

collection; *Title of Information Collection Request:* Initial Plan Data Collection to Support QHP Certification and other Financial Management and Exchange Operations; *Use:* As directed by the rule Establishment of Exchanges and Qualified Health Plans; Exchange Standards for Employers (77 FR 18310) (Exchange rule), each Exchange is responsible for the certification and offering of Qualified Health Plans (QHPs). To offer insurance through an certification standards, such as network adequacy, inclusion of Essential Community Providers (ECPs), and non-discrimination. The Exchange is responsible for ensuring that QHPs meet these minimum certification standards as described in the Exchange rule under 45 CFR 155 and 156, based on the Patient Protection and Affordable Care Act (PPACA), as well as other standards determined by the Exchange. Issuers can offer individual and small group market plans outside of the Exchanges that are not QHPs.

Issuers can offer individual and small group market plans outside of the Exchanges that are not QHPs. Such plans are referred to in this document as “non-Exchange.” For the risk adjustment program, administrative information is used to identify all non-grandfathered small group and individual market non-Exchange plan offerings eligible for the program. Risk adjustment also requires select data such as rating area, rating factors, and actuarial value (AV) level, to perform calculation of payments and charges.

This information collection request serves as a formal request for the revision of the data collection clearance. We intend to use the instruments in this information collection for the 2025 certification process and beyond, and believe that providing these instruments now will give issuers and other stakeholders more opportunity to familiarize themselves with the instruments before releasing the 2025 application. While we intend to use these instruments in 2025, we may propose further revisions to this data collection in the future as necessary which will include seeking comments through the full 60-day and 30-day public comment periods. *Form Number:* CMS-10433 (OMB control number: 0938-1187); *Frequency:* Annually; *Affected Public:* Private Sector—Business or other for-profits; State, Local, or Tribal Governments; *Number of Respondents:* 1,073; *Number of Responses:* 1,073; *Total Annual Hours:* 61,154. (For questions regarding this

collection, contact Alexandra Gribbin at (667) 290-9977.)

William N. Parham, III,

Director, Division of Information Collections and Regulatory Impacts, Office of Strategic Operations and Regulatory Affairs.

[FR Doc. 2026-05915 Filed 3-25-26; 8:45 am]

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DEPARTMENT OF HEALTH AND HUMAN SERVICES

Administration for Children and Families

[Office of Management and Budget (OMB) #: 0970-0248]

Submission for OMB Review; Annual Report on Temporary Assistance for Needy Families Programs and State Maintenance-of-Effort Programs—ACF-204 (Annual TANF and MOE Report)

AGENCY: Office of Family Assistance, Administration for Children and Families, U.S. Department of Health and Human Services.

ACTION: Request for public comments.

SUMMARY: The Administration for Children and Families (ACF) is requesting a 3-year extension of the ACF-204: Annual Temporary Assistance for Needy Families (TANF) and State Maintenance-of-Effort (MOE) Report (Office of Management and Budget #0970-0248, expiration March 31, 2026). There are no substantive changes requested to the form, but instructions have been updated, and burden estimates adjusted.

DATES: *Comments due* April 27, 2026.

ADDRESSES: The public may view and comment on this information collection request at: https://www.reginfo.gov/public/do/PRAViewICR?ref_nbr=202603-0970-007. You can also obtain copies of the proposed collection of information by emailing infocollection@acf.hhs.gov. Identify all emailed requests by the title of the information collection.

