

and the upper 30 megahertz via C-V2X deployment is speculative and similarly fails. Therefore, the Commission rejects 5GAA's claim that the Commission's decisions regarding protecting ITS operations in the upper 30 megahertz from unlicensed devices' OOBE are arbitrary and capricious, and the Commission declines to reconsider the indoor unlicensed device OOBE limits adopted in the First Report and Order.

#### Ordering Clauses

Accordingly, *it is ordered* that pursuant to 47 CFR 1.429, the Petition for Reconsideration filed on June 2, 2021 by Auto Innovators and the Petition for Partial Reconsideration filed on June 2, 2021 by 5GAA *are denied*.

Federal Communications Commission.

**Marlene Dortch,**  
Secretary.

[FR Doc. 2024-07428 Filed 4-8-24; 8:45 am]

BILLING CODE 6712-01-P

#### FEDERAL FINANCIAL INSTITUTIONS EXAMINATION COUNCIL

[Docket No. AS24-09]

#### Appraisal Subcommittee Notice of Meeting

**AGENCY:** Appraisal Subcommittee of the Federal Financial Institutions Examination Council

**ACTION:** Notice of Special Closed Meeting.

*Description:* In accordance with Section 1104(b) of Title XI of the Financial Institutions Reform, Recovery, and Enforcement Act of 1989, as amended, notice is hereby given that the Appraisal Subcommittee (ASC) met for a Special Closed Meeting on this date.

*Location:* Virtual meeting via Webex.

*Date:* April 3, 2024.

*Time:* 10:55 a.m. ET.

#### Action and Discussion Item

##### Personnel Matter

The ASC convened a Special Closed Meeting to discuss a personnel matter pursuant to section 1104(b) of Title XI (12 U.S.C. 3333(b)). No action was taken by the ASC.

**James R. Park,**  
Executive Director.

[FR Doc. 2024-07472 Filed 4-8-24; 8:45 am]

BILLING CODE 6700-01-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 received are subject to public disclosure. In general, comments received will be made available without change and will not be modified to remove personal or business information including confidential, contact, or other identifying information. Comments should not include any information such as confidential information that would not be appropriate for public disclosure.

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 April 24, 2024.

*A. Federal Reserve Bank of Minneapolis* (Stephanie Weber, Assistant Vice President) 90 Hennepin Avenue, Minneapolis, Minnesota 55480-0291. Comments may also be sent electronically to [MA@mpls.frb.org](mailto:MA@mpls.frb.org):

1. *Frederick C. Lewis II, Duluth, Minnesota*; to retain voting shares of North Shore Financial Corporation and thereby indirectly retain voting shares of North Shore Bank of Commerce, both of Duluth, Minnesota.

Board of Governors of the Federal Reserve System.

**Michele Taylor Fennell,**  
Deputy Associate Secretary of the Board.

[FR Doc. 2024-07506 Filed 4-8-24; 8:45 am]

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#### FEDERAL RESERVE SYSTEM

#### Formations of, Acquisitions by, and Mergers of Bank Holding Companies

The companies listed in this notice have applied to the Board for approval, pursuant to the Bank Holding Company Act of 1956 (12 U.S.C. 1841 *et seq.*) (BHC Act), Regulation Y (12 CFR part 225), and all other applicable statutes and regulations to become a bank holding company and/or to acquire the assets or the ownership of, control of, or the power to vote shares of a bank or bank holding company and all of the banks and nonbanking companies owned by the bank holding company, including the companies listed below.

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 the BHC Act (12 U.S.C. 1842(c)).

Comments received are subject to public disclosure. In general, comments received will be made available without change and will not be modified to remove personal or business information including confidential, contact, or other identifying information. Comments should not include any information such as confidential information that would not be appropriate for public disclosure.

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 May 9, 2024.

*A. Federal Reserve Bank of Atlanta* (Erien O. Terry, Assistant Vice President) 1000 Peachtree Street NE, Atlanta, Georgia 30309. Comments may also be submitted at [Applications.Comments@atl.frb.org](mailto:Applications.Comments@atl.frb.org):

1. *Volunteer State Bancshares, Inc., Portland, Tennessee*; to merge with Fourth Capital Holdings, Inc., and therefore indirectly acquire Fourth Capital Bank, both of Nashville, Tennessee.

Board of Governors of the Federal Reserve System.

**Michele Taylor Fennell,**

*Deputy Associate Secretary of the Board.*

[FR Doc. 2024-07504 Filed 4-8-24; 8:45 am]

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## GENERAL SERVICES ADMINISTRATION

[Notice-IE-2024-03; Docket No. 2024-0001; Sequence No. 9]

### Privacy Act of 1974; Rescindment of a System of Records

**AGENCY:** Office of the Chief Privacy Officer; General Services Administration, (GSA).

**ACTION:** Rescindment of a system of records notice.

**SUMMARY:** Pursuant to the Privacy Act of 1974 and Office of Management and Budget (OMB) Circular No. A-108, notice is hereby given that the GSA proposes to rescind the GSA/Transit-1, Transportation Benefits Records, System of Records Notice (SORN). This system of records contains information entered by GSA and provides transportation fringe benefits to employees who use mass transportation to commute to and from work.

**DATES:** Effective immediately.

**ADDRESSES:** Comments may be submitted to the Federal eRulemaking Portal, <http://www.regulations.gov>. Submit comments by searching for Notice-IE-2024-03, GSA/Transit-1.

