potential leased access programmers that have provided the following information:

(a) The desired length of a contract term;

(b) the anticipated commencement date for carriage; and

(c) the nature of the programming. All requests for leased access must be made in writing and must specify the date on which the request was sent to the operator. Operators must maintain supporting documentation to justify scheduled rates, including supporting contracts, calculations of the implicit fees, and justifications for all adjustments.

Cable system operators must disclose on their own websites, or through alternate means if they do not have their own websites, a contact name or title, telephone number, and email address for the person responsible for responding to requests for information about leased access channels.

47 CFR 76.971 requires cable operators to provide billing and collection services to leased access programmers unless they can demonstrate the existence of third-party billing and collection services which, in terms of cost and accessibility, offer leased access programmers an alternative substantially equivalent to that offered to comparable non-leased access programmers.

access programmers.
47 CFR 76.975(b) allows any person aggrieved by the failure or refusal of a cable operator to make commercial channel capacity available or to charge rates for such capacity in accordance with the relevant provisions of the statute or the implementing regulations to file a petition for relief with the Commission. Persons alleging that a cable operator's leased access rate is unreasonable must receive a determination of the cable operator's maximum permitted rate from an independent accountant prior to filing a petition. If parties cannot agree on a mutually acceptable accountant within five business days of the programmer's request for a review, they must each select an independent accountant on the sixth business day. These two accountants will then have five business days to select a third independent accountant to perform the review. To account for their more limited resources, operators of systems entitled to small system relief have 14 business days to select an independent accountant when no agreement can be reached.

47 CFR 76.975(c) requires that petitioners attach a copy of the final accountant's report to their petition where the petition is based on

allegations that a cable operator's leased access rates are unreasonable.

47 CFR 76.975(e) provides that the cable operator or other respondent will have 30 days from service of the petition to file an answer. If a leased access rate is disputed, the answer must show that the rate charged is not higher than the maximum permitted rate for such leased access, and must be supported by the affidavit of a responsible company official. If, after an answer is submitted, the staff finds a prima facie violation of our rules, the staff may require a respondent to produce additional information, or specify other procedures necessary for resolution of the proceeding. Replies to answers must be filed within fifteen (15) days after submission of the answer.

Federal Communications Commission. **Katura Jackson**,

Federal Register Liaison Officer. [FR Doc. 2022–20894 Filed 9–26–22; 8:45 am]

BILLING CODE 6712-01-P

FEDERAL RESERVE SYSTEM

Change in Bank Control Notices; Acquisitions of Shares of a Bank or Bank Holding Company

The notificants listed below have applied under the Change in Bank Control Act (Act) (12 U.S.C. 1817(j)) and § 225.41 of the Board's Regulation Y (12 CFR 225.41) to acquire shares of a bank or bank holding company. The factors that are considered in acting on the applications are set forth in paragraph 7 of the Act (12 U.S.C. 1817(j)(7)).

The public portions of the applications listed below, as well as other related filings required by the Board, if any, are available for immediate inspection at the Federal Reserve Bank(s) indicated below and at the offices of the Board of Governors. This information may also be obtained on an expedited basis, upon request, by contacting the appropriate Federal Reserve Bank and from the Board's Freedom of Information Office at https://www.federalreserve.gov/foia/ request.htm. Interested persons may express their views in writing on the standards enumerated in paragraph 7 of the Act.

Comments regarding each of these applications must be received at the Reserve Bank indicated or the offices of the Board of Governors, Ann E. Misback, Secretary of the Board, 20th Street and Constitution Avenue NW, Washington, DC 20551–0001, not later than October 12, 2022.

A. Federal Reserve Bank of Cleveland (Bryan S. Huddleston, Vice President) 1455 East Sixth Street, Cleveland, Ohio 44101–2566. Comments can also be sent electronically to

Comments.applications@clev.frb.org:

1. The Rebecca A. Moorman
Revocable Living Trust, Rebecca A.
Moorman, as trustee, Keith W. Moorman
and Pamela L. Suever, all of Ottoville,
Ohio; Neil R. Moorman, Saline,
Michigan; and Karen S. Andrew,
Brighton, Michigan; to join the Rebecca
Moorman Family Control Group, a
group acting in concert, to retain voting
shares of The Ottoville Bank Company,
Ottoville, Ohio.

Board of Governors of the Federal Reserve System.

