FOR FURTHER INFORMATION CONTACT: For additional information about the information collection, contact Cathy Williams at (202) 418–2918.

SUPPLEMENTARY INFORMATION:

OMB Control No.: 3060–1063.

Title: Global Mobile Personal Communications by Satellite (GMPCS) Authorization, Marketing and Importation Rules.

Form No.: Not Applicable.

Type of Review: Revision of a currently approved information collection.

Respondents: Business or other for-profit entities.

Number of Respondents: 17 respondents; 17 responses.

Estimated Time per Response: 1–24 hours per response.

Frequency of Response: On occasion reporting requirement.

Obligation To Respond: Required to obtain or retain benefits. The Commission has authority for this information collection pursuant to Sections 4(i), 301, 302(a), 303(e), 303(f), 303(g), 303(n) and 303(r) of the Communications Act of 1934, as amended; 47 U.S.C. 4(i), 301, 302(a), 303(e), 303(f), 303(g), 303(n) and 303(r).

Total Annual Burden: 595 hours.

Annual Cost Burden: None.

Privacy Act Impact Assessment: No impact(s).

Nature and Extent of Confidentiality: In general, there is no need for confidentiality with this collection of information.

Needs and Uses: On July 14, 2017, the Federal Communications Commission (''Commission'') released a First Report and Order titled, ''In the Matter of Amendment of Parts 0, 1, 2, 15 and 18 of the Commission's Rules Regarding Authorization of Radiofrequency Equipment,'' ET Docket No. 15–170 (FCC 17–93). In the First Report and Order, the Commission discontinued use of the ''Statement Regarding the Importation of Radio Frequency Devices Capable of Harmful Interference,'' (FCC Form 740) and eliminated 47 CFR 2.1205 and 2.1203(b), thus removing the Form 740 filing requirements. The agency concluded that there was no evidence indicating that the Form 740 filing process provided a substantial deterrent to illegal importation of RF devices, and that the existing filing requirement creates large burdens in light of the growth in the number and type of RF devices being imported, and that there is now a wider availability of product and manufacturer information, including that available to the FCC from the Custom and Border Protection's (CBP) database. The Form 740 was approved under OMB Control No. 3060–0059 and was under the purview of the Commission's Office of Engineering & Technology (OET).

The purposes of the revision of OMB Control No. 3060–1063 are to reflect a slight decrease in the number of satellite operators and/or GMPCS equipment manufacturers and changes resulting from the elimination of Form 740. Specifically, the number of respondents changed from 19 to 17 due to a decrease in the number of satellite operators and/or GMPCS equipment manufacturers. As a result of the elimination of the Form 740, the total annual burden hours changed from 684 to 595 and the total annual costs decreased from $13,110 to zero.

The purpose of this information collection is to maintain OMB approval of a certification requirement for portable GMPCS transceivers to prevent interference, reduce radio-frequency (''RF'') radiation exposure risk, and make regulatory treatment of portable GMPCS transceivers consistent with treatment of similar terrestrial wireless devices, such as cellular phones.

The Commission is requiring that applicants obtain authorization for the equipment by submitting an application and exhibits, including test data. If the Commission did not obtain such information, it would not be able to ascertain whether the equipment meets the FCC's technical standards for operation in the United States. Furthermore, the data is required to ensure that the equipment will not cause catastrophic interference to other telecommunications services that may impact the health and safety of American citizens.

Federal Communications Commission.

Marlene Dortch,

Secretary, Office of the Secretary.

[FR Doc. 2018–08377 Filed 4–20–18; 8:45 am]

BILLING CODE 6712–01–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 (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 May 10, 2018.

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:*

1. *Anna E. Hechler, Quincy, Illinois, individually and as part of a family control group that includes Joseph E. Gully, Barry, Illinois;* to retain shares of FNB Barry Bancorp, Inc., Barry, Illinois, and thereby retain shares of First National Bank of Barry, Barry, Illinois.

Board of Governors of the Federal Reserve System, April 17, 2018.

Ann Misback,

Secretary of the Board.

[FR Doc. 2018–08349 Filed 4–20–18; 8:45 am]

BILLING CODE 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 May 8, 2018.

A. Federal Reserve Bank of Boston (Prabal Chakrabarti, Senior Vice President) 600 Atlantic Avenue, Boston, Massachusetts 02210–2204. Comments can also be sent electronically to *BOS.SRC.Applications.Comments@bos.frb.org:*

1. *HarborOne Mutual Bancshares and its mid-tier stock holding company, HarborOne Bancorp, Inc., both of Brockton, Massachusetts;* to merge with Coastway Bancorp, Inc., and thereby indirectly acquire Coastway Community Bank, both of Warwick, Rhode Island.

Board of Governors of the Federal Reserve System, April 18, 2018.

Ann Misback,

Secretary of the Board.

[FR Doc. 2018–08384 Filed 4–20–18; 8:45 am]

BILLING CODE P

DEPARTMENT OF HEALTH AND HUMAN SERVICES

Centers for Disease Control and Prevention

[60Day–FY–0696; Docket No. CDC–2018–0035]

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 *National HIV Prevention Program Monitoring and Evaluation (NHM&E),* which collects standardized HIV prevention program evaluation data from health departments and community-based organizations (CBOs) who receive federal funds for HIV prevention activities.

DATES: CDC must receive written comments on or before June 22, 2018.

ADDRESSES: You may submit comments, identified by Docket No. CDC–2018–0035 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. 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 Leroy A. Richardson, 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

National HIV Prevention Program Monitoring and Evaluation (NHM&E) (OMB Control Number 0920–0696, Expiration Date 02/28/2019)—Revision—National Center for HIV/AIDS, Viral Hepatitis, STD, and TB Prevention (NCHHSTP), Centers for Disease Control and Prevention (CDC).

Background and Brief Description

CDC seeks to request a three-year Office of Management and Budget (OMB) approval to revise the previously approved project and continue the collection of standardized HIV prevention program evaluation data from health departments and community-based organizations (CBOs) who receive federal funds for HIV prevention activities. Health department grantees have the options to key-enter or upload data to a CDC-provided web-based software application (EvaluationWeb®). CBO grantees may only key-enter data to the CDC-provided web-based software application.

This revision includes changes to the data variables to adjust to the different monitoring and evaluation needs of new funding announcements without a substantial change in burden.

The evaluation and reporting process is necessary to ensure that CDC receives standardized, accurate, thorough evaluation data from both health department and CBO grantees. For these reasons, CDC developed standardized NHM&E variables through extensive consultation with representatives from health departments, CBOs, and national partners (*e.g.,* The National Alliance of State and Territorial AIDS Directors and Urban Coalition of HIV/AIDS Prevention Services).

CDC requires CBOs and health departments who receive federal funds for HIV prevention to report non-identifying, client-level and aggregate level, standardized evaluation data to: (1) Accurately determine the extent to which HIV prevention efforts are carried out, what types of agencies are providing services, what resources are allocated to those services, to whom services are being provided, and how these efforts have contributed to a reduction in HIV transmission; (2) improve ease of reporting to better meet these data needs; and (3) be accountable to stakeholders by informing them of