received from the public. EPA received 32 unique letters on Preliminary Plan 15 that generally provided suggestions for improving certain analyses as well as comments for EPA to consider as it continues work on the detailed studies and ongoing rulemakings discussed in the Preliminary Plan. EPA considered public comments and made appropriate revisions. EPA did not change course on any substantive areas of ELG work as a result.

Plan 15 provides updates on EPA's reviews of industrial wastewater discharges and treatment technologies discussed in Preliminary Plan 15 and presents EPA's 2021 annual review of effluent guidelines and pretreatment standards. With this Plan, EPA continued to focus on and evaluate the extent and nature of Per- and Polyfluoroalkyl Substances (PFAS) discharges and assess opportunities for limiting those discharges from multiple industrial categories, as outlined in EPA's 2021 PFAS Strategic Roadmap, found here: *https://www.epa.gov/pfas/pfas-strategic-roadmap-epas-commitments-action-2021-2024.*

Plan 15 announces that EPA, pending resource availability, intends to initiate one new rulemaking and several new studies. After collecting and analyzing data, EPA has determined that revisions to the effluent limitations guidelines and pretreatment standards for the Landfills Category (40 CFR part 445) are warranted, considering PFAS found in landfill leachate. Plan 15 also announces EPA's intent to expand the detailed study of the Textile Mills Category (40 CFR part 410) to gather information on the use, treatment, and discharge of PFAS from the industry. Plan 15 also announces EPA's intent to initiate a Publicly Owned Treatment Works (POTW) Influent Study focusing on collecting nationwide data pertaining to industrial discharges of PFAS to POTWs, including categories recently reviewed. Finally, Plan 15 announces EPA's intent to undertake a detailed study of the Concentrated Animal Feeding Operations (CAFO) Category (40 CFR part 412), which will focus on collecting further information to enable the agency to make an informed, reasoned decision whether to undertake rulemaking to revise the ELGs for CAFOs.

Plan 15 also announces that EPA is not pursuing further regulatory action for the Electrical and Electronic Components (E&EC) Category (40 CFR part 469) at this time but will continue monitoring this category for PFAS discharge data through the POTW Influent Study. Plan 15 announces that EPA will also continue to monitor PFAS applications and discharges from the Pulp, Paper, and Paperboard Category (40 CFR part 430) and airports. The agency intends to undertake the actions outlined in this Plan and summarized above. The commencement and pace of these activities depend on the agency's Fiscal Year 2023 appropriations and operating plan.

Finally, Plan 15 provides updates related to several actions that are included in EPA's Regulatory Agenda, including the Steam Electric Power Generating Category rulemaking to strengthen certain wastewater pollution discharge limitations for coal fired power plants that use steam to generate electricity; Meat and Poultry Products Category to address nutrient discharges; Organic Chemicals, Plastics & Synthetic Fibers Category to address PFAS discharges; and Metal Finishing and Electroplating Categories to address PFAS discharges.

Plan 15 can be found at *https://www.epa.gov/eg/effluent-guidelines-plan.*

Radhika Fox,

Assistant Administrator.

[FR Doc. 2023–01413 Filed 1–30–23; 8:45 am]

BILLING CODE 6560–50–P

FEDERAL ELECTION COMMISSION

Sunshine Act Meetings

FEDERAL REGISTER CITATION NOTICE OF PREVIOUS ANNOUNCEMENT: 88 FR 3412.

PREVIOUSLY ANNOUNCED TIME AND DATE OF THE MEETING: Tuesday, January 24, 2023 at 10:30 a.m. and its continuation at the conclusion of the open meeting on January 26, 2023.

CHANGES IN THE MEETING: The meeting also discussed:

Matters relating to internal personnel decisions, or internal rules and practices.

CONTACT FOR MORE INFORMATION: Judith Ingram, Press Officer, Telephone: (202) 694–1220.

(Authority: Government in the Sunshine Act, 5 U.S.C. 552b).

Vicktoria J. Allen,

Acting Deputy Secretary of the Commission.

[FR Doc. 2023–02076 Filed 1–27–23; 11:15 am]

BILLING CODE 6715–01–P

FEDERAL MARITIME COMMISSION

Privacy Act of 1974; System of Records

AGENCY: Office of the Secretary, Federal Maritime Commission.

