Colorado, by appointment. Comments and requests for a copy of the proposed agreement should be addressed to Matt Hogue, Enforcement Specialist, Superfund and Emergency Management Division, Environmental Protection Agency-Region 8, Mail Code 8SEM–PAC, 1595 Wynkoop Street, Denver, Colorado 80202, (303) 312–6591 and should reference the Richardson Flat Tailings Site.

FOR FURTHER INFORMATION CONTACT: Amelia Piggott, Senior Assistant Regional Counsel, Office of Regional Counsel, Environmental Protection Agency-Region 8, Mail Code 8ORC–LEC, 1595 Wynkoop Street, Denver, Colorado 80202, (303) 312–6410.

SUPPLEMENTARY INFORMATION: The proposed Settlement Agreement allows the Owners to make a cash payment: (1) To EPA and the State to resolve alleged civil CERCLA liability; and (2) to DOI and the State to resolve alleged natural resource damage liability. The proposed Settlement Agreement also facilitates the sale of the Property within the Site to Purchaser as a CERCLA Bona Fide Prospective Purchaser and provides for the performance of Work by Purchaser at the Property and for the payment of certain response costs incurred by the United States at or in connection with the Property. The Owners and Purchaser consent to and will not contest the authority of the United States to enter into the Agreement or to implement or enforce its terms. The Owners and Purchaser recognize that the Agreement has been negotiated in good faith and that the Agreement is entered into without the admission or adjudication of any issue of fact or law.

Dated: July 15, 2019.

Betsy Smidinger,

Division Director, Superfund and Emergency Management Division, U.S. Environmental Protection Agency, Region VIII.

[FR Doc. 2019–15852 Filed 7–24–19; 8:45 am]

BILLING CODE 6560–50–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 notices are set forth in paragraph 7 of the Act (12 U.S.C. 1817(j)(7)).

The notices are available for immediate inspection at the Federal Reserve Bank indicated. The notices also will be available for inspection at the offices of the Board of Governors. Interested persons may express their views in writing to the Reserve Bank indicated for that notice or to the offices of the Board of Governors. Comments must be received not later than August 9, 2019.

A. Federal Reserve Bank of Atlanta (Kathryn Haney, Assistant Vice President) 1000 Peachtree Street NE, Atlanta, Georgia 30309. Comments can also be sent electronically to *Applications.Comments@atl.frb.org:*

1. *Jacquelyn Lee Johnson, as co-trustee of the Zachary M. Johnson, Jr. Irrevocable Trust, Woodbine, Georgia; Ms. Jennifer J. Pope, as co-trustee, Macon, Georgia; Mr. Zachary M. Johnson, III, and Mr. Homer Jackson Johnson, as co-trustees, of the Zachary M. Johnson, Jr. Irrevocable Trust, all of Alma, Georgia;* to retain shares of First Bank Shares of the South East, Inc., and thereby indirectly retain shares of its subsidiary, FNB South (formerly known as First National Bank South), both of Alma, Georgia.

B. Federal Reserve Bank of Kansas City (Dennis Denney, Assistant Vice President) 1 Memorial Drive, Kansas City, Missouri 64198–0001:

1. *Mark Brase, Windsor, Colorado, individually and as trustee for the William S. Olson Trust, the Beth Brase Appointment Trust, the Christine Vanderliet Appointment Trust, and the Carla Lehman Appointment Trust, all of Windsor, Colorado;* to retain voting shares of O & F Cattle Company, and thereby indirectly retain shares of Nebraska State Bank, both in Oshkosh, Nebraska.

In addition, Christine Vanderliet, Angels Camp, California; Carla Lehman, Denver, Colorado; and Beth Brase, Windsor, Colorado; to join the Olson Family Group and retain voting shares of O & F Cattle Company.

C. Federal Reserve Bank of Minneapolis (Mark A. Rauzi, Vice President) 90 Hennepin Avenue, Minneapolis, Minnesota 55480–0291:

1. *Ted Gerber and Kelly Gerber, both of Grantsburg, Wisconsin;* to retain shares of Cameron Bancorp, Inc., Cameron, Wisconsin, and thereby indirectly retain shares of Community Bank of Cameron, Cameron, Wisconsin.

