PORTIONS CLOSED TO THE PUBLIC:

1. Staff Briefing, Bureau of Enforcement, Investigations, and Compliance Update

CONTACT PERSON FOR MORE INFORMATION: William Cody, Secretary, (202) 523–5725.

William Cody,

Secretary.

[FR Doc. 2023–08654 Filed 4–20–23; 11:15 am]

BILLING CODE 6730–02–P

FEDERAL MARITIME COMMISSION

Sunshine Act Meetings; Withdrawal

AGENCY: Federal Maritime Commission

ACTION: Notice; withdrawal.

SUMMARY: The Federal Maritime Commission published a document in the **Federal Register** of April 17, 2023 concerning the Sunshine Act Meetings for our April 19, 2023 Commission Meeting. The April 17, 2023 document contained dates and information for the meeting that was rescheduled.

FOR FURTHER INFORMATION CONTACT: William Cody, 202–523–5725.

SUPPLEMENTARY INFORMATION:

Withdrawal

This action withdraws the notice in the **Federal Register** of April 17, 2023, FR Doc. 2023–07907, at 88 FR 23422 concerning Sunshine Act Meetings.

The Sunshine Act Meeting was rescheduled to May 3, 2023 by the Commission.

Dated: April 18, 2023.

William Cody,

Secretary.

[FR Doc. 2023–08655 Filed 4–20–23; 4:15 pm]

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FEDERAL TRADE COMMISSION

Agency Information Collection Activities; Proposed Collection; Comment Request; Extension

AGENCY: Federal Trade Commission.

ACTION: Notice.

SUMMARY: The Federal Trade Commission (FTC or Commission) is seeking public comment on its proposal to extend an additional three years the current Paperwork Reduction Act clearance to participate in the Office of Management and Budget program ''Generic Clearance for the Collection of Qualitative Feedback on Service Delivery.'' The current clearance expires on July 31, 2023.

DATES: Comments must be received on or before June 23, 2023.

ADDRESSES: Interested parties may file a comment online or on paper by following the instructions in the Request for Comments part of the **SUPPLEMENTARY INFORMATION** section below. Write ''Paperwork Reduction Act Comment: FTC File No. P072108'' on your comment, and file your comment online at *https://www.regulations.gov* by following the instructions on the web-based form. If you prefer to file your comment on paper, mail your comment to the following address: Federal Trade Commission, Office of the Secretary, 600 Pennsylvania Avenue NW, Suite CC–5610 (Annex J), Washington, DC 20580, or deliver your comment to the following address: Federal Trade Commission, Office of the Secretary, Constitution Center, 400 7th Street SW, 5th Floor, Suite 5610 (Annex J), Washington, DC 20024.

FOR FURTHER INFORMATION CONTACT: Bridget Small, Federal Trade Commission, 600 Pennsylvania Avenue NW, CC–10402, Washington, DC 20580, (202) 326–3266.

SUPPLEMENTARY INFORMATION:

Title of Collection: Generic Clearance for the Collection of Qualitative Feedback on Agency Service Delivery.

OMB Control Number: 3084–0159.

Current Actions: Extension of approval for a collection of information.

Affected Public: Individuals and households, businesses and organizations, State, local or Tribal government.

Estimated Number of Annual Respondents: 5,700.

Estimated Total Annual Burden Hours: 900.

Abstract: The FTC seeks renewed Office of Management and Budget approval of its generic clearance to collect qualitative feedback on service delivery (*i.e.,* the products and services that the FTC provides to help consumers and businesses understand their rights and responsibilities, including websites, blogs, videos, print publications, and other content). ''Qualitative feedback'' denotes information that provides useful insight about public perceptions and opinions, but does not include statistical surveys that yield quantitative results that can be generalized to the population of study. The solicitation of feedback on service delivery will target areas such as timeliness, appropriateness, accuracy of information, courtesy, efficiency of service delivery, and resolution of issues with service delivery. The FTC will collect, analyze, and interpret information it gathers through this generic clearance program to identify strengths and weaknesses of current services and make improvements in service delivery based on feedback. This generic clearance for qualitative information will not be used for quantitative information collections that are designed to yield reliably actionable results, such as monitoring trends over time or documenting program performance.

The types of collections that the proposed generic clearance covers include, but are not limited to:

- Customer comment cards/ complaint forms;
- Small discussion groups;
- Focus groups of customers, potential customers, delivery partners, or other stakeholders;
- Cognitive laboratory studies, such as those used to refine questions or assess usability of a website;
- Qualitative customer satisfaction surveys (*e.g.,* post-transaction surveys; opt-out web surveys);
- In-person observation testing (*e.g.,* website or software usability tests).

An agency may not conduct or sponsor, and a person is not required to respond to, a collection of information unless it displays a currently valid OMB control number.

As required by section 3506(c)(2)(A) of the PRA, 44 U.S.C. 3506(c)(2)(A), the FTC is providing this opportunity for public comment before requesting that OMB extend the existing clearance for the information collection requirements contained in the Generic Clearance.

Request for Comments

Pursuant to section 3506(c)(2)(A) of the PRA, the FTC invites comments on: (1) 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) the accuracy of the agency's estimate of the burden of the proposed collection of information, including the validity of the methodology and assumptions used; (3) ways to enhance the quality, utility, and clarity of the information to be collected; and (4) ways to minimize the burden of maintaining records and providing disclosures to consumers. All comments must be received on or before June 23, 2023.

