rm. 1061, Rockville, MD 20852.

Dated: May 17, 2024.

Lauren K. Roth,

Associate Commissioner for Policy.

[FR Doc. 2024-11228 Filed 5-21-24; 8:45 am]

BILLING CODE 4164-01-P

DEPARTMENT OF HEALTH AND HUMAN SERVICES

Health Resources and Services Administration

Agency Information Collection Activities: Submission to OMB for Review and Approval; Public Comment Request: National Survey of Organ Donation Attitudes and Practices, OMB No. 0915-0290

AGENCY: Health Resources and Services Administration (HRSA), Department of Health and Human Services.

ACTION: Notice.

SUMMARY: In compliance with the Paperwork Reduction Act of 1995, HRSA submitted an Information Collection Request (ICR) to the Office of Management and Budget (OMB) for review and approval. Comments submitted during the first public review of this ICR will be provided to OMB. OMB will accept further comments from the public during the review and approval period. OMB may act on HRSA's ICR only after the 30-day comment period for this notice has closed.

DATES: Comments on this ICR should be received no later than June 21, 2024.

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. Find this particular information collection by selecting "Currently under Review—Open for Public Comments" or by using the search function.

FOR FURTHER INFORMATION CONTACT: To request a copy of the clearance requests submitted to OMB for review, email

Joella Roland, the HRSA Information Collection Clearance Officer, at paperwork@hrsa.gov or call (301) 443-3983.

SUPPLEMENTARY INFORMATION:

Information Collection Request Title: National Survey of Organ Donation Attitudes and Practices, OMB No. 0915-0290—Revision.

Abstract: The overall purpose of this study is to conduct an independent multi-mode (web and telephone) survey of public opinion regarding various issues related to organ donation. The survey will measure public opinion on issues such as willingness to become an organ donor, financial incentives for donation, living donation, impediments to donation, and level of public knowledge about donation. Previous National Survey of Organ Donation Attitudes and Practices were conducted during 1993, 2005, 2012, and 2019. Similar to the 2019 survey, the goal is to complete 10,000 interviews with adults (18 years of age or older) nationwide. Specifically, this will include 1,000 equal-probability of selection method computer-assisted telephone interviewing (CATI) interviews, 1,000 ethnic oversamples CATI interviews, and a supplemental web panel of 8,000 respondents. The final sample will include 1,000 interviews each with Black or African Americans, Asian Americans, Hispanic/Latino Americans, and American Indian/Alaskan Natives, and a statistically sufficient sample for meaningful comparisons across demographic levels of age group, education, and income groups. A total sample of 10,000 is necessary to achieve sufficiently large subgroups for statistical analysis across demographic groups.

A 60-day notice published in the **Federal Register** on January 18, 2024, vol. 89, No. 12; pp. 3409-3410. There were no public comments.

Need and Proposed Use of the Information: The Division of Transplantation, within the Health Systems Bureau of HRSA at the Department of Health and Human Services, is the primary federal entity responsible for oversight of the solid organ and blood stem cell transplant systems in the United States and for initiatives to increase organ donor registration and donation. Sponsorship of a national survey on the American public's donation attitudes and practices is one of the services that

Division of Transplantation provides for the larger donation community, consistent with its legal authority to establish a public education and awareness program (Section 377A of the Public Health Service Act, 42 U.S.C. 274f-1).

Patients in need of organ transplantation in the United States face a longstanding critical shortage of organs. Approximately 103,000 Americans were on the waiting list for transplantation by the end of 2022, but only 42,000 transplants were performed, which only meets two-fifths of the national need. While this represents an increase from the number of transplants performed in 2021, the organ shortage remains in the United States. Understanding public attitudes about organ donation and how the attitudes change over time is critical to addressing organ shortage through public awareness and education efforts.

The information from this survey will facilitate appropriate tailoring and targeting of donation outreach messages and strategies and provide an overall assessment of the impact of previous outreach messages and strategies. The data will also inform the development of policy related to organ donation and transplantation.

