has been posted on the *https://www.regulations.gov* website—as legally required by FTC Rule 4.9(b)—we cannot redact or remove your comment from that website, unless you submit a confidentiality request that meets the requirements for such treatment under FTC Rule 4.9(c), and the General Counsel grants that request.

Visit the FTC website at *https://www.ftc.gov* to read this document and the news release describing the proposed settlement. The FTC Act and other laws the Commission administers permit the collection of public comments to consider and use in this proceeding, as appropriate. The Commission will consider all timely and responsive public comments it receives on or before October 22, 2026. For information on the Commission's privacy policy, including routine uses permitted by the Privacy Act, see *https://www.ftc.gov/site-information/privacy-policy.*

Analysis of Proposed Consent Order To Aid Public Comment

The Federal Trade Commission (''Commission'') has accepted, subject to final approval, an agreement containing a consent order from Corpay, Inc. (formerly known as FleetCor Technologies, Inc.) and its CEO, Ronald Clarke (''Respondents''). The proposed consent order (''Proposed Order'') has been placed on the public record for 30 days for receipt of comments from interested persons. Comments received during this period will become part of the public record. After 30 days, the Commission will again review the agreement and the comments received, then decide whether it should withdraw from the agreement and take appropriate action or make final the agreement's Proposed Order.

The Commission's five-count complaint in this matter alleges that Respondents, who market and sell ''fuel cards'' that can be used to make purchases at gas stations and similar fueling locations, violated section 5 of the FTC Act in two principal ways. First, Respondents unfairly charged their customers, who overwhelmingly are small businesses, a variety of unauthorized fees (Counts IV & V). Specifically, Defendants charged late fees to customers who had paid on time and also charged a number of other unauthorized fees that they hid from their customers. Second, Respondents' marketing variously misrepresented the gas savings (Count I), fraud-control features (Count II), and fees (Count III) associated with Defendants' fuel cards.

The FTC alleged identical claims against these Respondents in a complaint filed in the United States District Court for the Northern District of Georgia. After more than two-and-a-half years of litigation, the district court determined that both Respondents had violated the FTC Act and entered a permanent injunction that requires consent before charging customers, prohibits misrepresentations, and bars other unlawful conduct. *FTC* v. *Fleetcor Techs., Inc.,* 620 F. Supp. 3d 1268 (N.D. Ga. 2022); *FTC* v. *FleetCor Techs., Inc.,* No. 19–5727, 2023 WL 5030099 (N.D. Ga. June 8, 2023).

The Court of Appeals for the Eleventh Circuit affirmed that Respondent Corpay, Inc. is liable on all five counts of the complaint and affirmed the permanent injunction against it. *FTC* v. *Corpay, Inc.,* 164 F.4th 807 (11th Cir. 2026). The court of appeals determined that Respondent Clarke is liable on Counts I, III, IV, and V, but not on Count II, and vacated the injunction against Clarke in light of this determination. Pursuant to the proposed Agreement Containing Consent Order, Respondents would not oppose the entry against Respondent Clarke of the same permanent injunction that the district court previously entered against him, except omitting as to Clarke two subparts that relate to Count II.

The Proposed Order contains monetary relief and related provisions to redress customers injured by Respondents' unfair and deceptive practices. Provision I requires Respondents to pay the Commission $100,000,000 in monetary relief. Provision II describes the procedures and legal rights related to that payment. Provision III requires Respondents to provide customer information to enable the Commission to efficiently administer consumer redress. Provision IV requires Respondents to submit acknowledgements of receipt of the Order. Provision V provides the effective dates of the order, including that, as long as Respondents have met all their obligations under the order, it will terminate in 20 years.

The purpose of this analysis is to aid public comment on the Proposed Order. It is not intended to constitute an official interpretation of the complaint or Proposed Order, or to modify in any way the Proposed Order's terms.

By direction of the Commission.

April J. Tabor,

Secretary.

[FR Doc. 2026–19289 Filed 9–21–26; 8:45 am]

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DEPARTMENT OF HEALTH AND HUMAN SERVICES

Centers for Medicare & Medicaid Services

[Document Identifiers: CMS–10142]

Agency Information Collection Activities: Proposed Collection; Comment Request

AGENCY: Centers for Medicare & Medicaid Services, 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' intention to collect information from the public. Under the Paperwork Reduction Act of 1995 (PRA), federal agencies are required to publish notice in the **Federal Register** concerning each proposed collection of information (including each proposed extension or reinstatement of an existing collection of information) and to allow 60 days for public comment on the proposed action. Interested persons are invited to send comments regarding our burden estimates or any other aspect of this collection of information, including the necessity and utility of the proposed information collection for the proper performance of the agency's functions, the accuracy of the estimated burden, ways to enhance the quality, utility, and clarity of the information to be collected, and the use of automated collection techniques or other forms of information technology to minimize the information collection burden.

DATES: Comments must be received by November 23, 2026.

