

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 December 4, 2020.

A. Federal Reserve Bank of Minneapolis (Chris P. Wangen, Assistant Vice President), 90 Hennepin Avenue, Minneapolis, Minnesota 55480-0291:

1. *Security Bancshares Company, Glencoe, Minnesota*; to acquire voting shares of Flagship Financial Group, Inc., Eden Prairie, Minnesota, and thereby indirectly acquire Flagship Bank Minnesota, Wayzata, Minnesota.

Board of Governors of the Federal Reserve System, October 30, 2020.

Michele Taylor Fennell,

Deputy Associate Secretary of the Board.

[FR Doc. 2020-24439 Filed 11-3-20; 8:45 am]

BILLING CODE 6210-01-P

DEPARTMENT OF DEFENSE

GENERAL SERVICES ADMINISTRATION

NATIONAL AERONAUTICS AND SPACE ADMINISTRATION

[OMB Control No. 9000-0001; Docket No. 2020-0053; Sequence No. 14]

Information Collection; Standard Form 28, Affidavit of Individual Surety

AGENCY: Department of Defense (DOD), General Services Administration (GSA), and National Aeronautics and Space Administration (NASA).

ACTION: Notice and request for comments.

SUMMARY: In accordance with section 3506(c)(2)(A) of the Paperwork Reduction Act of 1995, and the Office of Management and Budget (OMB) regulations, DoD, GSA, and NASA invite the public to comment on a revision and renewal concerning the Standard Form 28, Affidavit of Individual Surety. DoD, GSA, and NASA invite comments on: Whether the proposed collection of information is necessary for the proper performance of the functions of Federal Government acquisitions, including whether the information will have practical utility; the accuracy of the estimate of the burden of the proposed information collection; ways to enhance the quality, utility, and clarity of the information to be collected; and ways to minimize the burden of the information collection on respondents, including the use of automated collection techniques or other forms of information technology. OMB has approved this information collection for use through February 28, 2021. DoD, GSA, and NASA propose that OMB extend its approval for use for three additional years beyond the current expiration date.

DATES: DoD, GSA, and NASA will consider all comments received by January 4, 2021.

ADDRESSES: DoD, GSA, and NASA invite interested persons to submit comments on this collection through <http://www.regulations.gov> and follow the instructions on the site. This website provides the ability to type short comments directly into the comment field or attach a file for lengthier comments. If there are difficulties submitting comments, contact the GSA Regulatory Secretariat Division at 202-501-4755 or GSARegSec@gsa.gov.

Instructions: All items submitted must cite OMB Control No. 9000-0001, Standard Form 28, Affidavit of Individual Surety. Comments received generally will be posted without change to <http://www.regulations.gov>, including any personal and/or business confidential information provided. To confirm receipt of your comment(s), please check www.regulations.gov, approximately two-to-three days after submission to verify posting.

FOR FURTHER INFORMATION CONTACT: Zenaida Delgado, Procurement Analyst, at telephone 202-969-7207, or zenaida.delgado@gsa.gov.

SUPPLEMENTARY INFORMATION:

A. OMB Control Number, Title, and Any Associated Form(s)

9000-0001, Standard Form 28, Affidavit of Individual Surety.

B. Need and Uses

This clearance covers the information that offerors or contractors must submit to comply with the following Federal Acquisition Regulation (FAR) requirement:

- *Standard Form (SF) 28, Affidavit of Individual Surety.*

This form is used by all executive agencies, including the Department of Defense (DoD), to obtain information from individuals wishing to serve as sureties to Government bonds. Offerors and contractors may use an individual surety as security for bonds required under a solicitation or contract for supplies or services (including construction). It is an elective decision on the part of the offeror or contractor to use individual sureties instead of other available sources of surety or sureties for Government bonds.

The contracting officer uses the information on the SF 28 to determine the acceptability of individuals proposed as sureties.

C. Annual Burden

Respondents: 10.

Total Annual Responses: 20.

