

*Notice:* Section 309(a) of the Clean Air Act requires that EPA make public its comments on EISs issued by other Federal agencies. EPA's comment letters on EISs are available at: <http://www.epa.gov/compliance/nepa/eisdata.html>.

*EIS No. 20170023, Draft Supplement, DOS, MN, Line 67 Expansion (previously known as the Alberta Clipper Pipeline), Comment Period Ends: 03/27/2017, Contact:* Mary D. Hassell 202-736-7428.

*EIS No. 20170024, Revised Final, USFS, SC, Francis Marion Forest Plan Revision, Review Period Ends: 03/13/2017, Contact:* Mary Morrison 803-561-4058.

#### Amended Notices

*EIS No. 20160311, Draft, USACE, NE., Missouri River Recovery Management Plan, Comment Period Ends: 04/24/2017, Contact:* Aaron Quinn 402-995-2669, Revision to FR Notice Published 12/23/2016; Extending Comment Period from 2/24/2017 to 4/24/2017.

Dated: February 7, 2017.

#### Dawn Roberts,

*Management Analyst, NEPA Compliance Division, Office of Federal Activities.*

[FR Doc. 2017-02805 Filed 2-9-17; 8:45 am]

BILLING CODE 6560-50-P

## FEDERAL MARITIME COMMISSION

### Sunshine Act Meeting

**AGENCY HOLDING THE MEETING:** Federal Maritime Commission.

**TIME AND DATE:** February 15, 2017; 10:00 a.m.

**PLACE:** 800 N. Capitol Street NW., First Floor Hearing Room, Washington, DC.

**STATUS:** The meeting will be held in Open Session and will be streamed live at <http://fmc.capitolconnection.org/>.

#### MATTERS TO BE CONSIDERED:

#### Open Session

1. Briefing by Commissioner Doyle on the Competition Law in the Transport Sector Conference
2. Staff Update on Docket No. 16-05: Service Contracts and NVOCC Service Arrangements

**CONTACT PERSON FOR MORE INFORMATION:** Rachel E. Dickon, Assistant Secretary, (202) 523 5725.

**Rachel E. Dickon,**  
*Assistant Secretary.*

[FR Doc. 2017-02881 Filed 2-8-17; 4:15 pm]

BILLING CODE 6731-AA-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 applications listed below, as well as other related filings required by the Board, are available for immediate inspection at the Federal Reserve Bank indicated. The applications will also be available for inspection at the offices of the Board of Governors. Interested persons may express their views in writing on the standards enumerated in the BHC Act (12 U.S.C. 1842(c)). If the proposal also involves the acquisition of a nonbanking company, the review also includes whether the acquisition of the nonbanking company complies with the standards in section 4 of the BHC Act (12 U.S.C. 1843). Unless otherwise noted, nonbanking activities will be conducted throughout the United States.

Unless otherwise noted, comments regarding each of these applications must be received at the Reserve Bank indicated or the offices of the Board of Governors not later than March 9, 2017.

*A. Federal Reserve Bank of St. Louis (David L. Hubbard, Senior Manager) P.O. Box 442, St. Louis, Missouri 63166-2034. Comments can also be sent electronically to [Comments.applications@stls.frb.org](mailto:Comments.applications@stls.frb.org):*

1. *Tyronza Bancshares, Inc., Marked Tree, Arkansas;* to indirectly acquire 6.25 percent of the voting shares of Pinnacle Bancshares, Inc., and thereby indirectly acquire Pinnacle Bank, both of Rogers, Arkansas.

Board of Governors of the Federal Reserve System, February 7, 2017.