ADDRESSES: When commenting, please reference the document identifier or OMB control number. To be assured consideration, comments and recommendations must be submitted in any one of the following ways:

1. *Electronically.* You may send your comments electronically to *http://www.regulations.gov.* Follow the instructions for ''Comment or Submission'' or ''More Search Options'' to find the information collection document(s) that are accepting comments.

2. *By regular mail.* You may mail written comments to the following address: CMS, Office of Strategic Operations and Regulatory Affairs, Division of Regulations Development, Attention: Document Identifier: _____ OMB Control Number: ____, Room C4–26–05, 7500 Security Boulevard, Baltimore, Maryland 21244–1850.

To obtain copies of a supporting statement and any related forms for the proposed collection(s) summarized in this notice, please access the CMS PRA website by copying and pasting the following web address into your web browser: *https://www.cms.gov/Regulations-and-Guidance/Legislation/PaperworkReductionActof1995/PRA-Listing.*

FOR FURTHER INFORMATION CONTACT: William N. Parham at (410) 786–4669.

SUPPLEMENTARY INFORMATION:

Contents

This notice sets out a summary of the use and burden associated with the following information collections. More detailed information can be found in each collection's supporting statement and associated materials (see **ADDRESSES**).

Under the PRA (44 U.S.C. 3501–3520), federal agencies must obtain approval from the Office of Management and Budget (OMB) for each collection of information they conduct or sponsor. The term ''collection of information'' is defined in 44 U.S.C. 3502(3) and 5 CFR 1320.3(c) and includes agency requests or requirements that members of the public submit reports, keep records, or provide information to a third party. Section 3506(c)(2)(A) of the PRA requires federal agencies to publish a 60-day notice in the **Federal Register** concerning each proposed collection of information, including each proposed extension or reinstatement of an existing collection of information, before submitting the collection to OMB for approval. To comply with this requirement, CMS is publishing this notice.

Information Collections

1. *Type of Information Collection Request:* Revision of a currently approved collection; *Title of Information Collection:* Bid Pricing Tool (BPT) for Medicare Advantage (MA) Plans and Prescription Drug Plans (PDP); *Use:* Medicare Advantage organizations (MAO) and Prescription Drug Plans (PDP) are required to submit an actuarial pricing ''bid'' for each plan offered to Medicare beneficiaries for approval by CMS. The MAOs and PDPs use the Bid Pricing Tool (BPT) software to develop their actuarial pricing bid. The competitive bidding process defined by the ''The Medicare Prescription Drug, Improvement, and Modernization Act'' (MMA) applies to both the MA and Part D programs. It is an annual process that encompasses the release of the MA rate book in April, the bid's that plans submit to CMS in June, and the release of the Part D and RPPO benchmarks, which typically occurs in August. *Form Number:* CMS–10142 (OMB control number: 0938–0944); *Frequency:* Annually; *Affected Public:* Private Sector, Business or other for profits, and Not for profits institutions; *Number of Respondents:* 460; *Total Annual Responses:* 11,700; *Total Annual Hours:* 406,000. (For questions regarding this collection contact Rachel Shevland at 410–786–3026 or *Rachel.shevland@cms.hhs.gov.*)

William N. Parham, III,

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

[FR Doc. 2026–19366 Filed 9–21–26; 8:45 am]

BILLING CODE 4169–69–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; Generic Information Collection Request for Health Resources and Services Administration Hotlines, Chatlines, and Online Portals

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 October 22, 2026.

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 Samantha Miller, the HRSA Information Collection Clearance Officer, at *paperwork@hrsa.gov* or call (301) 443–9094.

SUPPLEMENTARY INFORMATION:

Information Collection Request Title: Generic Information Collection Request for Collections Related to HRSA Hotlines, Chatlines, and Online Portals, OMB No. 0906–New.

HRSA administers approximately 20 hotlines, chatlines, and online portals for use by customers, members of the public, and HRSA funding recipients. These hotlines, chatlines, and online portals are administered by HRSA or a contractor on behalf of HRSA. The purpose of information collections under this umbrella ICR package is to allow HRSA to collect information on the operation of HRSA hotlines, chatlines, and online portals to assist HRSA in improving their operation and determining if these services are helpful. In addition to collecting basic demographic information, the information collections would include questions such as reasons for inquiry, topics covered by inquiry, and feedback on provided guidance or the hotline/chatline/online portal user experience. Individual collections approved under this umbrella ICR are referred to as generic or fast-track collections.

An illustrative, but not exhaustive, list of examples of information collection activities that would fall under this collection include standardized questions that are asked during the inquiry; surveys about the inquiry; and information collected about the customer's experience in the use of hotlines, chatlines, and online portals. This umbrella collection covers responses to standardized and survey questions relating to the public's use of HRSA's hotlines, chatlines, and online portals. Responses to any generic information collections under this umbrella collection will be voluntary and low burden. Any collections that raise substantive or policy issues would be approved by a standard ICR package, with the standard clearance periods, as opposed to a generic information collection.

A 60-day notice published in the **Federal Register** on November 18, 2024, vol. 89, No. 22; pp. 90708–09. There were no public comments.

Need and Proposed Use of the Information: The purpose of collections under

this umbrella ICR is for accountability, program management,