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 April 26, 2023.

A. Federal Reserve Bank of Chicago (Colette A. Fried, Assistant Vice President) 230 South LaSalle Street, Chicago, Illinois 60690–1414:

1. *David Oren Nelson Living Trust dated January 28, 2022, Memphis, Tennessee; David Oren Nelson, as trustee, Somerville, Tennessee;* to retain voting shares of A.M. Saylor, Incorporated, and thereby indirectly retain voting shares of First Hampton Bank, both of Hampton, Iowa.

Board of Governors of the Federal Reserve System.

Ann E. Misback,

Secretary of the Board.

[FR Doc. 2023–07577 Filed 4–10–23; 8:45 am]

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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 April 25, 2023.

A. Federal Reserve Bank of Dallas (Karen Smith, Director, Mergers & Acquisitions) 2200 N Pearl St., Dallas, Texas 75201. Comments can also be sent electronically to *Comments.applications@dal.frb.org:*

1. *Jane Cheever Powell and Suzanne Cheever Goudge, individually and as co-voting person of the Jane Cheever Powell Trust under the Last Will and Testament of Charles E. Cheever, Sr., the Jane Cheever Powell Trust Under the Last Will & Testament of Elizabeth D. Cheever; and Cecelia Daley Cheever, individually and as trustee of the Cecelia Daley Cheever 2012 Trust; and Jean Mary Cheever, individually and as trustee of the Cece Cheever 2019 Stock Trust, the Jean Mary Cheever 2012 Trust, the Charles E. Cheever, III 2020 Family Trust, the Hope Eileen Cheever Descendant's Trust, and the Emmett Hunter Cheever Descendant's Trust; and Joan McKinney Cheever, individually, as trustee of the Joan McKinney Cheever 2012 Trust, Joan M. Cheever, and as co-trustee of the Joan M. Cheever Irrevocable Trust, and the Joan McKinney Cheever 2012 Trust; and Christopher Hance Cheever, individually and as trustee of the Christopher Hance Cheever 2012 Trust; all of San Antonio, Texas;*

Helen Elizabeth Cheever, individually, and Jean Cheever, individually, as trustee of the Helen Elizabeth Cheever Descendant's Trust, all of San Diego, California; and as custodian of the Hope E. Cheever Texas Uniform Transfer to Minors Act (UTMA), and the Emmett Hunter Cheever Texas UTMA; and Hope Eileen Cheever, individually, all of Dallas, Texas;

Dennis C. Quinn, individually and as co-trustee of the Joan M. Cheever Irrevocable Trust, both of San Antonio, Texas; as trustee of the Elizabeth Daley Quinn Descendant's Trust, and Elizabeth Daley Quinn, individually, both of New York, New York; and as trustee of the Austin McKinney Quinn Descendant's Trust, and Austin McKinney Quinn, individually, both of Los Angeles, California;

Sara E. Goudge Brouillard, individually, as trustee of the Sara Goudge Brouillard Descendant's Trust, the John Cyril Goudge Descendant's Trust, the Suzanne Cheever Goudge 2012 Trust, the Katherine McKinney Goudge Descendant's Trust, the Carrie Goudge Dyer Descendant's Trust, and as custodian of the Minor Children A, B and C under the Texas Uniform Transfer to Minors Act, all of San Antonio, Texas;

Carrie Patricia Goudge Dyer, individually and as custodian of the Minor Children D, E and F, under the Texas UTMA, and Nick Dyer, all of Austin, Texas;

John Cyril Goudge, individually, as trustee of the Jean Cheever 2109 Stock Trust, and as custodian of the Minor Children G and H under the Texas UTMA; and James D. Goudge, Jeff Brouillard, and Laura M. Goudge, all individually, all of San Antonio, Texas;

Katherine McKinney Goudge Ankumah, individually and as custodian of the Minor Children I, and J under the Texas UTMA; and Kobi Ankumah, individually, all of Nashville, Tennessee;

Cheever Partners, as authorized signer to the following GST Trusts: the Charles E. Cheever, III GST Trust, both of New Canaan, Connecticut; the Suzanne C. Goudge GST Trust, the Cecelia Daley Cheever GST Trust, the Jean Mary Cheever GST Trust, the Joan M. Cheever GST Trust, and the Christopher Hance Cheever GST Trust, all of San Antonio, Texas; and

Charles Emmett Cheever, III, individually and trustee of the Charles Emmett Cheever, III 2012 Trust, and the Chris Cheever 2109 Stock Trust, all of San Antonio, Texas; and *Regina Cheever, New Canaan, Connecticut;* a group acting in concert to retain voting shares of Broadway Bancshares, Inc., and thereby indirectly retain voting shares of Broadway National Bank, both of San Antonio, Texas.