Michele Taylor Fennell,

Deputy Associate Secretary of the Board. [FR Doc. 2022–20897 Filed 9–26–22; 8:45 am]

FEDERAL RESERVE SYSTEM

Privacy Act of 1974; System of Records

AGENCY: Board of Governors of the Federal Reserve System.

ACTION: Notice of a modified system of records.

SUMMARY: Pursuant to the provisions of the Privacy Act of 1974, notice is given that the Board of Governors of the Federal Reserve System (Board) proposes to modify an existing system of records, entitled BGFRS-3, "FRB—Medical Records." BGFRS-3 includes information relating to medical examinations and drug testing of current and prospective employees, and any other medical-related information that may be submitted by employees, contractors, candidates for Board employment, and members of the public.

DATES: Comments must be received on or before October 27, 2022. This new system of records will become effective October 27, 2022, without further notice, unless comments dictate otherwise.

The Office of Management and Budget (OMB), which has oversight responsibility under the Privacy Act, requires a 30-day period prior to publication in the **Federal Register** in which to review the system and to provide any comments to the agency. The public is then given a 30-day period in which to comment, in accordance with 5 U.S.C. 552a(e)(4) and (11).

ADDRESSES: You may submit comments, identified by *BGFRS-3*, "*FRB—Medical Records*," by any of the following methods:

- Agency website: https:// www.federalreserve.gov. Follow the instructions for submitting comments at https://www.federalreserve.gov/apps/ foia/proposedregs.aspx.
- Email: regs.comments@ federalreserve.gov. Include SORN name and number in the subject line of the message.
- Fax: (202) 452–3819 or (202) 452–3102.
- *Mail*: Ann E. Misback, Secretary, Board of Governors of the Federal Reserve System, 20th Street and Constitution Avenue NW, Washington, DC 20551.

All public comments will be made available on the Board's website at https://www.federalreserve.gov/apps/foia/proposedregs.aspx as submitted. Accordingly, comments will not be edited to remove any identifying or contact information. Public comments may also be viewed electronically and in-person in Room M–4365A, 2001 C St. NW, Washington, DC 20551, between 9:00 a.m. and 5:00 p.m. during federal business weekdays.

FOR FURTHER INFORMATION CONTACT:

David B. Husband, Senior Counsel, (202) 530–6270, or david.b.husband@frb.gov, Legal Division, Board of Governors of the Federal Reserve System, 20th Street and Constitution Avenue NW, Washington, DC 20551. If you are deaf, hard of hearing, or have a speech disability, please dial 7–1–1 to access telecommunication relay services.

SUPPLEMENTARY INFORMATION: The Board is modifying this system of records to reflect that the Board has eliminated its in-house health unit and to further clarify the scope of the records included in the system. Specifically, the Board is updating the "Categories of Records in the System" section as follows: (a) to remove the outdated references to the collection of information by the Board's Health Unit, which is now defunct; (b) to clarify that the system includes any medical records that an employee, contractor, applicant, or other individual may provide directly to the Board, including medical records relating to certain sick leave and other leave requests and reasonable accommodation requests; (c) to remove the reference to records relating to employees' workers' compensation claims, as those records are maintained by the Department of Labor's Office of Workers' Compensation Programs; (d) to remove the reference to records regarding employees' use of the Board's exercise facilities as the Board does not collect such medical information; and (e) to remove the unclear and redundant reference to records regarding employees' participation in an occupational health services program. In light of these changes, the Board has made corresponding revisions to system's purpose, categories of individuals covered by the system, and the policies and practices for retention and disposal of records section.

The Board is also making general updates to the system. Specifically, the Board is updating the system location, the system manager, the authority for maintenance of the system, the record source categories, and the policies and practices for storage of records. The Board is also taking the opportunity to update the "Routine Uses" section to incorporate a link to the Board's general routine uses. The Board is not amending or establishing any new routine uses.

The Board is also making technical changes to BGFRS-3 consistent with the template laid out in OMB Circular No. A-108. Accordingly, the Board is making technical corrections and nonsubstantive language revisions to the following categories: "Policies and Practices for Storage of Records, "Policies and Practices for Retrieval of Records," "Policies and Practices for Retention and Disposal of Records," "Administrative, Technical and Physical Safeguards," "Record Access Procedures," "Contesting Record Procedures," and "Notification Procedures." The Board is also adding the following new fields: "Security Classification" and "History."

SYSTEM NAME AND NUMBER:

BGFRS-3 "FRB-Medical Records".

SECURITY CLASSIFICATION:

Unclassified.