Total Burden Hours: 6.

Obtaining Copies: Requesters may obtain a copy of the information collection documents from the GSA Regulatory Secretariat Division by calling 202-501-4755 or emailing GSARegSec@gsa.gov. Please cite OMB Control No. 9000-0001, Standard Form 28, Affidavit of Individual Surety.

William F. Clark,

Director, Office of Government-wide Acquisition Policy, Office of Acquisition Policy, Office of Government-wide Policy.

[FR Doc. 2020-24417 Filed 11-3-20; 8:45 am]

BILLING CODE 6820-EP-P

DEPARTMENT OF HEALTH AND HUMAN SERVICES

Centers for Disease Control and Prevention

[30Day-21-1215]

Agency Forms Undergoing Paperwork Reduction Act Review

In accordance with the Paperwork Reduction Act of 1995, the Centers for Disease Control and Prevention (CDC) has submitted the information collection request titled Awardee Lead Profile Assessment (ALPA) to the Office

of Management and Budget (OMB) for review and approval. CDC previously published a “Proposed Data Collection Submitted for Public Comment and Recommendations” notice on July 20, 2020 to obtain comments from the public and affected agencies. CDC did not receive comments related to the previous notice. This notice serves to allow an additional 30 days for public and affected agency comments.

CDC will accept all comments for this proposed information collection project. The Office of Management and Budget is particularly interested in comments that:

- (a) 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;
- (b) Evaluate the accuracy of the agencies estimate of the burden of the proposed collection of information, including the validity of the methodology and assumptions used;
- (c) Enhance the quality, utility, and clarity of the information to be collected;
- (d) 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 submission of responses; and
- (e) Assess information collection costs.

To request additional information on the proposed project or to obtain a copy of the information collection plan and instruments, call (404) 639-7570. 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. Direct written comments and/or suggestions regarding the items contained in this notice to the Attention: CDC Desk Officer, Office of

Management and Budget, 725 17th Street NW, Washington, DC 20503 or by fax to (202) 395-5806. Provide written comments within 30 days of notice publication.

Proposed Project

Awardee Lead Profile Assessment (ALPA) (OMB Control No. 0920-1215, Exp. 02/28/2021)—Revision—National Center for Environmental Health (NCEH), Centers for Disease Control and Prevention (CDC).

Background and Brief Description

The Centers for Disease Control and Prevention (CDC) is requesting Paperwork Reduction Act (PRA) clearance for a three-year revised information collection request (ICR) titled “Awardee Lead Profile Assessment (ALPA)” (OMB Control No. 0920-1215; expiration date 02/28/2021). The goal of this ICR is to build on the CDC’s existing childhood lead poisoning prevention program. Based on program successes over the past three years, CDC has made ALPA an annual reporting requirement for ongoing and new CDC Childhood Lead Poisoning Prevention Programs (CLPPPs), including the FY17 “Lead Poisoning Prevention—Childhood Lead Poisoning Prevention—financed partially by Prevention and Public Health Funds” (CDC-RFA-EH17-1701PPHF17); the FY18 “Childhood Lead Poisoning Prevention Projects, State and Local Childhood Lead Poisoning Prevention and Surveillance of Blood Lead Levels in Children” (CDC-RFA-EH18-1806); and the FY20 “Childhood Lead Poisoning Prevention and Surveillance of Blood Lead Levels in Children” (CDC-RFA-EH20-2001). This annual information collection will be used to: (1) identify common characteristics of funded childhood lead poisoning prevention programs, and (2) inform guidance and resource development in support of the ultimate program goal, which is blood lead elimination in children.

The dissemination of these ALPA results will ensure that both funded and non-funded jurisdictions are able to; (1) identify policies and other factors that

support or hinder childhood lead poisoning prevention efforts; (2) understand what strategies are being used by funded public health agencies to implement childhood lead poisoning prevention activities; and (3) use this knowledge to develop and apply similar strategies to support the national agenda to eliminate childhood lead poisoning.