**FOR FURTHER INFORMATION CONTACT:** Call or email Richard Speidel, Chief Privacy Officer at 202-969-5830 and [gsa.privacyact@gsa.gov](mailto:gsa.privacyact@gsa.gov).

**SUPPLEMENTARY INFORMATION:** GSA proposes to rescind a System of Records, GSA/Transit-1. This Notice is being rescinded due to the records of GSA/Transit-1 being maintained under DOT/ALL-8, Parking and Transit Benefit System, managed by the Department of Transportation (DOT). The records under GSA/Transit-1 were transitioned to the DOT in 2017 and are now being maintained under DOT/ALL-8.

## SYSTEM NAME AND NUMBER:

Transportation Benefits Records, GSA/TRANSIT-1.

## HISTORY:

A SORN was previously published in the **Federal Register** at 76 FR 56762 on October 14, 2011.

**Richard Speidel,**

*Chief Privacy Officer, Office of the Deputy Chief Information Officer, General Services Administration.*

[FR Doc. 2024-07430 Filed 4-8-24; 8:45 am]

**BILLING CODE 6820-AB-P**

## DEPARTMENT OF HEALTH AND HUMAN SERVICES

### Food and Drug Administration

[Docket No. FDA-2024-N-1569]

#### Determination That NALFON (Fenoprofen Calcium) Oral Capsules, Equivalent to 300 Milligram Base, and Other Drug Products Were Not Withdrawn From Sale for Reasons of Safety or Effectiveness

**AGENCY:** Food and Drug Administration, HHS.

**ACTION:** Notice.

**SUMMARY:** The Food and Drug Administration (FDA or Agency) has determined that the drug products listed in this document were not withdrawn from sale for reasons of safety or effectiveness. This determination means that FDA will not begin procedures to withdraw approval of abbreviated new drug applications (ANDAs) that refer to these drug products, and it will allow FDA to continue to approve ANDAs that refer to the products as long as they meet relevant legal and regulatory requirements.

#### FOR FURTHER INFORMATION CONTACT:

Stacy Kane, Center for Drug Evaluation and Research, Food and Drug Administration, 10903 New Hampshire Ave., Bldg. 51, Rm. 6236, Silver Spring, MD 20993-0002, 301-796-8363, [Stacy.Kane@fda.hhs.gov](mailto:Stacy.Kane@fda.hhs.gov).

**SUPPLEMENTARY INFORMATION:** Section 505(j) of the Federal Food, Drug, and

Cosmetic Act (FD&C Act) (21 U.S.C. 355(j)) allows the submission of an ANDA to market a generic version of a previously approved drug product. To obtain approval, the ANDA applicant must show, among other things, that the generic drug product: (1) has the same active ingredient(s), dosage form, route of administration, strength, conditions of use, and (with certain exceptions) labeling as the listed drug, which is a version of the drug that was previously approved, and (2) is bioequivalent to the listed drug. ANDA applicants do not have to repeat the extensive clinical testing otherwise necessary to gain approval of a new drug application (NDA).

Section 505(j)(7) of the FD&C Act requires FDA to publish a list of all approved drugs. FDA publishes this list as part of the “Approved Drug Products With Therapeutic Equivalence Evaluations,” which is generally known as the “Orange Book.” Under FDA regulations, a drug is removed from the list if the Agency withdraws or suspends approval of the drug’s NDA or ANDA for reasons of safety or effectiveness or if FDA determines that the listed drug was withdrawn from sale for reasons of safety or effectiveness (21 CFR 314.162).

Under § 314.161(a) (21 CFR 314.161(a)), the Agency must determine whether a listed drug was withdrawn from sale for reasons of safety or effectiveness: (1) before an ANDA that refers to that listed drug may be approved, (2) whenever a listed drug is voluntarily withdrawn from sale and ANDAs that refer to the listed drug have been approved, and (3) when a person petitions for such a determination under 21 CFR 10.25(a) and 10.30. Section 314.161(d) provides that if FDA determines that a listed drug was withdrawn from sale for safety or effectiveness reasons, the Agency will initiate proceedings that could result in the withdrawal of approval of the ANDAs that refer to the listed drug.

FDA has become aware that the drug products listed in the table are no longer being marketed.

Application No.	Drug name	Active ingredient(s)	Strength(s)	Dosage form/route	Applicant
NDA 017604	NALFON	Fenoprofen Calcium	Equivalent to (EQ) 300 Milligrams (mg) Base.	Capsule; Oral	Xspire Pharma.
NDA 017087	ETHRANE	Enflurane	99.9%	Liquid; Inhalation	Baxter Healthcare Corp.
NDA 018801	STERILE WATER FOR INJECTION.	Sterile Water For Injection	100% (1 Milliliter (mL)); 100% (5.2 mL).	Liquid; N/A	Hospira, A Pfizer Company.
NDA 019152	CALAN SR	Verapamil Hydrochloride	120 mg; 180 mg, 240 mg	Tablet, Extended Release; Oral.	Pfizer Inc.
NDA 019885	ACCUPRIL	Quinapril Hydrochloride	EQ 5 mg Base; EQ 10 mg Base; EQ 20 mg Base; EQ 40 mg Base.	Tablet; Oral	Pfizer Pharmaceuticals Ltd.