Board of Governors of the Federal Reserve System.

Ann E. Misback,

Secretary of the Board.

[FR Doc. 2023–07482 Filed 4–10–23; 8:45 am]

BILLING CODE P

DEPARTMENT OF HEALTH AND HUMAN SERVICES

Centers for Medicare & Medicaid Services

[Document Identifiers CMS–10224 & CMS–10242]

Agency Information Collection Activities: Submission for OMB Review; 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 a second opportunity for public comment on the notice. Interested persons are invited to send comments regarding the burden estimate 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 on the collection(s) of information must be received by the OMB desk officer by May 11, 2023.

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 30-day Review—Open for Public Comments'' or by using the search function.

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 Parham at (410) 786–4669.

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. 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 (44 U.S.C. 3506(c)(2)(A)) requires federal agencies to publish a 30-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 that summarizes the following proposed collection(s) of information for public comment:

1. *Type of Information Collection Request:* Revision of a currently approved collection; *Title of Information Collection:* CMS HCPCS Modification to Code Set Form; *Use:* The Healthcare Common Procedure Coding System (HCPCS) Level II code set is one of the standard code sets used for this purpose. The HCPCS Level II code set, also referred to as alpha-numeric codes, is a standardized coding system that is used primarily to identify items, supplies, and services not included in the HCPCS Level I Current Procedural Terminology (CPT®) codes, such as ambulatory services and durable medical equipment, prosthetics, orthotics, and supplies when used in the home or outpatient setting as well as certain drugs and biologicals. Because Medicare and other insurers cover a variety of these services and supplies, HCPCS Level II codes were established for assignment by insurers to identify items on claims. HCPCS Level II classifies similar items or services that are medical in nature into categories for the purpose of efficient claims processing. For each alpha-numeric HCPCS code, there is descriptive terminology that identifies a category of like items.

As stated in 42 CFR Sec. 414.40 (a) CMS establishes uniform national definitions of services, codes to represent services, and payment modifiers to the codes. The HCPCS code set has been maintained and distributed via modifications of codes, modifiers and descriptions, as a direct result of data received from applicants. Thus, information collected in the application is significant to code set maintenance. The HCPCS code set maintenance is an ongoing process, as changes are implemented and updated quarterly (for drug and biological products) and biannual (for non-drug and non-biological items or services); therefore, the process requires continual collection of information from applicants on a quarterly and bi-annual basis. As new technology evolves and new devices, drugs and supplies are introduced to the market, applicants submit applications to CMS requesting modifications to the HCPCS Level II code set. *Form Number:* CMS–10244 (OMB control number: 0938–1042); *Frequency:* Quarterly; *Affected Public:* Private sector, Business or other for-profit; *Number of Respondents:* 250; *Total Annual Responses:* 250; *Total Annual Hours:* 2,500. (For policy questions regarding this collection contact Sundus Ashar at 410–786–0750.)

2. *Type of Information Collection Request:* Extension of a currently approved collection; *Title of Information Collection:* Emergency Ambulance Transports and Beneficiary Signature; *Use:* The statutory authority requiring a beneficiary's signature on a claim submitted by a provider is located in section 1835(a) and in 1814(a) of the Social Security Act (the Act), for Part B and Part A services, respectively. The authority requiring a beneficiary's signature for supplier claims is implicit in sections 1842(b) (3) (B) (ii) and in 1848(g) (4) of the Act. Federal regulations at 42 CFR 424.32(a) (3) state that all claims must be signed by the beneficiary or on behalf of the Beneficiary (in accordance with 424.36). Section 424.36(a) states that the beneficiary's signature is required on a claim unless the beneficiary has died or the provisions of 424.36(b), (c), or (d) apply.

