

Burden Estimate

The proposal would establish regulatory requirements for FDIC-supervised PPSIs as mandated by the GENIUS Act, as well as provide further clarity for FDIC-supervised custodians. As discussed in the proposal, the FDIC recognizes the significant uncertainty regarding estimates of the number of FDIC-supervised PPSIs and IDIs that would seek to issue payment stablecoins or engage in other permitted payment stablecoin activities. For the purposes of providing a conservative estimate, the

FDIC assumes that approximately 30 FDIC-supervised IDIs would perform these activities under the proposal. The FDIC assumes that: (1) 10 FDIC-supervised IDIs would seek approval for their subsidiary to become a PPSI each year over the first three years after the proposal is finalized and (2) there would be approximately 30 FDIC-supervised custodians, some of which may also be FDIC-supervised PPSIs themselves.

The total burden estimates for this collection of information were

published in the proposal. The proposal included burden estimates for the weekly form (information collection 7 for implementation burden and information collection 36 for ongoing burden in the proposal) and for the quarterly form (information collection 8 for implementation burden and 37 for ongoing burden in the proposal). This notice is re-publishing the estimates for the weekly and quarterly reporting burden below for informational purposes.

TABLE 1—SUMMARY OF ESTIMATED ANNUAL BURDEN FOR WEEKLY AND QUARTERLY REPORTING
[OMB No. 3064–NEW]

Information Collection (IC) (obligation to respond)	Type of burden (frequency of response)	Number of respondents	Number of responses per respondent	Average time per response (HH:MM)	Annual burden (hours)
Implementation Burden Reporting Requirements					
Confidential weekly report to the FDIC—Section 350.7(g) (Mandatory).	Reporting	10	1	16:00	160
Provide quarterly reports of financial condition—Section 350.7(h) (Mandatory).	Reporting	10	1	80:00	800
Ongoing Burden Reporting Requirements					
Provide confidential weekly report to the FDIC—Section 350.7(g) (Mandatory).	Reporting (Weekly)	20	52	2:00	2,080
Provide quarterly reports of financial condition to the FDIC—Section 350.7(h) (Mandatory).	Reporting (Quarterly)	20	4	10:00	800
Total for weekly and quarterly reporting burden (Hours).					3,840

The weekly and quarterly reporting information collections and their corresponding estimates were introduced in the *GENIUS Act Requirements and Standards for FDIC-Supervised Permitted Payment Stablecoin Issuers and Insured Depository Institutions* notice of proposed rule-making published on April 10, 2026 (91 FR 18534). The estimated annual burden for this collection is 22,110 hours. The estimated annual burden for the weekly and quarterly reporting is 3,840 hours.

In addition to the questions posed through the document, comments are also invited on: (a) whether the collections of information are necessary for the proper performance of the FDIC's functions, including whether the information has practical utility; (b) the accuracy of the estimates of the burden of the information collections, including the validity of the methodology and assumptions used; (c) ways to enhance the quality, utility, and clarity of the information to be collected; and (d) ways to minimize the burden of the collections of information on respondents, including through the use of automated collection techniques or other forms of information technology; and (e) estimates of capital or start-up costs and costs of operation, maintenance, and purchase of services to provide information. All comments will become a matter of public record.

Federal Deposit Insurance Corporation.

Dated at Washington, DC, on July 16, 2026.

Debra A. Decker,

Executive Secretary.

[FR Doc. 2026–14589 Filed 7–17–26; 8:45 am]

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FEDERAL MARITIME COMMISSION

[Docket No. 26–02]

Nancy Prior, Complainant v. AMOOV Group; FreightLead LLC; and Air 7 Seas Transport Logistics, Inc., Respondents; Notice of Filing of Amended Complaint

Notice is given that an amended complaint has been filed with the Federal Maritime Commission (the “Commission”) by Nancy Prior (the “Complainant”) against AMOOV Group; FreightLead LLC; and Air 7 Seas Transport Logistics, Inc. (the “Respondents”). Complainant states that the Commission has personal and subject-matter jurisdiction over this

complaint pursuant to 46 U.S.C. 41102(c), 41301, and 41305 and 46 CFR part 502.

Complainant is an individual residing in Fort Lauderdale, Florida.

Complainant identifies Respondent AMOOV Group as an international freight forwarder and shipping logistics provider with place of business in Lille, France.

Complainant identifies Respondent FreightLead LLC as a licensed non-vessel-operating common carrier with its corporate and operational headquarters in Parsippany, New Jersey.

Complainant identifies Respondent Air 7 Seas Transport Logistics, Inc. as a licensed non-vessel-operating common carrier with its corporate and operational headquarters in Milpitas, California.

Complainant alleges that Respondents violated 46 U.S.C. 41102(c). Complainant alleges these violations arose from Respondents withholding household goods, demanding payment

of unsupported charges as a condition of release despite no contractual obligation, withholding shipping documentation, and other acts or omissions of Respondents.

Pursuant to the order issued by the presiding judge on June 30, 2026, titled “Order Denying Respondent Freightlead LLC’s Motion to Dismiss Without Prejudice and Directing Complaint to File Amended Complaint,” answers to the amended complaint must be filed with the Commission no later than July 28, 2026.

The full text of the amended complaint can be found in the Commission’s electronic Reading Room at <https://www2.fmc.gov/readingroom/proceeding/26-02/>.