Additionally, Mary Gerber, Timothy Gerber, Heather Gerber, Caralyn Duerkop, Justin Duerkop, all of Cameron, Wisconsin; Ernest Tyler Gerber, Menasha, Wisconsin; Nancy Gerber, Exeland, Wisconsin; and Mercedes Gerber, Rice Lake, Wisconsin; to retain shares and be approved as members of the Gerber Family group acting in concert, to retain shares of Cameron Bancorp, Inc., Cameron, Wisconsin.

Board of Governors of the Federal Reserve System, July 22, 2019.

Yao-Chin Chao,

Assistant Secretary of the Board.

[FR Doc. 2019–15835 Filed 7–24–19; 8:45 am]

BILLING CODE 6210–01–P

DEPARTMENT OF HEALTH AND HUMAN SERVICES

Centers for Disease Control and Prevention

[60Day–19–19BIW; Docket No. CDC–2019–0060]

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 effort to reduce public burden and maximize the utility of government information, invites the general public and other Federal agencies the opportunity to comment on a proposed and/or continuing information collection, as required by the Paperwork Reduction Act of 1995. This notice invites comment on a proposed information collection project titled Evaluation of the DP18–1801 Healthy Schools Program. This evaluation will examine three selected DP18–1801 Healthy Schools Program (DP18–1801) grantees to provide a comprehensive picture of implementation activities, context, successes and challenges, key partnerships, lessons learned, and impact on program outcomes.

DATES: CDC must receive written comments on or before September 23, 2019.

ADDRESSES: You may submit comments, identified by Docket No. CDC–2019–0060 by any of the following methods:

- *Federal eRulemaking Portal: Regulations.gov*. Follow the instructions for submitting comments.
- *Mail:* Jeffrey M. Zirger, 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. CDC will post, without change, all relevant comments to *Regulations.gov*.

Please note: Submit all comments 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 Jeffrey M. Zirger, 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.*

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 the OMB for approval. To comply with this requirement, we are publishing this notice of a proposed data collection as described below.

The OMB is particularly interested in comments that will help:

1. Evaluate whether the proposed collection of information is necessary for the proper performance of the functions of the agency, including whether the information will have practical utility;

2. Evaluate the accuracy of the agency's estimate of the burden of the proposed collection of information, including the validity of the methodology and assumptions used;

3. Enhance the quality, utility, and clarity of the information to be collected; and

4. Minimize the burden of the collection of information on those who are to respond, including through the use of appropriate automated, electronic, mechanical, or other technological collection techniques or other forms of information technology, *e.g.,* permitting electronic submissions of responses.

5. Assess information collection costs.

Proposed Project

Evaluation of the DP18–1801 Healthy Schools Program—New—Division of Population Health (DPH) National Center for Chronic Disease Prevention and Health Promotion (NCCDPHP), Centers for Disease Control and Prevention (CDC).

Background and Brief Description

The Centers for Disease Control and Prevention's (CDC) School Health Branch (SHB) requests a three-year OMB approval to conduct a new information collection entitled DP18–1801 Healthy Schools (DP18–1801) Program Evaluation. The DP18–1801 Healthy Schools Program builds upon previous CDC efforts designed to enhance the capacity of state education agencies (SEAs) to adopt and implement evidence-based policies, practices, and programs that support health among the nation's youth. The purpose of the DP18–1801 Healthy Schools Program is to: (1) Increase the number of students who consume nutritious food and beverages (*i.e.,* those aligned with the *Dietary Guidelines for Americans*); (2) increase the number of students who participate in daily physical education and physical activity; and (3) increase the number of students who can effectively manage their chronic health conditions. The evaluation approach is a multisite, embedded case study design, consisting of both process and outcome components, focusing on three 1801 state grantees and a subset of their targeted LEAs and schools. The process component will assess implementation of strategies and activities at the state, local, and school levels and their integration across levels; fidelity of implementation; implementation facilitators and barriers; and contributions of national and state level TA towards program achievements. Three primary data collection methods will be used: (1) Key informant interviews (KII) conducted during in-person site visits or by phone, (2) Web-based surveys, and (3) review of secondary data sources.