**Yao-Chin Chao,**

*Assistant Secretary of the Board.*

[FR Doc. 2017-02800 Filed 2-9-17; 8:45 am]

BILLING CODE P

## DEPARTMENT OF HEALTH AND HUMAN SERVICES

### Centers for Disease Control and Prevention

[60Day-17-170B: Docket No. ATSDR-2017-0002]

### Proposed Data Collection Submitted for Public Comment and Recommendations

**AGENCY:** Agency for Toxic Substances and Disease Registry (ATSDR), Department of Health and Human Services (HHS).

**ACTION:** Notice with comment period.

**SUMMARY:** Agency for Toxic Substances and Disease Registry (ATSDR), as part of its continuing efforts to reduce public burden and maximize the utility of government information, invites the general public and other Federal agencies to take this opportunity to comment on proposed and/or continuing information collections, as required by the Paperwork Reduction Act of 1995. This notice invites comment on a proposed information collection project titled "Characterization of Exposure Potential during Activities Conducted on Synthetic Turf with Crumb Rubber Infill." The purpose of the proposed study is to evaluate and characterize human exposure potential to constituents in crumb rubber infill.

**DATES:** Written comments must be received on or before April 11, 2017.

**ADDRESSES:** You may submit comments, identified by Docket No. ATSDR-2017-0002 by any of the following methods:

- *Federal eRulemaking Portal:* [Regulations.gov](http://Regulations.gov). Follow the instructions for submitting comments.

- *Mail:* Leroy A. Richardson, Information Collection Review Office, Centers for Disease Control and Prevention, 1600 Clifton Road NE., MS-D74, Atlanta, Georgia 30329.

*Instructions:* All submissions received must include the agency name and Docket Number. All relevant comments received will be posted without change to [Regulations.gov](http://Regulations.gov), including any personal information provided. For access to the docket to read background documents or comments received, go to [Regulations.gov](http://Regulations.gov).

*Please note: All public comment should be submitted through the Federal eRulemaking portal ([Regulations.gov](http://Regulations.gov)) or by U.S. mail to the address listed above.*

**FOR FURTHER INFORMATION CONTACT:** To request more information on the proposed project or to obtain a copy of

the information collection plan and instruments, contact the Information Collection Review Office, Centers for Disease Control and Prevention, 1600 Clifton Road NE., MS-D74, Atlanta, Georgia 30329; phone: 404-639-7570; Email: [omb@cdc.gov](mailto:omb@cdc.gov).

**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. In addition, the PRA also requires Federal agencies to provide a 60-day notice in the **Federal Register** concerning each proposed collection of information, including each new proposed collection, each proposed extension of existing collection of information, and each reinstatement of previously approved information collection before submitting the collection to OMB for approval. To comply with this requirement, we are publishing this notice of a proposed data collection as described below.

Comments are invited on: (a) Whether the proposed collection of information is necessary for the proper performance of the functions of the agency, including whether the information shall have practical utility; (b) the accuracy of the agency's estimate of the burden of the proposed collection of information; (c) ways to enhance the quality, utility, and clarity of the information to be collected; (d) ways to minimize the burden of the collection of information on respondents, including through the use of automated collection techniques or other forms of information technology; and (e) estimates of capital or start-up costs and costs of operation, maintenance, and purchase of services to provide information. Burden means the total time, effort, or financial resources expended by persons to generate, maintain, retain, disclose or provide information to or for a Federal agency. This includes the time needed to review instructions; to develop, acquire, install and utilize technology and systems for the purpose of collecting, validating and verifying information, processing and maintaining information, and disclosing and providing information; to train personnel and to be able to respond to a collection of information, to search data sources, to complete and review the collection of information; and to transmit or otherwise disclose the information.

### Proposed Project

Characterization of Exposure Potential during Activities Conducted on Synthetic Turf with Crumb Rubber Infill—New—Agency for Toxic Substances and Disease Registry (ATSDR).

#### Background and Brief Description

Currently in the United States, there are more than 12,000 synthetic turf fields in use. While the Synthetic Turf Council has set guidelines for the content of crumb rubber used as infill in synthetic turf fields, manufacturing processes result in differences among types of crumb rubber. Additionally, the chemical composition may vary highly between different processes and source materials and may vary even within granules from the same origin.