SUPPLEMENTARY INFORMATION:

Description: The Annual TANF and MOE Report is used to collect descriptive program characteristics information on the programs operated by states and territories in association with their TANF programs. All state and territory expenditures claimed toward states and territories MOE requirements must be appropriate, *i.e.*, meet all applicable MOE requirements. The Annual TANF and MOE Report provides the ability to learn about and to monitor the nature of state and

territory expenditures used to meet states and territories MOE requirements, and it is an important source of information about the different ways that states and territories are using their resources to help families attain and maintain self-sufficiency. In addition, the report is used to obtain state and territory program characteristics for ACF's annual report to Congress, and the report serves as a useful resource to use in Congressional hearings about how TANF programs are evolving, in assessing state and territory MOE expenditures, and in assessing the need for legislative changes. This information collection will support administration and agency priorities to enhance oversight mechanisms and reinforce accountability across programs. Further, this information collection addresses a recommendation issued by the U.S. Government Accountability Office (GAO) in its final report, GAO-25-107235, that the Secretary of the Department of Health and Human Services should ensure ACF conducts a review of all TANF reporting requirements and forms and make appropriate changes within its statutory authority to enhance reporting for oversight of TANF funds. The Annual TANF and MOE Report requires states to relate each TANF and MOE benefit or service program to the third and fourth statutory purposes of TANF, which is responsive to GAO's recommendation and aligns with the agency's values of program integrity and fiscal stewardship. This request includes clarifications within instructions, a reduction in burden hours, and a change in collection title to better reflect the content of the collection.

Respondents: The 50 states of the U.S., the District of Columbia, Guam, Puerto Rico, and the Virgin Islands.

Annual Burden Estimates

ACF proposes a 32 percent reduction in estimated average time per response to reflect the intent of existing regulations. 45 CFR 265.9(d) indicates that states that have already incorporated the information required in their TANF State Plan may meet the annual reporting requirements by reference in lieu of completing a re-submission of a particular data element. If the information in the annual report has not changed since the previous annual report, the state may reference this information in lieu of re-submission of a particular data element. Instructions have been updated to make clear both opportunities to reference in lieu of re-submission.

Instrument	Total number of respondents	Annual number of responses per respondent	Average burden hours per response	Annual burden hours
ACF-204; Annual TANF and MOE Report	54	1	80	4,320

Authority: 42 U.S.C. 602 and 611; the Personal Responsibility and Work Opportunity Reconciliation Act of 1996, Public Law 104-193, 110 Stat. 2105.

Mary C. Jones,
ACF/OPRE Certifying Officer.

[FR Doc. 2026-05855 Filed 3-25-26; 8:45 am]

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DEPARTMENT OF HEALTH AND HUMAN SERVICES

Food and Drug Administration

[FDA-2025-N-3346]

Elite Laboratories, Inc., et al.; Withdrawal of Approval of 72 Abbreviated New Drug Applications; Correction

AGENCY: Food and Drug Administration, HHS.

ACTION: Notice; correction.

SUMMARY: The Food and Drug Administration (FDA) is correcting a notice that appeared in the **Federal Register** on September 24, 2025 (90 FR 183), appearing on page 45942 in FR Doc. 2025-18453. The document announced the withdrawal of approval of 72 abbreviated new drug applications (ANDAs) from multiple applicants, withdrawn as of October 23, 2025. The document indicated that FDA was withdrawing approval of the ANDA 070631 for valproic acid, capsule, 250 milligrams, held by Upsher-Smith Laboratories, LLC, 6701 Evenstad Dr., Maple Grove, MN 55369. Before FDA withdrew the approval of this ANDA, Upsher-Smith Laboratories, LLC, 6701 Evenstad Dr., Maple Grove, MN 55369, informed FDA that they did not want the approval of the ANDA withdrawn. Because Upsher-Smith Laboratories, LLC, timely requested that approval of the ANDA not be withdrawn, the approval is still in effect. This notice corrects this error.

FOR FURTHER INFORMATION CONTACT: Martha Nguyen, Center for Drug Evaluation and Research, Food and Drug Administration, 10903 New Hampshire Ave., Bldg. 75, Rm. 1676, Silver Spring, MD 20993-0002, 301-796-3471, Martha.Nguyen@fda.hhs.gov.