SYSTEM LOCATION:

Human Resources, Management Division, Board of Governors of the Federal Reserve System, 20th Street and Constitution Avenue NW, Washington, DC 20551. Certain records are also maintained on the Board's behalf by Medical Advisory Services, 1140 19th Street NW, Suite 700, Washington, DC 20036 and Workpartners, 600 Grant Street, 8th Floor, Pittsburgh, PA 15219.

SYSTEM MANAGER:

John Forbes, Program Manager— Employee Life, Human Resources, Management Division, Board of Governors of the Federal Reserve System, 20th Street and Constitution Avenue NW, Washington, DC 20551, (202) 974–7052, or john.b.forbes@ frb.gov.

AUTHORITY FOR MAINTENANCE OF THE SYSTEM:

Sections 10 and 11 of the Federal Reserve Act (12 U.S.C. 244 and 248).

PURPOSE(S) OF THE SYSTEM:

These records are collected and maintained to assist the Board in determining an employee's fitness for duty and eligibility for benefits based on medical information, to respond to reasonable accommodation requests, to assist the Board in providing a safe and healthy working environment, and to comply with Executive Order 12564, Drug-Free Federal Workplace.

CATEGORIES OF INDIVIDUALS COVERED BY THE SYSTEM:

Past and present Board employees, contractors, candidates for Board employment, and members of the public.

CATEGORIES OF RECORDS IN THE SYSTEM:

The majority of records are maintained by the Board's vendor on behalf of the Board and relate to occupational medical examinations of current and prospective employees, including employees subject to fit-forduty requirements, and the drug testing of current and prospective employees under the Board's Drug-Free Workplace Plan. The Board's vendor also maintains historical records relating to the preventive health screenings that the Board previously offered employees through its now-defunct in-house health unit. The Board's Management Division maintains medical records that an employee, contractor, applicant, or other individual may provide directly to the Board, including records relating to certain sick leave and other leave requests and reasonable accommodation requests. The Management Division also maintains information regarding any failed drug testing administered by the Board's vendor and information regarding whether an employee or applicant has passed or failed the medical examinations administered by the Board's vendor.

RECORD SOURCE CATEGORIES:

Information may be provided by the individual to whom the record pertains, medical professionals, and diagnostic laboratories.

ROUTINE USES OF RECORDS MAINTAINED IN THE SYSTEM, INCLUDING CATEGORIES OF USERS AND PURPOSES OF SUCH USES:

General routine uses A, B, C, D, F, G, H, I, and J apply to this system. These general routine uses are located at https://www.federalreserve.gov/files/SORN-page-general-routine-uses-of-board-systems-of-records.pdf and are published in the Federal Register at 83

FR 43872 at 43873–74 (August 28, 2018). Employee medical information that is obtained under the Rehabilitation Act may be used only in accordance with the confidentiality provisions of the Rehabilitation Act. Records may also be used:

- 1. To disclose information to the Board's Thrift Plan, the Board's Group Life Insurance administrators, the Department of Labor, Department of Veterans Affairs, Social Security Administration, Federal Retirement Thrift Investment Board, or a national, state, or local Social Security-type agency, when necessary to adjudicate a claim (filed by or on behalf of the individual) under a retirement, insurance, or health benefit program;
- 2. To disclose information to a federal, state, or local agency to the extent necessary to comply with laws governing reporting of communicable disease or when it is reasonably believed that an individual might have contracted an illness or been exposed to or suffered from a health hazard while employed in the federal workforce;
- 3. To disclose to health insurance carriers that provide a health benefits plan under the Federal Employees Health Benefits Program information that is necessary to verify eligibility for payment of a claim for health benefits; and
- 4. To disclose information to the executor of an individual's estate, the government entity probating a will, a designated beneficiary, or to any person who is responsible for the care of an individual to the extent necessary when the individual to whom a record pertains is deceased, mentally incompetent, or under other legal disability, and to disclose information to an individual's emergency contact, or, if the emergency contact is unavailable, to any person who the Board believes is assisting the individual, when necessary to assist that individual in obtaining any employment benefit or any working condition, such as an accommodation under the Rehabilitation Act of 1973.

POLICIES AND PRACTICES FOR STORAGE OF RECORDS:

Paper records in this system are stored in locked file cabinets with access limited to staff with a need to know. Electronic records are stored on secure servers with access limited to staff with a need to know.

POLICIES AND PRACTICES FOR RETRIEVAL OF RECORDS:

Records can be retrieved by name.