ACTION: Notice of a new system of records.

SUMMARY: The Federal Maritime Commission (FMC or Commission) sponsors advisory committees in accordance with the Federal Advisory Committee Act when the FMC or Congress deems it would be beneficial to obtain advice or recommendations on issues or policies under consideration by the FMC. The Commission appoints advisory committee members which may include special government employees, Federal employees, or representatives of outside organizations, such as trade groups or industry. The FMC collects information about applicants to assess their qualifications to serve as a committee member, such as employment history, education, references, and other information relevant to the applicant's qualifications.

DATES: Written comments must be received no later than March 2, 2023. The revisions will become effective as proposed on March 2, 2023 unless the Commission receives comments that would result in a contrary determination.

ADDRESSES: Submit written comments to the Secretary, Federal Maritime Commission, 800 N Capitol Street NW, Washington, DC 20573–0001; or email comments to: *Secretary@fmc.gov* (email comments as an attachment in MS Word or PDF). Include in the Subject Line: Comments on Systems of Records Notice FMC–44.

FOR FURTHER INFORMATION CONTACT: William Cody, Office of the Secretary, 800 N Capitol Street NW, Suite 1046, Washington, DC 20573–0001. (202) 523–5725. *Secretary@fmc.gov.*

SUPPLEMENTARY INFORMATION: This is a new system of records for records related to Federal advisory committees that the Commission supports.

SYSTEM NAME AND NUMBER:

FMC–44 Federal Advisory Committee Files.

SECURITY CLASSIFICATION:

Unclassified.

SYSTEM LOCATION:

Office of the Secretary, Federal Maritime Commission, 800 North Capitol Street NW, Washington, DC 20573–0001.

SYSTEM MANAGER(S):

Secretary, Federal Maritime Commission, 800 N Capitol Street NW, Washington, DC 20573–0001, *Secretary@fmc.gov.*

AUTHORITY FOR MAINTENANCE OF THE SYSTEM:

Federal Advisory Committee Act, 5 U.S.C. appendix 2.

46 U.S.C. chapter 425 (establishing the National Shipper Advisory Committee).

PURPOSE(S) OF THE SYSTEM:

The purpose of this system is to evaluate and select members of advisory committees within the jurisdiction of the FMC, and for the management of advisory committees, including the preparation of reports, documenting membership, and the nomination and appointment of members, member terms, vacancies, acceptance, and separation.

CATEGORIES OF INDIVIDUALS COVERED BY THE SYSTEM:

Appointed members of the Commission's advisory committees, applicants or nominees to the Commission's advisory committees, and individual participants in the Commission's advisory committees' working groups or subcommittees who are not necessarily appointed members of the advisory committees.

CATEGORIES OF RECORDS IN THE SYSTEM:

Applications containing descriptions of educational and professional experience and qualifications, and letters of reference which may also include the following:

- name;
- title;
- home address;
- business address;
- employer and employment history;
- organizational affiliation;
- phone number;
- email address;
- educational institutions attended, and degrees held;
- references/letters of recommendation, and other information relevant to an individual's qualifications to serve on an advisory committee;
- date of birth;
- social security number;
- gender;
- race;
- drivers license number and state of issuance;
- prior residences for purposes of obtaining a credit check and criminal background check and the results of those checks;
- information about the member's position on the committee, including documentation of their appointment, date of appointment, term, date of separation, and reason for separation.

RECORD SOURCE CATEGORIES:

Records are obtained from applicants who seek to serve on advisory committees, references for applicants, Members of Congress, and applicants' former employers. Information may also be obtained from publicly available sources with the applicant's consent.

ROUTINE USES OF RECORDS MAINTAINED IN THE SYSTEM, INCLUDING

CATEGORIES OF USERS AND PURPOSES OF SUCH USES:

In addition to those disclosures generally permitted under 5 U.S.C. 552a(b), the Commission may:

1. Use the records to determine suitability for Committee membership of an applicant or retention of a current member or to make a determination whether to appoint an individual to an advisory committee or retain the individual as an advisory committee member.