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)
SEA staff	Web-Survey	3	1	75/60	4
	Key-Informant Interview	9	1	75/60	11
LEA staff	Web-Survey	30	1	75/60	38
	Key-Informant Interview	12	1	75/60	15
School staff	Web-Survey	210	1	75/60	263
	Key-Informant Interview	54	1	75/60	68
Total					398

Jeffrey M. Zirger,

Lead, Information Collection Review Office, Office of Scientific Integrity, Office of Science, Centers for Disease Control and Prevention.

[FR Doc. 2019–15816 Filed 7–24–19; 8:45 am]

BILLING CODE 4163–18–P

DEPARTMENT OF HEALTH AND HUMAN SERVICES

Centers for Disease Control and Prevention

[60Day–19–19BJD; Docket No. CDC–2019–0059]

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 effort to reduce public burden and maximize the utility of government information, invites the general public and other Federal agencies the opportunity to comment on a proposed and/or continuing information collection, as required by the Paperwork Reduction Act of 1995. This notice invites comment on a proposed information collection project titled ''Monitoring and reporting for the Overdose Data to Action Cooperative Agreement.'' This new data collection effort is to collect information from grantees funded under the Overdose Data to Action (CDC–RFA–CE19–1904) funding opportunity. The information collected will be used to monitor the progress on set performance activities, and progress towards stated grant objectives.

DATES: CDC must receive written comments on or before September 23, 2019.

ADDRESSES: You may submit comments, identified by Docket No. CDC–2010–0059 by any of the following methods:

- *Federal eRulemaking Portal: Regulations.gov*. Follow the instructions for submitting comments.
- *Mail:* Jeffrey M. Zirger, 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. CDC will post, without change, all relevant comments to *Regulations.gov*.

Please note: Submit all comments 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 Jeffrey M. Zirger, 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.*

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 the OMB for approval. To comply with this requirement, we are publishing this notice of a proposed data collection as described below.

The OMB is particularly interested in comments that will help:

1. Evaluate whether the proposed collection of information is necessary for the proper performance of the functions of the agency, including whether the information will have practical utility;

2. Evaluate the accuracy of the agency's estimate of the burden of the proposed collection of information, including the validity of the methodology and assumptions used;

3. Enhance the quality, utility, and clarity of the information to be collected; and

4. Minimize the burden of the collection of information on those who are to respond, including through the use of appropriate automated, electronic, mechanical, or other technological collection techniques or other forms of information technology, *e.g.,* permitting electronic submissions of responses.

5. Assess information collection costs.

Proposed Project

Monitoring and reporting for the Overdose Data to Action Cooperative Agreement—New—National Center for Injury Prevention and Control (NCIPC), Centers for Disease Control and Prevention (CDC).

Background and Brief Description

This new request for a data collection effort is to collect information from grantees funded under the Overdose Data to Action. OMB approval is requested for three years for this new collection. Drug overdose deaths in the United States increased by 18% per year from 2014 to 2016. Opioid overdose deaths have increased fivefold from 1999 to 2016 and in 2017, there were more than 47,000 deaths attributed to opioids. The purpose of the Overdose Data to Action funding opportunity is to support funded grantees in getting high quality, complete, and timely data on opioid prescribing and overdoses, and to use those data to inform prevention and response efforts. There are two required components of this award, a surveillance component, and a prevention component. The intent is to ensure that funded grantees are well equipped to do rigorous work under both components.

The information collected will provide crucial data to CDC for program monitoring and budget tracking, to improve timely CDC-recipient communications, and to inform technical assistance and guidance documents produced by CDC to support program implementation among funded grantees. It will also provide CDC with the capacity to respond in a timely manner to requests for information about the program from the Department of Health and Human Services (HHS), the White House, Congress, and other sources.

Data collection will include 100% of the funded grantee population, with no sampling. The data will be analyzed using descriptive, summary statistics, and qualitative summary. CDC requests approval for 1,320 annualized burden hours. There are no costs to the respondents other than their time.