

Dated: July 16, 2026.

David M. Swegle,

*Director Office for Family Assistance,
Administration for Children and Families.*

[FR Doc. 2026–14587 Filed 7–17–26; 8:45 am]

BILLING CODE 4184–42–P

DEPARTMENT OF HEALTH AND HUMAN SERVICES

[Document Identifier: OS–0990–0279]

Agency Information Collection Request; 30-Day Public Comment Request

AGENCY: Office for Human Research Protections (OHRP), Office of the Assistant Secretary for Health (OASH), 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, submitted an Information Collection Request (ICR) to the Office of Management and Budget (OMB) for review and approval. OMB will accept further comments from the public during the review and approval period.

DATES: Comments on the ICR must be received on or before August 19, 2026.

ADDRESSES: Written comments and recommendations for the proposed information collection should be sent within 30 days of publication of this notice via www.reginfo.gov/public/do/PRAMain. Find this information collection by selecting “Currently under Review” and “Select Agency: Department of Health and Human Services”.

FOR FURTHER INFORMATION CONTACT: Natalie Klein, Natalie.Klein@hhs.gov or

(240) 453–6900. If requesting information, please include the document identifier 0990–0279–30D and project title (Department of Health and Human Services (HHS) Registration of an Institutional Review Board Form) 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: Department of Health and Human Services (HHS) Registration of an Institutional Review Board Form.

Type of Collection: Reinstatement with changes OMB No. 0990–0279.

Abstract: The Office for Human Research Protections (OHRP) and the Food and Drug Administration (FDA) are requesting reinstatement with changes for the Office of Management and Budget (OMB) No. 0990–0279, Department of Health and Human Services (HHS) Institutional Review Board (IRB) Registration Form. The previous information collection was approved by OMB on June 27, 2022, and expired on June 30, 2025. The request for reinstatement with changes involves implementing a burden reducing change. Specifically, OHRP is seeking to remove the IRB membership roster information collection from the IRB registration form. This change will align the IRB registration form with the 2018

Requirements at 45 CFR 46.103. The change, when implemented, is anticipated to result in a shorter, simplified IRB registration process for respondents. Updates to the software applications OHRP uses to manage IRB registration will be deployed to enable such changes.

The purpose of the form is to provide a simplified procedure for: (1) IRBs to satisfy the requirements for IRB registration at 45 CFR part 46, subpart E; and (2) IRBs in the United States (US) to satisfy the FDA requirements for IRB registration at 21 CFR 56.106. Institutions engaged in nonexempt human subjects research conducted or supported by HHS, or another Common Rule department or agency, are required by the terms of the Federalwide Assurance (FWA) to rely upon only IRBs registered with OHRP for review of research to which the FWA applies. In this way, OHRP’s FWA process, established pursuant to the requirements for assurances at 45 CFR 46.103, is linked to the regulatory requirements for IRB registration.

The respondents for this information collection are institutions or organizations operating IRBs that review human subjects research conducted or supported by HHS; or, in the case of FDA’s requirements, each IRB in the United States that reviews clinical investigations regulated by FDA under sections 505(i) or 520(g) of the Federal Food, Drug and Cosmetic Act; and each IRB in the United States that reviews clinical investigations that are intended to support applications for research or marketing permits for FDA-regulated products. Many of the IRBs also review research conducted or supported by other Common Rule departments and agencies.

ANNUALIZED BURDEN HOUR TABLE

Form	Respondents	Number of respondents	Number of responses per respondent	Average burden per response (in hours)	Total burden hours
IRB Registration Form 0990–0279 ...	Institutions or Organizations that operate IRBs.	5,350	1	0.5	2,675
Total		5,350	1	0.5	2,675

The estimate of the number of respondents is based upon the current number of IRBs registered with HHS, approximately 4,988 (as of July 13, 2026), and projecting that the number of respondents may increase to 5,350. Of the approximately 4,988 registered IRBs,

100% submitted their registration information electronically.

Of the 5,350 projected respondent IRBs, approximately 350 institutions or organizations are expected to submit IRB registration information for the first time, and nearly 5,000 institutions or

organizations are expected to update or renew their existing IRB registration once each year. The burden is estimated to average 0.5 hours for both an initial IRB registration and for updating or renewing the registration of a previously registered IRB. If on average 5,000

previous respondents submit information once each year, and on average 350 submit new registration information one time a year, the total annual burden hours are projected to be 2,675 hours. Consistent with 5 CFR 1320.5(a)(1)(iv)(5), therefore, we believe this estimate represents the total annual reporting and recordkeeping burden that will result from the collection of information.