For emergency and nonemergency ambulance transport services, where the beneficiary is physically or mentally incapable of signing the claim (and the beneficiary's authorized representative is unavailable or unwilling to sign the claim), that it is impractical and infeasible to require an ambulance provider or supplier to later locate the beneficiary or the person authorized to sign on behalf of the beneficiary, before submitting the claim to Medicare for payment. Therefore, an exception was created to the beneficiary signature requirement with respect to emergency and nonemergency ambulance transport services, where the beneficiary is physically or mentally incapable of signing the claim, and if certain documentation requirements are met. Thus, we added subsection (6) to paragraph (b) of 42 CFR 424.36. The information required in this ICR is needed to help ensure that services were in fact rendered and were rendered as billed. *Form Number:* CMS–10242 (OMB control number: 0938–1049); *Frequency:* Occasionally; *Affected Public:* Private sector, Business or other for-profit, Not-for-profits institutions; *Number of Respondents:* 10,233; *Total Annual Responses:* 10,954,288; *Total Annual Hours:* 912,492. (For policy questions regarding this collection

contact Sabrina Teferi at 404–562–7251).

Dated: April 5, 2023.

William N. Parham, III,

Director, Paperwork Reduction Staff, Office of Strategic Operations and Regulatory Affairs.

[FR Doc. 2023–07525 Filed 4–10–23; 8:45 am]

BILLING CODE 4120–01–P

DEPARTMENT OF HEALTH AND HUMAN SERVICES

Administration for Children and Families

Submission for OMB Review; Child Care and Development Fund (CCDF) Consumer Education Website and Reports of Serious Injuries and Death

AGENCY: Office of Child Care, Administration for Children and Families, Department of Health and Human Services.

ACTION: Request for Public Comments.

SUMMARY: The Office of Child Care (OCC), Administration for Children and Families (ACF), U.S. Department of Health and Human Services (HHS), is requesting a 3-year extension of the CCDF Consumer Education website and Reports of Serious Incidents and Death (Office of Management and Budget (OMB) #: 0970–0473, expiration date: April 30, 2023). There are no changes requested to the reporting requirements.

DATES: *Comments due within 30 days of publication.* OMB must make a decision about the collection of information between 30 and 60 days after publication of this document in the **Federal Register**. Therefore, a comment is best assured of having its full effect if OMB receives it within 30 days of publication.

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 30-day Review—Open for Public Comments'' or by using the search function. You can also obtain copies of the proposed collection of information by emailing *infocollection@acf.hhs.gov.* Identify all emailed requests by the title of the information collection.

SUPPLEMENTARY INFORMATION:

Description: The existing Consumer Education Website reporting requirement will not be modified and requires states and territories to include information about state or territory policies (related to licensing, monitoring, and background checks) and provider-specific information, including results of monitoring and inspection reports and, if available, information about quality. The existing Reporting of Serious Injuries and Death reporting requirement will not be modified. CCDF Lead Agencies must establish procedures that require child care providers that care for children receiving CCDF subsidies to report to a designated state, territorial, or tribal entity any serious injuries or deaths of children occurring in child care. There are no standard federal forms associated with these reporting requirements.

Respondents: The Consumer Education website information collection requirement applies to the 50 states, the District of Columbia, and 5 territories that receive CCDF grants. Reporting of Serious Injuries and Death is a requirement for child care providers.

ANNUAL BURDEN ESTIMATES

Instrument	Total number of respondents	Total number of responses per respondent	Average burden hours per response	Total burden hours	Annual burden hours
Consumer Education Website	56	1	300	50,400	16,800
Reporting of Serious Injuries and Death	10,000	1	1	30,000	10,000

Estimated Total Annual Burden Hours: 26,800.

Authority: Pub. L. 113–186; 42 U.S.C. 9858 *et seq.*

Mary B. Jones,

ACF/OPRE Certifying Officer.

[FR Doc. 2023–07513 Filed 4–10–23; 8:45 am]

BILLING CODE 4184–43–P

DEPARTMENT OF HEALTH AND HUMAN SERVICES

Food and Drug Administration

[Docket No. FDA–2023–N–1006]

Agency Information Collection Activities; Proposed Collection; Comment Request; Medical Devices; Reports of Corrections and Removals

AGENCY: Food and Drug Administration, HHS.

ACTION: Notice.

SUMMARY: The Food and Drug Administration (FDA, Agency, or we) is announcing an opportunity for public comment on the proposed collection of certain information by the Agency. 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 of an existing collection of information, and to allow 60 days for public comment in response to the notice. This notice solicits comments on information collection associated with reports of removals and corrections for medical and radiation emitting products regulated by FDA's Center for Devices and Radiological Health.

DATES: Either electronic or written comments on the collection of information must be submitted by June 12, 2023.

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 June 12, 2023. 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