Likely Respondents: A nationally representative sample of adults over the age of 18 with a higher number of responses from populations of interest such as racial-ethnic minorities, including Black or African American, Asian American, American Indian/Alaskan Native, and Hispanic/Latino American respondents, as well as respondents of all age groups and education levels.

Burden Statement: Burden in this context means the time expended by persons to generate, maintain, retain, disclose, or provide the information requested. This includes the time needed to review instructions; to develop, acquire, install, and utilize technology and systems for the purpose of collecting, validating, and verifying information, processing and maintaining information, and disclosing and providing information; to train personnel and to be able to respond to a collection of information; to search data sources; to complete and review the collection of information; and to transmit or otherwise disclose the information. The total annual burden hours estimated for this ICR are summarized in the table below.

TOTAL ESTIMATED ANNUALIZED BURDEN HOURS

Form name	Number of respondents	Number of responses per respondent	Total responses	Average burden per response (in hours)	Total burden hours
The National Survey of Organ Donation Attitudes and Practices—Telephone (English and Spanish versions)	2,000	1	2,000	0.37	740
The National Survey of Organ Donation Attitudes and Practices—Web (English and Spanish versions)	8,000	1	8,000	0.27	2,160
Total	10,000	2,900

Amy P. McNulty,
Deputy Director, Executive Secretariat.
[FR Doc. 2024–11246 Filed 5–21–24; 8:45 am]
BILLING CODE 4165–15–P

DEPARTMENT OF HEALTH AND HUMAN SERVICES

National Institutes of Health

Submission for OMB Review; 30-Day Comment Request; Application and Impact of Clinical Research Training on Healthcare Professionals in Academia and Clinical Research (Office of the Director)

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. 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: To request more information on the proposed project or to obtain a copy of the data collection plans and instruments, contact: Dr. Anne Zajicek, Program Director, Office of Clinical Research Education and Collaboration Outreach, OD, NIH, Building 1, Room 201, 1 Center Drive, Bethesda, MD 20892, or call non-toll-free number (301) 480–9913 or Email your request, including your address to: zajiceka@mail.nih.gov.

SUPPLEMENTARY INFORMATION: This proposed information collection was previously published in the **Federal Register** on Monday, July 31, 2023, Volume 88, pages 49472–49473 (64 FR 16184) 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 Office of Clinical Research Education and Collaboration Outreach, Office of the Director, 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: Application and Impact of Clinical Research Training on Healthcare Professionals in Academia and Clinical Research, 0925–0764-expiration date, 02/28/2026, Office of Clinical Research Education and Collaboration Outreach (OCRECO), National Institutes of Health (NIH), Office of the Director (OD).

Need and Use of Information Collection: The purpose of this survey is to assess the long-term impact and outcomes of clinical research training programs provided by the Office of Clinical Research Education and Collaboration Outreach located in the NIH Office of the Director (OD) over a ten-year follow-up period. The information received from respondents will provide insight on the following: impact of the courses on (a) promotion of professional competence, (b) research productivity and independence, and (c) future career development within clinical, translational and academic research settings. These surveys will provide preliminary data and guidance in (1) developing recommendations for collecting outcomes to assess the effectiveness of the training courses, and (2) tracking the impact of the curriculum on participants’ ability to perform successfully in academic, non-academic, research, and non-research settings.

OMB approval is requested for 3 years. There are no costs to respondents other than their time. The total estimated annualized burden hours are 1,773.

ESTIMATED ANNUALIZED BURDEN HOURS

Form name	Type of respondents	Estimated number of respondents	Number of responses per respondent	Average burden per response (in hours)	Total annual burden hours
OCRECO Learning Portal Registration (Attachment 1)	Healthcare Professionals	2,000	1	5/60	167
	Students	2,000	1	5/60	167
	General Public	1,000	1	5/60	83
IPPCR Lecture Evaluation (Attachment 2)	Healthcare Professionals	1,000	1	5/60	83
	Students	2,000	1	5/60	167
	General Public	1,000	1	5/60	83