SUPPLEMENTARY INFORMATION: In the **Federal Register** of Wednesday

September 24, 2025 (90 FR 183), appearing on page 45942 in FR Doc. 2025-18453, the following correction is made:

On page 45943, in the table, the entry for ANDA 070631 is removed.

Grace R. Graham,
Deputy Commissioner for Policy, Legislation, and International Affairs.

[FR Doc. 2026-05913 Filed 3-25-26; 8:45 am]

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DEPARTMENT OF HEALTH AND HUMAN SERVICES

National Institutes of Health

National Eye Institute; Notice of Closed Meeting

Pursuant to section 1009 of the Federal Advisory Committee Act, as amended, notice is hereby given of a meeting of the Board of Scientific Counselors, National Eye Institute.

The meeting will be closed to the public as indicated below in accordance with the provisions set forth in sections 552b(c)(4) and 552b(c)(6), Title 5 U.S.C., as amended for the review, discussion, and evaluation of individual intramural programs and projects conducted by the National Eye Institute, including consideration of personnel qualifications and performance, and the competence of individual investigators, the disclosure of which would constitute a clearly unwarranted invasion of personal privacy.

Name of Committee: Board of Scientific Counselors, National Eye Institute.

Date: April 6, 2026.

Time: 2:00 p.m. to 3:00 p.m.

Agenda: To review and evaluate personnel qualifications and performance, and competence of individual investigators.

Address: National Eye Institute, 31 Center Drive, Bethesda, MD 20892.

Meeting Format: Virtual Meeting.

Contact Person: David M. Schneeweis, Ph.D., Acting Scientific Director, National Eye Institute, National Institutes of Health, Building 31, Room 6A22, Bethesda, MD 20892, 301-451-6763, David.schneeweis@nih.gov.

Information is also available on the Institute's/Center's home page: <https://www.nei.nih.gov/about/advisory-committees>, where an agenda and any additional information for the meeting will be posted when available.

(Catalogue of Federal Domestic Assistance Program No. 93.867, Vision Research, National Institutes of Health, HHS)

Dated: March 23, 2026.

Rosalind M. Niamke,
Program Analyst, Office of Federal Advisory Committee Policy.

[FR Doc. 2026-05852 Filed 3-25-26; 8:45 am]

BILLING CODE 4140-01-P

DEPARTMENT OF HOMELAND SECURITY

Finding of Mass Influx of Aliens

On January 23, 2025, the Acting Secretary of Homeland Security issued a Finding of Mass Influx of Aliens. This finding went into effect immediately (on January 23, 2025) and remained in effect for 60 days (until March 23, 2025). The Acting Secretary's finding published in the **Federal Register** on January 29, 2025. *See* 90 FR 8399. On March 21, 2025, I extended the January 23, 2025, Finding of Mass Influx for 180 days (until September 17, 2025). My decision extending the Finding of Mass Influx published in the **Federal Register** on March 25, 2025. *See* 90 FR 13622. On September 17, 2025, I extended the March 23, 2025, Finding of Mass Influx for 180 days (until March 21, 2026). My decision extending the Finding of Mass Influx published in the **Federal Register** on September 22, 2025. *See* 90 FR 45396. Upon review of the current situation at the border, I am extending that finding.

The Immigration and Nationality Act (INA), at 8 U.S.C. 1103(a), provides an expansive grant of authority, stating that in the event of a mass influx of aliens off the coast of the United States or a land border, the Secretary may authorize a State or local law enforcement officer, with the consent of the officer's superiors, to perform duties of immigration officers under the INA. In turn, section 65.83 of Title 28 of the Code of Federal Regulations allows the Secretary¹ to "request assistance from a

¹ Although the regulations reference the "Attorney General," Congress has, since the publication of these regulations, transferred the authority and responsibility for administering and enforcing the immigration laws to the Secretary of Homeland Security. *See* Homeland Security Act of 2002 471, 6 U.S.C. 291 (abolishing the former Immigration and Naturalization Service); id. S 441,

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