POLICIES AND PRACTICES FOR RETENTION AND DISPOSAL OF RECORDS:

Records pertaining to occupational medical examination of employees are retained for 30 years after the employee's separation or when the Official Personnel Folder (OPF) is destroyed, whichever is longer. All medical records of applicants who do not become Board employees are destroyed 3 years after the end of the year in which the position is filled or the vacancy announcement is closed, whichever is later, unless longer retention is authorized for business use. Medical records relating to reasonable accommodation requests are retained 3 years after employee separation from the Board or when all appeals are concluded whichever is later, but longer retention is authorized if required for business use. Records of positive drug test results in connection with drug testing of employees are destroyed when the employee leaves the Board or when 3 years old, whichever is later. Other medical records are generally destroyed when 3 years old in accordance with applicable records schedules.

ADMINISTRATIVE, TECHNICAL, AND PHYSICAL SAFEGUARDS:

Electronic files are stored on secure servers. The system has the ability to track individual user actions within the system. The audit and accountability controls are based on NIST and Board standards which, in turn, are based on applicable laws and regulations. The controls assist in detecting security violations and performance or other issues in the system. Access to the system is restricted to authorized users within the Board who require access for official business purposes. Users are classified into different roles and common access and usage rights are established for each role. User roles are used to delineate between the different types of access requirements such that users are restricted to data that is required in the performance of their duties. Periodic assessments and reviews are conducted to determine whether users still require access, have the appropriate role, and whether there have been any unauthorized changes.

RECORD ACCESS PROCEDURES:

The Privacy Act allows individuals the right to access records maintained about them in a Board system of records. Your request for access must: (1) contain a statement that the request is made pursuant to the Privacy Act of 1974; (2) provide either the name of the Board system of records expected to contain the record requested or a concise description of the system of

records; (3) provide the information necessary to verify your identity; and (4) provide any other information that may assist in the rapid identification of the record you seek.

Current or former Board employees may make a request for access by contacting the Board office that maintains the record. The Board handles all Privacy Act requests as both a Privacy Act request and as a Freedom of Information Act request. The Board does not charge fees to a requestor seeking to access or amend his/her Privacy Act records.

You may submit your Privacy Act request to the—

Secretary of the Board, Board of Governors of the Federal Reserve System, 20th Street and Constitution Avenue NW, Washington, DC 20551

You may also submit your Privacy Act request electronically by using the Board's request portal at: https://foia.federalreserve.gov/.

CONTESTING RECORD PROCEDURES:

The Privacy Act allows individuals to seek amendment of information that is erroneous, irrelevant, untimely, or incomplete and is maintained in a system of records that pertains to them. To request an amendment to your record, you should clearly mark the request as a "Privacy Act Amendment Request." You have the burden of proof for demonstrating the appropriateness of the requested amendment and you must provide relevant and convincing evidence in support of your request.

Your request for amendment must: (1) provide the name of the specific Board system of records containing the record you seek to amend; (2) identify the specific portion of the record you seek to amend; (3) describe the nature of and reasons for each requested amendment; (4) explain why you believe the record is not accurate, relevant, timely, or complete; and (5) unless you have already done so in a related Privacy Act request for access or amendment, provide the necessary information to verify your identity.

NOTIFICATION PROCEDURES:

Same as "Access procedures" above. You may also follow this procedure in order to request an accounting of previous disclosures of records pertaining to you as provided for by 5 U.S.C. 552a(c).

EXEMPTIONS PROMULGATED FOR THE SYSTEM:

None.

HISTORY:

This SORN was previously published in the ${\bf Federal\ Register}$ at 73 FR 24984

at 24988 (May 6, 2008). The SORN was also amended to incorporate two new routine uses required by OMB at 83 FR 43872 (August 28, 2018).

Board of Governors of the Federal Reserve System.

Ann E. Misback,

Secretary of the Board.

[FR Doc. 2022–20887 Filed 9–26–22; 8:45 am]

BILLING CODE P

DEPARTMENT OF DEFENSE

GENERAL SERVICES ADMINISTRATION

NATIONAL AERONAUTICS AND SPACE ADMINISTRATION

[OMB Control No. 9000-0136; Docket No. 2022-0053; Sequence No. 18]

Submission for OMB Review; Commercial Acquisitions

AGENCY: Department of Defense (DOD), General Services Administration (GSA), and National Aeronautics and Space Administration (NASA).

ACTION: Notice.