This program management information collection has been revised in several ways. Due to an increase in funding and program growth, CDC is requesting an increase in the number of respondents, defined as state and local governments, or their bona fide agents.

CDC will continue to use two data collection modes, a web survey and an email survey. We anticipate that most of the respondents (n=60; 98 percent) will use the web survey. The estimates of the number and percentage of respondents by mode of data collection are based on previous data collections. In the past, respondents only used the email survey if they had technical difficulties with the web survey, which was rare. For this purpose, we estimate that only 2% (n=1) of the respondents may need to submit an email survey. This represents a change in distribution from the 2018 estimates, which were initially assumed as 83.3% for the web survey and 16.7% for the email survey.

A redistribution by mode of collection will not affect the total time burden requested as the time per response is the same for either mode; however, the time to take the survey has increased from seven minutes in 2018 to 47 minutes per response due to a revision of the survey. This revised time estimate per response is based on pilot tests of the revised survey among nine respondents, and includes the time needed to review the ALPA Training Manual, which is a new addition in this revision ICR.

Thus, CDC is requesting an increase in the annual number of respondents from 48 to a maximum of 61 recipients (n=13 more respondents), and an increase in the total annual time burden from six hours in 2018 to 48 hours (n=42 more hours).

ESTIMATED ANNUALIZED BURDEN HOURS

Type of respondents	Form name	Number of respondents	Number of responses per respondent	Average burden per response (in hours)
State or Local Governments (or their bona fide fiscal agents).	ALPA Web Survey	60	1	47/60
	ALPA Email Survey	1	1	47/60

Jeffrey M. Zirger,

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

[FR Doc. 2020-24474 Filed 11-3-20; 8:45 am]

BILLING CODE 4163-18-P

DEPARTMENT OF HEALTH AND HUMAN SERVICES

Centers for Disease Control and Prevention

Framework for Conditional Sailing and Initial Phase COVID-19 Testing Requirements for Protection of Crew

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

ACTION: Notice.

SUMMARY: The Centers for Disease Control and Prevention (CDC), a component of the Department of Health and Human Services (HHS), announces a framework for a phased resumption of cruise ship passenger operations. CDC also announces requirements for the initial phases of this framework regarding testing of crew members for COVID-19, an integral part of the initial phases prior to resuming passenger operations. This Order applies to cruise ship operators with cruise ships operating in U.S. waters and cruise ship operators who are operating cruise ships outside of U.S. waters, but intend for their cruise ships to return to operating in U.S. waters while this Order remains in effect.

DATES: This action is effective October 30, 2020.

FOR FURTHER INFORMATION CONTACT: Jennifer Buigut, Division of Global Migration and Quarantine, Centers for Disease Control and Prevention, 1600 Clifton Road NE, MS H16-4, Atlanta, GA 30329. Phone: 404-498-1600. Email: dgmqpolicyoffice@cdc.gov.

SUPPLEMENTARY INFORMATION: This Order establishes a framework for a phased approach to resuming cruise ship passenger operations in U.S. waters. This phased approach will include: (1) Establishment of laboratory testing of crew onboard cruise ships in U.S. waters; (2) simulated voyages designed to test a cruise ship operators' ability to mitigate COVID-19 onboard cruise ships; (3) a certification process; and (4) a return to passenger voyages in a manner that mitigates the risk of COVID-19 introduction, transmission, or spread among passengers and crew onboard ships and ashore to communities.

As part of the initial crew testing phases, this Order additionally contains requirements for: (1) Shoreside COVID-19 laboratory screening testing of all crew currently onboard cruise ships; (2) onboard diagnostic testing capabilities for symptomatic travelers (crew and future passengers); (3) shoreside COVID-19 laboratory screening testing of all newly embarking crew; and (4) continued compliance with complete, accurate, and acknowledged, No Sail Order Response Plans.

A copy of the Order is provided below and a copy of the signed order can be found at <https://www.cdc.gov/quarantine/cruise/index.html>.

