the Communications Act of 1934, as amended, 47 U.S.C. 151, 154(i)–(j), 201–205, 211, 214, 219–220, 303(r), 309 and 403

*Total Annual Burden:* 6.5 hours.

*Annual Cost Burden:* $1,850

*Needs and Uses:* The Federal Communications Commission (''Commission'') is requesting that the Office of Management and Budget (OMB) to approve a revision to OMB Control No. 3060–1029—Data Network Identification Code (DNIC). The Commission is developing revised and new electronic forms for this collection as part of the Commission's modernization of its online, web-based electronic filing system—the International Bureau filing system (IBFS). This Supporting Statement seeks approval for the new and revised forms to request an International Signaling Point Code (ISPC), and reflects changes in the costs and burdens associated with these applications.

A Data Network Identification Code (DNIC) is a unique, four-digit number designed to provide discrete identification of individual public data networks. The DNIC is intended to identify and permit automated switching of data traffic to particular networks. The DNIC is the central device of the international data numbering plan developed by the International Telecommunications Union (ITU) and set forth in Recommendation X.121. Prior to the availability of electronic web-based application forms in 1999, the Commission used an informal process for assigning DNICs. In the informal system, a company desiring a code would notify the Commission that it wishes one assigned and demonstrate that it has the ability to originate and terminate international traffic (e.g,, by showing an interconnection arrangement with a U.S. international carrier) and the Commission would assign a DNIC. In 1986, the Commission established procedures for the assignment of DNICs to interested data network operators. Today, the operators of public data networks file an application for a DNIC in IBFS. The DNIC is obtained on a one-time only basis unless there is a change in ownership or the owner chooses to relinquish the code to the Commission.

*IBFS Modernization of DNIC Electronic Forms.* The Commission seeks OMB approval of revisions to its DNIC application form and the addition of new forms that will be electronically filed through IBFS. The new online forms will ensure the Commission collects the information required by the Commission's rules. The use of such online forms will reduce costs and administrative burdens on applicants, resulting in greater efficiencies, and improve transparency to the public. Once the Commission receives approval for the new forms from OMB, as required by section 1.10006 of the Commission's rules, we will announce the availability of mandated e-forms and their effective dates.

Federal Communications Commission.

**Marlene Dortch,**

*Secretary, Office of the Secretary.*

[FR Doc. 2023–07147 Filed 4–5–23; 8:45 am]

**BILLING CODE 6712–01–P**

## 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 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 5, 2023.

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

1. *Grinnell Bancshares, Inc., Grinnell, Iowa;* to acquire The Colorado Bank & Trust Company of La Junta, La Junta, Colorado.

Board of Governors of the Federal Reserve System.

**Michele Taylor Fennell,**

*Deputy Associate Secretary of the Board.*

[FR Doc. 2023–07162 Filed 4–5–23; 8:45 am]

**BILLING CODE P**

## FEDERAL TRADE COMMISSION

### Agency Information Collection Activities; Submission for OMB Review; Comment Request; Extension

**AGENCY:** Federal Trade Commission.

**ACTION:** Notice.

**SUMMARY:** The Federal Trade Commission (''FTC'' or ''Commission'') requests that the Office of Management and Budget (''OMB'') extend for an additional three years the current Paperwork Reduction Act (''PRA'') clearance for information collection requirements associated with its Funeral Industry Practice Rule (''Funeral Rule'' or ''Rule''). That clearance expires on July 31, 2023.

**DATES:** Comments must be filed by May 8, 2023.

**ADDRESSES:** Interested parties may file a comment online or on paper, by following the instructions in the Request for Comment part of the **SUPPLEMENTARY INFORMATION** section below. 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.

**FOR FURTHER INFORMATION CONTACT:** Melissa Dickey, Division of Marketing Practices, Bureau of Consumer Protection, Federal Trade Commission, 600 Pennsylvania Ave. NW, Washington, DC 20580, *mdickey@ftc.gov,* (202) 326–2662.

**SUPPLEMENTARY INFORMATION:**

*Title of Collection:* Funeral Industry Practice Rule, 16 CFR part 453.

*OMB Control Number:* 3084–0025.

*Type of Review:* Extension without change of currently approved collection.

*Abstract:* The Funeral Rule ensures that consumers who are purchasing funeral goods and services have access to accurate itemized price information so they can purchase only the funeral goods and services they want or need. Among other things, the Rule requires a funeral provider to: (1) provide consumers a copy of the funeral provider's General Price List that

itemizes the goods and services it offers; (2) show consumers a Casket Price List and an Outer Burial Container Price List at the outset of any discussion of those items or their prices, and in any event before showing consumers caskets or vaults; (3) provide price information from its price lists over the telephone; and (4) give consumers a Statement of Funeral Goods and Services Selected after determining the funeral arrangements with consumers. The Rule requires that funeral providers disclose this information to consumers and maintain records documenting their compliance with the Rule.

*Affected Public:* Private Sector: Businesses and other for-profit entities.

*Estimated Number of Annual Respondents:* 18,874.

*Estimated Annual Burden Hours:* 173,936.

*Estimated Annual Labor Costs:* $5,387,875.