SUMMARY: Under the provisions of the Paperwork Reduction Act, the Regulatory Secretariat Division has submitted to the Office of Management and Budget (OMB) a request to review and approve an extension of a previously approved information collection requirement regarding commercial acquisitions.

DATES: Submit comments on or before October 27, 2022.

ADDRESSES: Written comments and recommendations for this 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.

Additionally, submit a copy to GSA through https://www.regulations.gov and follow the instructions on the site. This website provides the ability to type short comments directly into the comment field or attach a file for lengthier comments.

Instructions: All items submitted must cite OMB Control No. 9000–0136, Commercial Acquisitions. Comments received generally will be posted without change to https://www.regulations.gov, including any personal and/or business confidential information provided. To confirm receipt of your comment(s), please check www.regulations.gov,

approximately two-to-three days after submission to verify posting. If there are difficulties submitting comments, contact the GSA Regulatory Secretariat Division at 202–501–4755 or GSARegSec@gsa.gov.

FOR FURTHER INFORMATION CONTACT:

Zenaida Delgado, Procurement Analyst, at telephone 202–969–7207, or zenaida.delgado@gsa.gov.

SUPPLEMENTARY INFORMATION:

A. OMB Control Number, Title, and Any Associated Form(s)

9000–0136, Commercial Acquisitions.

B. Need and Uses

This clearance covers the information that offerors must submit to comply with the following Federal Acquisition Regulation (FAR) requirements:

FAR 52.212-3, Offeror Representations and Certifications— Commercial Products and Commercial Services. Paragraph (b)(2) requires offerors to identify the applicable paragraphs at (c) through (v) of this provision that the offeror has completed for the purposes of the relevant solicitation only, if any. The provision stipulates that any changes provided by the offeror under paragraph (b)(2) are applicable to that specific solicitation only, and do not result in an update to the representations and certifications posted electronically in the System for Award Management. The contracting officer will use the information to determine a contractor's eligibility for award, and to incorporate appropriate terms and conditions into the contract award.

C. Annual Burden

Respondents: 140,055. Total Annual Responses: 414,909. Total Burden Hours: 207,455.

D. Public Comment

A 60-day notice was published in the **Federal Register** at 87 FR 43039, on July 19, 2022. No comments were received.

Obtaining Copies: Requesters may obtain a copy of the information collection documents from the GSA Regulatory Secretariat Division by calling 202–501–4755 or emailing GSARegSec@gsa.gov. Please cite OMB Control No. 9000–0136, Commercial Acquisitions.

Janet Fry,

Director, Federal Acquisition Policy Division, Office of Governmentwide Acquisition Policy, Office of Acquisition Policy, Office of Governmentwide Policy.

[FR Doc. 2022–20811 Filed 9–26–22; 8:45 am]

BILLING CODE 6820-EP-P

DEPARTMENT OF HEALTH AND HUMAN SERVICES

Food and Drug Administration [Docket No. FDA-2022-N-0571]

Ortho-phthalates for Food Contact Use; Reopening of Comment Period; Request for Information

AGENCY: Food and Drug Administration, HHS.

ACTION: Notice; reopening of comment period; request for information.

SUMMARY: The Food and Drug Administration (FDA or we) is reopening the comment period for the notice titled "Ortho-phthalates for Food Contact Use; Request for Information," which published in the Federal Register of May 20, 2022. We are taking this action in response to a request from stakeholders to extend the comment period to allow additional time for interested parties to develop and submit data, other information, and comments for this request for information.

DATES: FDA is reopening the comment period on the notice "Ortho-phthalates for Food Contact Use; Request for Information," which published in the **Federal Register** on May 20, 2022 (87 FR 31090). Submit either electronic or written comments by December 27, 2022.

ADDRESSES: You may submit comments as follows. Please note that late, untimely filed comments will not be considered. The https://www.regulations.gov electronic filing system will accept comments until 11:59 p.m. Eastern Time at the end of December 27, 2022. Comments received by mail/hand delivery/courier (for written/paper submissions) will be considered timely if they are received on or before that date.

Electronic Submissions

Submit electronic comments in the following way:

• Federal eRulemaking Portal: https://www.regulations.gov. Follow the instructions for submitting comments. Comments submitted electronically, including attachments, to https:// www.regulations.gov will be posted to the docket unchanged. Because your comment will be made public, you are solely responsible for ensuring that your comment does not include any confidential information that you or a third party may not wish to be posted, such as medical information, your or anyone else's Social Security number, or confidential business information, such as a manufacturing process. Please note