U.S. Department of Health and Human Services (HHS)—Centers for Disease Control and Prevention (CDC)

Order Under Sections 361 & 365 of the Public Health Service Act (42 U.S.C. 264, 268) and 42 Code of Federal Regulations Part 70 (Interstate) and Part 71 (Foreign): Framework for Conditional Sailing and Initial Phase COVID-19 Testing Requirements for Protection of Crew

Executive Summary

The Centers for Disease Control and Prevention (CDC), a component of the U.S. Department of Health and Human Services (HHS), announces this framework for a phased resumption of cruise ship passenger operations. Considering the continued spread of COVID-19 worldwide and increased risk of COVID-19 on cruise ships, a careful approach is needed to safely resume cruise ship passenger operations. CDC is establishing requirements to mitigate the COVID-19 risk to passengers and crew, prevent the further spread of COVID-19 from cruise ships into U.S. communities, and protect public health and safety. After expiration of CDC's No Sail Order (NSO) on October 31, 2020, CDC will take a phased approach to resuming cruise ship passenger operations in U.S. waters.

The initial phases will consist of testing and additional safeguards for crew members. CDC will ensure cruise ship operators have adequate health and safety protections for crew members while these cruise ship operators build the laboratory capacity needed to test future passengers. Subsequent phases will include simulated voyages to test cruise ship operators' ability to mitigate COVID-19 risk, certification for ships that meet specific requirements, and a phased return to cruise ship passenger voyages in a manner that mitigates COVID-19 risk among passengers, crew members, and U.S. communities. These

phases are subject to change based on public health considerations and cruise ship operators' demonstrated ability to mitigate COVID-19 risk. CDC will issue additional orders as needed that will be published in the **Federal Register** and technical instructions that will be subsequently posted on CDC's website.

This Order additionally announces requirements for the initial phases relating to crew testing. CDC considers adequate crew safeguards as demonstrated through laboratory testing for SARS coronavirus 2 (SARS-CoV-2), the virus that causes COVID-19, an integral part of the initial phases prior to resuming passenger operations.

Previous Orders and Incorporation by Reference

The findings and other evidence relied upon in issuing the No Sail Order and Other Measures Related to Operations signed by the CDC Director on March 14, 2020,¹ as further modified and extended effective April 15, 2020,² July 16, 2020,³ and September 30, 2020⁴—are incorporated herein by reference.

Statement of Intent

This Order shall be interpreted and implemented in a manner as to achieve the following paramount objectives:

- Preserving human life;
- Preserving the health and safety of cruise ship crew members, port personnel, and communities;
- Preventing the further introduction, transmission, and spread of COVID-19 into and throughout the United States;
- Preserving the public health and other critical resources of Federal, State, and local governments;

¹ No Sail Order and Suspension of Further Embarkation. <https://www.federalregister.gov/documents/2020/03/24/2020-06166/no-sail-order-and-suspension-of-further-embarkation>. Last accessed October 19, 2020.

² No Sail Order and Suspension of Further Embarkation; Notice of Modification and Extension and Other Measures Related to Operations. <https://www.federalregister.gov/documents/2020/04/15/2020-07930/no-sail-order-and-suspension-of-further-embarkation-notice-of-modification-and-extension-and-other>. Last accessed October 19, 2020.

³ No Sail Order and Suspension of Further Embarkation, Second Modification and Extension of No Sail Order and Other Measures Related to Operations. <https://www.federalregister.gov/documents/2020/07/21/2020-15810/no-sail-order-and-suspension-of-further-embarkation-second-modification-and-extension-of-no-sail>. Last accessed October 19, 2020.

⁴ No Sail Order and Suspension of Further Embarkation; Third Modification and Extension of No Sail Order and Other Measures Related to Operations. <https://www.federalregister.gov/documents/2020/10/05/2020-22030/no-sail-order-and-suspension-of-further-embarkation-third-modification-and-extension-of-no-sail>. Last accessed October 19, 2020.