2. Provide these records to the Executive Office of the President, the Office of Management and Budget, or the General Services Administration, when necessary, in the administration of the FMC's advisory committees, including complying with reporting obligations;

3. Share advisory committee records with the public when the FMC deems it necessary to inform the public of advisory committee membership or activities. This routine use does not permit disclosure of an individual's home address, date of birth, social security number, gender, race, school or university transcripts, drivers license number and state of issuance, or prior residences.

4. Provide such records during an appropriate proceeding before a court, grand jury, or administrative or adjudicative body or official, when the FMC or other Agency representing the FMC determines the records are relevant and necessary to the proceeding to which the government is a party; or in an appropriate proceeding before an administrative or adjudicative body when the adjudicator determines the records to be relevant to the proceeding,

5. Provide such records to contractors, grantees, experts, consultants, students, and others performing or working on a contract, service, grant, cooperative agreement, or other assignment for the Federal Government when necessary to accomplish an agency function related to this System of Records.

6. Provide such records to appropriate agencies, entities, and persons when (1) the FMC suspects or confirms that there has been a breach of the system of records; (2) the FMC determines that as a result of the suspected or confirmed breach there is a risk of harm to individuals, the FMC (including its information systems, programs, and operations), the Federal Government, or national security; and (3) the disclosure made to such agencies, entities, and persons is reasonably necessary to assist in connection with the FMC's efforts to respond to the suspected or confirmed breach or to prevent, minimize, or remedy such harm.

7. Provide such records to another Federal agency or Federal entity, when the FMC determines that information from this system of records is reasonably necessary to assist the recipient agency or entity in (1) responding to a suspected or confirmed breach or (2) preventing, minimizing, or remedying the risk of harm to individuals, the recipient agency or entity (including its information systems, programs, and operations), the Federal Government, or national security, resulting from a suspected or confirmed breach.

POLICIES AND PRACTICES FOR STORAGE OF RECORDS:

- Physical records are maintained in file folders in a limited access location.
- Electronic records are maintained within the confines of the FMC General Support System (FMC GSS).

POLICIES AND PRACTICES FOR RETRIEVAL OF RECORDS:

- Physical records are indexed alphabetically by name or committee name.
- Electronic records are retrievable by individual name or committee name.

POLICIES AND PRACTICES FOR RETENTION AND DISPOSAL OF RECORDS:

Resumes and references of members and applicants are retained and disposed of when they are superseded, obsolete, or no longer needed, in accordance with General Records Schedule 6.2, Item 050. Records related to the selection and membership of committee members are permanent and transferred to the National Archives and Records Administration when they are 15 years old or the committee is terminated, whichever is sooner, in accordance with General Records Schedule 6.2, Item 010. *https://www.archives.gov/files/records-mgmt/grs/trs30-sch-only.pdf*

ADMINISTRATIVE, TECHNICAL, AND PHYSICAL SAFEGUARDS:

- Access to physical records in this system are limited to those individuals who have a need to know the information for performance of their official duties and who have appropriate clearances or permission.
- Electronic files are safeguarded to meet multiple National Institute of Standards and Technology (NIST)

Security Standards with password and identification protections. File access is limited to individuals who have a need to know the information for performance of their official duties and who have appropriate clearances or permission.

RECORD ACCESS PROCEDURES:

Requests for access to a record should be directed to the Secretary listed at the above address. Requests may be in person or by mail and shall meet the requirements set out in out in 46 CFR 503.65.

CONTESTING RECORD PROCEDURES:

An individual desiring to amend a record shall direct such a request to the Secretary at the above listed address. Such requests shall specify the desired amendments and the reasons therefore and shall meet the requirements set out in of 46 CFR 503.66.

NOTIFICATION PROCEDURES:

Any individual shall be informed whether or not any Commission system of records contains a record pertaining to him or her when requested in accordance with the requirements of 46 CFR 503.63(a).

EXEMPTIONS PROMULGATED FOR THE SYSTEM:

None.

HISTORY:

N/A.

William Cody,

Secretary.