Catherine Howard,

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

[FR Doc. 2026–14537 Filed 7–17–26; 8:45 am]

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DEPARTMENT OF HEALTH AND HUMAN SERVICES

National Institutes of Health

Office of the Secretary; Amended Notice of Meeting

Notice is hereby given of a change in the meeting of the Interagency Autism Coordinating Committee, July 31, 2026, 9:00 a.m. to 5:00 p.m. ET. National Institutes of Mental Health (NIMH), Neuroscience Center (NSC), First Floor Conference Rooms, 6001 Executive Boulevard, Rockville, MD 20852, was published in the **Federal Register** on July 07, 2026, FR Doc. 2026–13714, 91 FR 41645.

This Notice is being amended to extend the public comment period from July 17, 2026, through July 24, 2026. The meeting is open to the public.

Dated: July 15, 2026.

Rosalind M Niamke,

Program Analyst, Office of Federal Advisory Committee Policy.

[FR Doc. 2026–14599 Filed 7–17–26; 8:45 am]

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DEPARTMENT OF HOMELAND SECURITY

U.S. Immigration and Customs Enforcement

[OMB Control Number 1653–0055]

Flight Manifest/Billing Agreement

AGENCY: U.S. Immigration and Customs Enforcement, Department of Homeland Security.

ACTION: 30-Day notice.

SUMMARY: In accordance with the Paperwork Reduction Act (PRA) of 1995 the Department of Homeland Security (DHS), U.S. Immigration and Customs Enforcement (ICE) will submit the

following Information Collection Request (ICR) to the Office of Management and Budget (OMB) for review and clearance. This information collection was previously published in the **Federal Register** on December 17, 2025, allowing for a 60-day comment period. ICE received several comments which have been addressed in the supporting statement available on www.reginfo.gov/public/do/PRAMain. The purpose of this notice is to allow an additional 30 days for public comments.

DATES: Comments are encouraged and will be accepted until August 19, 2026.

ADDRESSES: Written comments and recommendations for this 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 under e-Docket ID number ICEB–2021–0005–0003.

FOR FURTHER INFORMATION CONTACT: For specific question related to collection activities, please contact U.S. Immigration and Customs Enforcement at ICEOCFOPolicy@ice.dhs.gov.

SUPPLEMENTARY INFORMATION:

Comments

Written comments and suggestions from the public and affected agencies concerning the proposed collection of information should address one or more of the following four points:

(1) Evaluate whether the proposed collection of information is necessary for the proper performance of the functions of the agency, including whether the information will have practical utility;

(2) Evaluate the accuracy of the agencies estimate of the burden of the proposed collection of information, including the validity of the methodology and assumptions used;

(3) Enhance the quality, utility, and clarity of the information to be collected; and

(4) Minimize the burden of the collection of information on those who are to respond, including through the use of appropriate automated, electronic, mechanical, or other technological collection techniques or other forms of information technology, e.g., permitting electronic submission of responses.

Overview of This Information Collection

(1) Type of Information Collection: Extension of a Currently Approved Collection.

(2) Title of the Form/Collection: Flight Manifest/Billing Agreement.

(3) Agency form number, if any, and the applicable component of the Department of Homeland Security sponsoring the collection: U.S. Immigration and Customs Enforcement.

(4) Affected public who will be asked or required to respond, as well as a brief abstract: Primary: Individuals or households. The Flight Manifest/Billing Agreement collects information for the purpose of confirming Space Available passengers on any ICE-chartered flight and to facilitate the effective billing of those passengers for the full coach fare of their seats on the flight.

(5) An estimate of the total number of responses and the amount of time estimated for an average respondent to respond: The estimated total number of respondents for the information collection is 250 and the estimated hour burden per response is .25 hours.

(6) An estimate of the total public burden (in hours) associated with the collection: 63 annual burden hours.

(7) An estimate of the total public burden (in cost) associated with the collection: The estimated total annual cost burden associated with this collection of information is \$2,391.

Dated: May 27, 2026.

Ashley Pearce,

Deputy Chief Data Officer.

[FR Doc. 2026–14560 Filed 7–17–26; 8:45 am]

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DEPARTMENT OF THE INTERIOR

Geological Survey

[Docket No. USGS–2026–0199; OMB Control Number 1028–0137; GX26EN05ES90000]

Agency Information Collection Activities; Submission to the Office of Management and Budget for Review and Approval; Evaluation of the Arctic Rivers Project

AGENCY: U.S. Geological Survey, Interior.

ACTION: Notice of information collection; request for comment.

SUMMARY: In accordance with the Paperwork Reduction Act of 1995 (PRA), the U.S. Geological Survey (USGS) is proposing an extension of an information collection.

DATES: Interested persons are invited to submit comments on or before August 19, 2026.

ADDRESSES: You may submit comments by one of the following methods:

■ *Internet:* <https://www.regulations.gov>. Search for and