In July, 2016, the Agency for Toxic Substances and Disease Registry (ATSDR) and the United States Environmental Protection Agency (USEPA) were granted an emergency Paperwork Reduction Act (PRA) clearance for a research study titled "Collections Related to Synthetic Turf Fields with Crumb Rubber Infill" (OMB Control No. 0923-0054, expiration date 01/31/2017). The research goals for the three activities in the protocol are pilot-level investigations to evaluate and characterize: (1) The chemical composition and use of crumb rubber infill in synthetic turf using a convenience sample of nine tire recycling manufacturing plants and 40 facilities that use synthetic turf fields (Activity 1); (2) the human exposure potential to constituents in crumb rubber infill among a convenience sample of 60 field users (Activity 2); and (3) collection of biological specimens (blood and urine) from 45 participants from Activity 2 (Activity 3).

By December, 2016, ATSDR and USEPA completed Activity 1 which was aimed at characterizing the chemical composition and use of synthetic turf fields with tire crumb rubber infill. The agencies successfully consented and sampled 40 synthetic turf fields with crumb rubber infill across the United States. The activities are reported in the "Status Report on the Federal Research Action Plan on Recycled Tire Crumb Used on Playing Fields and Playgrounds." The Status Report was released on December 30, 2016.

During Activity 1, ATSDR and USEPA obtained permission to return to participating fields to complete the human exposure characterization. Due

to the limited time constraints and field activity schedules, ATSDR and USEPA chose to begin Activity 2 data collection and Activity 3 specimen collection in 2017.

The agencies are requesting a new information collection request (ICR) for a two-year OMB clearance to complete Activity 2 and Activity 3, now subtitled "Characterization of Exposure Potential during Activities Conducted on Synthetic Turf with Crumb Rubber Infill." This will be the first assessment of activities conducted on synthetic turf for the purpose of characterizing potential exposure patterns. The study will include persons who use synthetic turf with crumb rubber infill (*e.g.*, facility users) and who routinely perform activities that would result in a high level of contact to crumb rubber. This will allow for evaluation of potential high-end exposures to constituents in synthetic turf among this group of users. The respondents will be administered a detailed questionnaire on activity patterns on synthetic turf with crumb rubber infill. This instrument, along with extant videography of persons engaged in activities of interest (see below), will be used to characterize exposure scenarios, including the nature and duration of potential exposures.

Additionally, we will conduct an exposure characterization sub-study among a subset of the respondents. We will use a subset of the facilities sampled in the first study to conduct activities for the exposure characterization of facility users. The exposure characterization sub-study will include field environment sampling, personal air monitoring, dermal sampling, and urine and blood collection. Video data collection of facility user activities will be performed for a subset of respondents. It is likely that some of the collection items will not be analyzed in the current project time frame but will be archived for future analysis.

The research study will screen a total of 75 participants for eligibility. The sample size for the exposure characterization study is 60 respondents and 45 respondents for the exposure measurements sub-study. The total burden hours for the research study is 174 hours among all of the 75 respondents. There is no cost to the respondents other than their time in the study.

## ESTIMATED ANNUALIZED BURDEN HOURS

| Type of respondents                           | Form name                                     | Number of respondents | Number of responses per respondent | Average burden per response (in hours) | Total burden (in hours) |
|---|---|-----------------------|------------------------------------|--|-------------------------|
| Adult/Adolescent Field Users .....            | Eligibility Screening Script .....            | 41                    | 1                                  | 5/60                                   | 4                       |
|   | Adult and Adolescent Questionnaire .....      | 36                    | 1                                  | 30/60                                  | 18                      |
|   | Exposure Measurement Form .....               | 27                    | 1                                  | 3                                      | 81                      |
|   | Phlebotomist Safety Exclusion Questions Form. | 27                    | 1                                  | 2/60                                   | 1                       |
| Parents/Guardians of Youth/Child Field Users. | Eligibility Screening Script .....            | 34                    | 1                                  | 5/60                                   | 3                       |
|   | Youth and Child Questionnaire .....           | 24                    | 1                                  | 30/60                                  | 12                      |
|   | Phlebotomist Safety Exclusion Questions Form. | 18                    | 1                                  | 2/60                                   | 1                       |
| Youth/Child Field Users .....                 | Exposure Measurement Form .....               | 18                    | 1                                  | 3                                      | 54                      |
| Total .....                                   | .....   | .....                 | .....                              | .....                                  | 174                     |

**Leroy A. Richardson,**  
Chief, Information Collection Review Office,  
Office of Scientific Integrity, Office of the  
Associate Director for Science, Office of the  
Director, Centers for Disease Control and  
Prevention.