This proceeding has been assigned to the Office of Administrative Law Judges. The initial decision of the presiding judge shall be issued by January 14, 2027, and the final decision of the Commission shall be issued by July 28, 2027.

(Authority: 46 U.S.C. 41301; 46 CFR 502.61(c).)

Served: July 15, 2026.

David Eng,
Secretary.

[FR Doc. 2026–14541 Filed 7–17–26; 8:45 am]

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DEPARTMENT OF HEALTH AND HUMAN SERVICES

Centers for Medicare & Medicaid Services

[CMS–4219–N]

Medicare Program; Inflation Reduction Act of 2022 (IRA) Medicare Drug Price Negotiation Program Draft Guidance; Comment Request

AGENCY: Centers for Medicare & Medicaid Services (CMS), Department of Health and Human Services (HHS).

ACTION: Notice.

SUMMARY: The Centers for Medicare & Medicaid Services (CMS) is announcing an opportunity for the public to comment on CMS’ draft guidance for the Medicare Drug Price Negotiation Program regarding manufacturer effectuation of the maximum fair price in 2028 for the implementation of the Inflation Reduction Act of 2022 (IRA) (Pub. L. 117–169). CMS’ draft guidance and other IRA-related guidance can be viewed on the CMS website at <https://www.cms.gov/priorities/medicare-prescription-drug-affordability/medicare-prescription-drug-affordability>.

DATES: Comments must be received by September 18, 2026.

ADDRESSES: Written comments should be sent to IRAREbateandNegotiation@cms.hhs.gov with the subject line, “Medicare Drug Price Negotiation Program Draft Guidance”.

FOR FURTHER INFORMATION CONTACT:

Elisabeth Daniel,
IRAREbateandNegotiationprogram@cms.hhs.gov, or (667) 290–8793.

SUPPLEMENTARY INFORMATION: The Inflation Reduction Act of 2022 (IRA) was signed into law on August 16, 2022. Sections 11001 and 11002 of the IRA established the Medicare Drug Price Negotiation Program (hereafter the “Negotiation Program”) to negotiate maximum fair prices (MFPs) for certain high expenditure, single source drugs and biological products. The requirements for this program are described in sections 1191 through 1198 of the Social Security Act as added by sections 11001 and 11002 of the IRA. The draft guidance specifies the requirements for manufacturer effectuation of the MFPs in 2028.

To obtain copies of the draft guidance and other IRA-related documents, please access the CMS website by copying and pasting the following web address into your web browser: <https://www.cms.gov/inflation-reduction-act-and-medicare>. If interested in receiving CMS Inflation Reduction Act updates by email, individuals may sign up for CMS’ IRA email updates at <https://www.cms.gov/priorities/medicare-prescription-drug-affordability/medicare-prescription-drug-affordability>.

The Administrator of the Centers for Medicare & Medicaid Services (CMS), Dr. Mehmet Oz, having reviewed and approved this document, authorizes Chyana Woodyard, who is the Federal Register Liaison, to electronically sign this document for purposes of publication in the **Federal Register**.

Chyana Woodyard,

Federal Register Liaison, Centers for Medicare & Medicaid Services.

[FR Doc. 2026–14583 Filed 7–16–26; 4:15 pm]

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DEPARTMENT OF HEALTH AND HUMAN SERVICES

Administration for Children and Families

Privacy Act of 1974; System of Records

AGENCY: Office of Family Assistance (OFA), Administration for Children and

Families (ACF), Department of Health and Human Services (HHS).

ACTION: Extension of comment period for and the effective date of the new routine use under modified system of records.

SUMMARY: The Department of Health and Human Services (HHS) is extending the comment period for and the effective date of the new routine use under the modified system of records maintained by the Office of Family Assistance (OFA) within HHS’ Administration for Children and Families (ACF), System No. 09–80–0375, Temporary Assistance for Needy Families (TANF) Data that appeared in the **Federal Register** of June 23, 2026. We are taking this action in response to an issue that prevented the public from commenting through the Federal eRulemaking Portal until July 10, 2026.

DATES: ACF is extending the comment period for and the effective date of the new routine use under the modified system of records published June 23, 2026 (91 FR 37406). In accordance with 5 U.S.C. 552a(e)(4) and (11), the notice was effective June 23, 2026, with the exception of subparagraph a under routine use 1 and the new routine use 10, which will be effective August 11, 2026. Please submit any comments on the notice by August 11, 2026.

ADDRESSES: Comments may be submitted to the Federal eRulemaking Portal electronically at <http://www.regulations.gov> or mailed to John Talieri, Senior Official for Privacy, Administration for Children and Families, 330 C Street SW, Washington DC 20201. Please include “09–80–0375” in the subject line or [regulations.gov](https://www.regulations.gov) comment. Comments received will be available at [regulations.gov](https://www.regulations.gov) for public viewing, inspection or copies. ACF does not edit personally identifiable information from submissions; therefore, commenters should submit only information that they wish to make publicly available.

FOR FURTHER INFORMATION CONTACT:

General questions about the modified system of records may be submitted by mail or email to TANF Data Division, Office of Family Assistance, Administration for Children and Families, 330 C Street SW, Washington, DC 20201, or tanfdata@acf.hhs.gov; or may be submitted by telephone to John Talieri, Senior Official for Privacy, at (202) 969–3581.