*Request for Comment:*

On December 19, 2022, the FTC sought public comment on the information collection requirements in the Funeral Rule. 87 FR 77610 (Dec. 19, 2022). No relevant comments were received during the public comment period. Pursuant to OMB regulations, 5 CFR part 1320, that implement the PRA, 44 U.S.C. 3501 *et seq.,* the FTC is providing this second opportunity for public comment while seeking OMB approval to renew the pre-existing clearance for the Rule. For more details about the Rule requirements and the basis for the calculations summarized below, see 87 FR 77610.

Your comment—including your name and your state—will be placed on the public record of this proceeding. Because your comment will be made public, you are solely responsible for making sure that your comment does not include any sensitive personal information, such as anyone'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 ensuring 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 ''[t]rade secret or any commercial or financial information which is . . . privileged or confidential''—as provided in section 6(f) of the FTC Act 15 U.S.C. 46(f), and FTC Rule 4.10(a)(2), 16CFR 4.10(a)(2)—including, in particular, competitively sensitive information, such as costs, sales statistics, inventories, formulas, patterns devices, manufacturing processes, or customer names.

**Josephine Liu,**

*Assistant General Counsel for Legal Counsel.*

[FR Doc. 2023–07186 Filed 4–5–23; 8:45 am]

**BILLING CODE 6750–01–P**

## DEPARTMENT OF HEALTH AND HUMAN SERVICES

### Agency for Healthcare Research and Quality

### Supplemental Evidence and Data Request on Psychosocial and Pharmacologic Interventions for Disruptive Behavior in Children and Adolescents

**AGENCY:** Agency for Healthcare Research and Quality (AHRQ), HHS.

**ACTION:** Request for Supplemental Evidence and Data Submissions.

**SUMMARY:** The Agency for Healthcare Research and Quality (AHRQ) is seeking scientific information submissions from the public. Scientific information is being solicited to inform our review on *Psychosocial and Pharmacologic Interventions for Disruptive Behavior in Children and Adolescents,* which is currently being conducted by the AHRQ's Evidence-based Practice Centers (EPC) Program. Access to published and unpublished pertinent scientific information will improve the quality of this review.

**DATES:** *Submission Deadline* on or before May 8, 2023.

**ADDRESSES:**

*Email submissions: epc@ahrq.hhs.gov.*

*Print submissions:*

*Mailing Address:* Center for Evidence and Practice Improvement, Agency for Healthcare Research and Quality, ATTN: EPC SEADs Coordinator 5600 Fishers Lane, Mail Stop 06E53A, Rockville, MD 20857.

*Shipping Address (FedEx, UPS, etc.):* Center for Evidence and Practice Improvement, Agency for Healthcare Research and Quality, ATTN: EPC SEADs Coordinator, 5600 Fishers Lane, Mail Stop 06E53A, Rockville, MD 20857.

**FOR FURTHER INFORMATION CONTACT:** Kelly Carper, Telephone: 301–427–1656 or Email: *epc@ahrq.hhs.gov.*

**SUPPLEMENTARY INFORMATION:** The Agency for Healthcare Research and Quality has commissioned the Evidence-based Practice Centers (EPC) Program to complete a review of the evidence for *Psychosocial and Pharmacologic Interventions for Disruptive Behavior in Children and Adolescents.* AHRQ is conducting this systematic review pursuant to section 902 of the Public Health Service Act, 42 U.S.C. 299a.

The EPC Program is dedicated to identifying as many studies as possible that are relevant to the questions for each of its reviews. In order to do so, we are supplementing the usual manual and electronic database searches of the literature by requesting information from the public (*e.g.,* details of studies conducted). We are looking for studies that report on Psychosocial and Pharmacologic Interventions for Disruptive Behavior in Children and Adolescents, including those that describe adverse events. The entire research protocol is available online at: *https://effectivehealthcare.ahrq.gov/products/disruptive-behavior/protocol.*

This is to notify the public that the EPC Program would find the following information on Psychosocial and Pharmacologic Interventions for Disruptive Behavior in Children and Adolescents helpful:

- A list of completed studies that your organization has sponsored for this indication. In the list, please *indicate whether results are available on ClinicalTrials.gov along with the ClinicalTrials.gov trial number.*
- *For completed studies that do not have results on ClinicalTrials.gov,* a summary, including the following elements: study number, study period, design, methodology, indication and diagnosis, proper use instructions, inclusion and exclusion criteria, primary and secondary outcomes, baseline characteristics, number of patients screened/eligible/enrolled/lost to follow-up/withdrawn/analyzed, effectiveness/efficacy, and safety results.
- *A list of ongoing studies that your organization has sponsored for this indication.* In the list, please provide the *ClinicalTrials.gov* trial number or, if the trial is not registered, the protocol for the study including a study number, the study period, design, methodology, indication and diagnosis, proper use instructions, inclusion and exclusion criteria, and primary and secondary outcomes.
- Description of whether the above studies constitute *ALL Phase II and above clinical trials* sponsored by your organization for this indication and an index outlining the relevant information in each submitted file.

Your contribution is very beneficial to the Program. Materials submitted must be publicly available or able to be made public. Materials that are considered confidential; marketing materials; study types not included in the review; or