You can file a comment online or on paper. For the FTC to consider your comment, we must receive it on or before June 23, 2023. Write ''Paperwork Reduction Act Comment: FTC File No. P072108'' on your comment. Your comment—including your name and your state—will be placed on the public

record of this proceeding, including the *https://www.regulations.gov* website.

Due to the public health emergency in response to the COVID–19 outbreak and the agency's heightened security screening, postal mail addressed to the Commission will be subject to delay. We encourage you to submit your comments online through the *https://www.regulations.gov* website.

If you prefer to file your comment on paper, write ''Paperwork Reduction Act Comment: FTC File No. P072108'' on your comment and on the envelope, and mail your comment to the following address: Federal Trade Commission, Office of the Secretary, 600 Pennsylvania Avenue NW, Suite CC–5610 (Annex J), Washington, DC 20580; or deliver your comment to the following address: Federal Trade Commission, Office of the Secretary, Constitution Center, 400 7th Street SW, 5th Floor, Suite 5610 (Annex J), Washington, DC 20024. If possible, submit your paper comment to the Commission by courier or overnight service.

Because your comment will become publicly available at *https://www.regulations.gov,* you are solely responsible for making sure that your comment does not include any sensitive or confidential information. In particular, your comment should not include any sensitive personal information, such as your or anyone else's Social Security number; date of birth; driver's license number or other state identification number, or foreign country equivalent; passport number; financial account number; or credit or debit card number. You are also solely responsible for making sure that your comment does not include any sensitive health information, such as medical records or other individually identifiable health information. In addition, your comment should not include any ''trade secret or any commercial or financial information which . . . is privileged or confidential''—as provided by Section 6(f) of the FTC Act, 15 U.S.C. 46(f), and FTC Rule 4.10(a)(2), 16 CFR 4.10(a)(2)—including in particular competitively sensitive information such as costs, sales statistics, inventories, formulas, patterns, devices, manufacturing processes, or customer names.

Comments containing material for which confidential treatment is requested must be filed in paper form, must be clearly labeled ''Confidential,'' and must comply with FTC Rule 4.9(c). In particular, the written request for confidential treatment that accompanies the comment must include the factual and legal basis for the request, and must identify the specific portions of the comment to be withheld from the public record. *See* FTC Rule 4.9(c). Your comment will be kept confidential only if the General Counsel grants your request in accordance with the law and the public interest. Once your comment has been posted publicly at *www.regulations.gov,* we cannot redact or remove your comment 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.

The FTC Act and other laws that 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 that it receives on or before June 23, 2023. 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.*

Josephine Liu,

Assistant General Counsel for Legal Counsel.

[FR Doc. 2023–08533 Filed 4–21–23; 8:45 am]

BILLING CODE 6750–01–P

DEPARTMENT OF HEALTH AND HUMAN SERVICES

Centers for Disease Control and Prevention

[60Day–23–0740; Docket No. CDC–2023–0026]

Proposed Data Collection Submitted for Public Comment and Recommendations

AGENCY: Centers for Disease Control and Prevention (CDC), Department of Health and Human Services (HHS).

ACTION: Notice with comment period.

SUMMARY: The Centers for Disease Control and Prevention (CDC), as part of its continuing effort to reduce public burden and maximize the utility of government information, invites the general public and other federal agencies the opportunity to comment on a continuing information collection, as required by the Paperwork Reduction Act of 1995. This notice invites comment on a proposed information collection project titled Medical Monitoring Project (MMP). The purpose of this data collection is to guide national and local HIV-related service organization and delivery, and monitor receipt of HIV treatment and prevention services and clinical outcomes.

DATES: CDC must receive written comments on or before June 23, 2023.

ADDRESSES: You may submit comments, identified by Docket No. CDC–2023–0026 by either of the following methods:

- *Federal eRulemaking Portal: www.regulations.gov.* Follow the instructions for submitting comments.
- *Mail:* Jeffrey M. Zirger, Information Collection Review Office, Centers for Disease Control and Prevention, 1600 Clifton Road NE, MS H21–8, Atlanta, Georgia 30329.

Instructions: All submissions received must include the agency name and Docket Number. CDC will post, without change, all relevant comments to *www.regulations.gov.*

Please note: Submit all comments through the Federal eRulemaking portal (*www.regulations.gov*) or by U.S. mail to the address listed above.

FOR FURTHER INFORMATION CONTACT: To request more information on the proposed project or to obtain a copy of the information collection plan and instruments, contact Jeffrey M. Zirger, Information Collection Review Office, Centers for Disease Control and Prevention, 1600 Clifton Road NE, MS H21–8, Atlanta, Georgia 30329; Telephone: 404–639–7570; Email: *omb@cdc.gov.*

SUPPLEMENTARY INFORMATION: Under the Paperwork Reduction Act of 1995 (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. In addition, the PRA also requires federal agencies to provide a 60-day notice in the **Federal Register** concerning each proposed collection of information, including each new proposed collection, each proposed extension of existing collection of information, and each reinstatement of previously approved information collection before submitting the collection to the OMB for approval. To comply with this requirement, we are publishing this notice of a proposed data collection as described below.

The OMB is particularly interested in comments that will help:

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 agency's 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;