[FR Doc. 2017-02760 Filed 2-9-17; 8:45 am]

BILLING CODE 4163-18-P

## DEPARTMENT OF HEALTH AND HUMAN SERVICES

### Centers for Disease Control and Prevention

[60Day-17-17ND; Docket No. CDC-2017-0007]

#### Proposed Data Collection Submitted for Public Comment and Recommendations

**AGENCY:** Centers for Disease Control and Prevention (CDC), Department of Health and Human Services (HHS).

**ACTION:** Notice with comment period.

**SUMMARY:** The Centers for Disease Control and Prevention (CDC), as part of its continuing efforts to reduce public burden and maximize the utility of government information, invites the general public and other Federal agencies to take this opportunity to comment on proposed and/or continuing information collections, as required by the Paperwork Reduction Act of 1995. This notice invites comment on a proposed information collection entitled "Grants for Injury Control Research Centers Annual Progress Report (APR)." CDC will collect information from grantees funded under Grants for Injury Control and Research Centers (ICRC) for the Annual Progress Report (APR). The APR is used to monitor the ICRCs' progress on set performance indicators, activities,

and progress towards stated grant objectives.

**DATES:** Written comments must be received on or before April 11, 2017.

**ADDRESSES:** You may submit comments, identified by Docket No. CDC-2017-0007 by any of the following methods:

- *Federal eRulemaking Portal:* *Regulations.gov.* Follow the instructions for submitting comments.

- *Mail:* Leroy A. Richardson, Information Collection Review Office, Centers for Disease Control and Prevention, 1600 Clifton Road NE., MS-D74, Atlanta, Georgia 30329.

*Instructions:* All submissions received must include the agency name and Docket Number. All relevant comments received will be posted without change to *Regulations.gov*, including any personal information provided. For access to the docket to read background documents or comments received, go to *Regulations.gov*.

**Please note:** All public comment should be submitted through the Federal eRulemaking portal (*Regulations.gov*) or by U.S. mail to the address listed above.

**FOR FURTHER INFORMATION CONTACT:** To request more information on the proposed project or to obtain a copy of the information collection plan and instruments, contact the Information Collection Review Office, Centers for Disease Control and Prevention, 1600 Clifton Road NE., MS-D74, Atlanta, Georgia 30329; phone: 404-639-7570; Email: [omb@cdc.gov](mailto:omb@cdc.gov).

**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. In addition, the PRA also requires Federal agencies to provide a 60-day notice in the **Federal Register**

concerning each proposed collection of information, including each new proposed collection, each proposed extension of existing collection of information, and each reinstatement of previously approved information collection before submitting the collection to OMB for approval. To comply with this requirement, we are publishing this notice of a proposed data collection as described below.

Comments are invited on: (a) Whether the proposed collection of information is necessary for the proper performance of the functions of the agency, including whether the information shall have practical utility; (b) the accuracy of the agency's estimate of the burden of the proposed collection of information; (c) ways to enhance the quality, utility, and clarity of the information to be collected; (d) ways to minimize the burden of the collection of information on respondents, including through the use of automated collection techniques or other forms of information technology; and (e) estimates of capital or start-up costs and costs of operation, maintenance, and purchase of services to provide information. Burden means the total time, effort, or financial resources expended by persons to generate, maintain, retain, disclose or provide information to or for a Federal agency. This includes the time needed to review instructions; to develop, acquire, install and utilize technology and systems for the purpose of collecting, validating and verifying information, processing and maintaining information, and disclosing and providing information; to train personnel and to be able to respond to a collection of information, to search data sources, to complete and review the collection of information; and to transmit or otherwise disclose the information.