[FR Doc. 2023–01975 Filed 1–30–23; 8:45 am]

BILLING CODE 6730–02–P

DEPARTMENT OF HEALTH AND HUMAN SERVICES

Food and Drug Administration

[Docket No. FDA–2022–N–0863]

Agency Information Collection Activities; Submission for Office of Management and Budget Review; Comment Request; Monthly Monitoring Study

AGENCY: Food and Drug Administration, HHS.

ACTION: Notice.

SUMMARY: The Food and Drug Administration (FDA) is announcing that a proposed collection of information has been submitted to the Office of Management and Budget (OMB) for review and clearance under the Paperwork Reduction Act of 1995.

DATES: Submit written comments (including recommendations) on the collection of information by March 2, 2023.

ADDRESSES: To ensure that comments on the information collection are received, OMB recommends that written comments be submitted to *https://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. The title of this information collection is ''Monthly Monitoring Study.'' Also include the FDA docket number found in brackets in the heading of this document.

FOR FURTHER INFORMATION CONTACT: JonnaLynn Capezzuto, Office of Operations, Food and Drug Administration, Three White Flint North, 10A–12M, 11601 Landsdown St., North Bethesda, MD 20852, 301–796–3794, *PRAStaff@fda.hhs.gov.*

SUPPLEMENTARY INFORMATION: In compliance with 44 U.S.C. 3507, FDA has submitted the following proposed collection of information to OMB for review and clearance.

Monthly Monitoring Study

OMB Control Number 0910–NEW

This information collection supports the development and implementation of FDA public education campaigns related to tobacco use. To reduce the public health burden of tobacco use in the United States and educate the public—especially young people—about the dangers of tobacco use, FDA's Center for Tobacco Products (CTP) is developing and implementing multiple public education campaigns.

FDA launched ''The Real Cost'' in February 2014, seeking to reduce tobacco use among at-risk youth ages 12 to 17 years old in the United States who are open to smoking cigarettes and/or using electronic nicotine delivery systems (ENDS) products, or have already experimented with cigarettes and/or ENDS products. Given the rapidly evolving tobacco landscape in the United States, frequent and nimble data collection strategies are needed to keep pace and provide relevant information to FDA to inform its tobacco prevention media campaign development about changes in tobacco use and emerging products among youth and young adults.

In an effort to inform specified recommendations around ''The Real Cost'' and FDA's other public education programs to reduce tobacco-related death and disease, more research is needed to understand the trends in use and perceptions of novel and emerging tobacco products, as well as awareness and preferences related to emerging tobacco products and specific brands and device types so that FDA can develop new media campaign messages that resonate with youth and young adults. The purpose of the Monthly Monitoring Study is to collect primary data from youth and young adults, ages 15 to 24 years old, in the United States to monitor perceptions and use of emerging and novel tobacco products and emerging trends in brand and device awareness and use.

The study will be conducted using web-based surveys that are self-administered on personal computers or web enabled mobile devices. The study will use an online survey to collect data from up to 27,000 youth and young adults ages 15 to 24 years to monitor perceptions about and trends in use of ENDS and other emerging tobacco products. Participants will be recruited through social media advertisements. To achieve the required pace of data collection, the study will not contact parents of youth under 18 years old for parental permission and will obtain a waiver of parental permission from the institutional review board. The study will include questions about marijuana use to allow the study team to differentiate between use of current and emerging tobacco products and marijuana, which can be used in tobacco products such as ENDS and little cigar/cigarillos. The survey will take approximately 20 minutes to complete per participant. This survey will ask participants to provide feedback on tobacco use and quitting behavior, as well as brand and device preferences, tobacco information sources, peer influence and perceptions, and marijuana use.

The aim of the Monthly Monitoring Study is to answer the following questions:

- What are the trends in brand and device use for ENDS products and other emerging tobacco products among youth and young adults ages 15 to 24 years in the United States? What are their perceptions of these products?
- How is respondent tobacco use affected by environmental factors, including peer influence and other external factors such as COVID–19?
- What are the primary sources of new product information and where are these products purchased/acquired?
- What are the primary sources of health information for ENDS and other emerging tobacco products?

In support of the provisions of the Family Smoking Prevention and Tobacco Control Act that require FDA to protect the public health and to reduce