as stated in the IUP and grant agreement. The CWA and SDWA require this information to ensure the national accountability, adequate public review and comment, fiscal integrity, and consistent management needed to achieve public health and CWA and SDWA compliance objectives.

Additional information about the CWSRFs and DWSRFs are available at *http://www.epa.gov/cwsrf/learn-about-clean-water-state-revolving-fund-cwsrf* and *https://www.epa.gov/dwsrf/how-drinking-water-state-revolving-fund-works#tab-1,* respectively.

This ICR renews the Office of Management and Budget (OMB) Number 2040–0185 DWSRF ICR and provides updated estimates of the reporting burden associated with the information collection activities for both DWSRF ICR and CWSRF ICR.

Form numbers: None.

Respondents/affected entities: Entities affected by this action are states and local governments.

Respondent's obligation to respond: Required to obtain or retain a benefit per the Clean Water Act Title VI and the Safe Drinking Water Act Section 1452.

Estimated number of respondents: 2,836 (total).

Frequency of response: Varies by requirement (*i.e.,* quarterly, semi-annually, annually).

Total estimated burden: 108,519 hours (per year). Burden is defined at 5 CFR 1320.03(b)

Total estimated cost: $12,874,081 (per year), which includes $6,354,600 annualized capital or operation & maintenance costs.

Changes in the estimates: There is a decrease of 37,500 hours in the total estimated respondent burden compared with the two separate ICRs currently approved by OMB. This change in hourly burden is primarily due to a decrease in annual hourly burden in the DWSRF from the elimination of the application review estimates to align with the estimates from CWSRF.

Courtney Kerwin,

Director, Regulatory Support Division.

[FR Doc. 2023–11008 Filed 5–23–23; 8:45 am]

BILLING CODE 6560–50–P

FEDERAL MARITIME COMMISSION

[Docket No. FMC–2023–0012]

Agency Information Collection Activities: 60-Day Public Comment Request

AGENCY: Federal Maritime Commission.

ACTION: Sixty-day notice; request for comments.

SUMMARY: As part of our continuing effort to reduce paperwork and respondent burden, and as required by the Paperwork Reduction Act of 1995, the Federal Maritime Commission (Commission) invites comments on a new data collection concerning empty container volumes at intermodal locations. The collection also implements certain provisions of the Ocean Shipping Reform Act of 2022.

DATES: Written comments must be submitted on or before July 24, 2023.

ADDRESSES: Submit comments for the proposed information collection requests to Lucille L. Marvin, Managing Director at email: *omd@fmc.gov.* The FMC will summarize any comments received in response to this notice in a subsequent notice and include them in its information collection submission to OMB for approval.

FOR FURTHER INFORMATION CONTACT: Copies of the information collections and instructions, or copies of any comments received, may be obtained by contacting Tara Nielsen at 202–523–5800 or *omd@fmc.gov.*

SUPPLEMENTARY INFORMATION:

Request for Comments

The Commission, as part of its continuing effort to reduce paperwork and respondent burden, invites the general public and other Federal agencies to comment on the continuing information collections listed in this notice, as required by the Paperwork Reduction Act of 1995 (44 U.S.C. 3501 *et seq.*).

Comments submitted in response to this notice will be included or summarized in our request for Office of Management and Budget (OMB) approval of the relevant information collection. All comments are part of the public record and subject to disclosure. Please do not include any confidential or inappropriate material in your comments. We invite comments on: (1) the necessity and utility of the proposed information collection for the proper performance of the agency's functions; (2) the accuracy of the estimated burden; (3) ways to enhance the quality, utility, and clarity of the information to be collected; and (4) the use of automated collection techniques or other forms of information technology to minimize the information collection burden.

Information Collections Open for Comment

Title: Empty Containers Ready for Use by Location.

OMB Approval Number: 3072–XXXX.

Abstract: The Ocean Shipping Reform Act of 2022 (OSRA 2022) mandated new data collections for Federal agencies to address key gaps in available data. *See* 46 U.S.C. 41110. The Federal Maritime Commission (FMC) was instructed to collect and report on vessel-level tonnage as well as full and empty containers entering and leaving U.S. ports in international trade. The U.S. Department of Transportation's Bureau of Transportation Statistics (BTS) was instructed to collect operational data on intermodal equipment and dwell times. Both agencies have work underway on these data collections. The FMC and BTS have further identified data gaps related to location of intermodal shipping containers (as defined in ISO 668—Series 1 Freight Containers) ready for use by shippers, particularly exporters, at key intermodal locations. Exporters would benefit from current information on the geographic areas where empty ocean intermodal equipment is positioned. FMC is seeking to collect daily data on empty containers on a weekly basis as well as a two-week outlook for containers ready for use at 30 intermodal locations, which will include inland dry ports, intermodal container transfer facilities, and marine terminals.

The information collected would be used to compile and publish a weekly report on the throughput and availability of multiple container types (*e.g.,* refrigerated and dry) and sizes (*e.g.,* 20-foot, 40-foot, and 45-foot) at key intermodal locations. The universe of respondents is the top 12 ocean common carriers, measured by container volume.

Current Actions: This information being submitted contains a new data collection.

Type of Review: New data collection.

Needs and Uses: The Federal Maritime Commission and Bureau of Transportation Statistics will use collected data to produce a weekly report and to monitor current industry trends.

Frequency: This information will be collected weekly each year. There are 52 weeks in a year.

Type of Respondents: The universe will be the 12 largest ocean common carriers, measured by the total volume of containers moving through U.S. ports in international common carriage.

Number of Annual Respondents: The FMC estimates an annual respondent universe of 12 ocean common carriers, each of which will provide data weekly. The total number of responses will be 624. The FMC expects the estimated number of annual respondents to remain constant in the future.

Estimated Time per Response: The time per response is estimated at 2.5 person-hours for reporting requirements.

Total Annual Burden: For the 12 carriers with weekly reporting, the burden is calculated as $12 \times 52 \times 2.5$ hours = 1,560 hours.

William Cody,

Secretary.

[FR Doc. 2023–11082 Filed 5–23–23; 8:45 am]

BILLING CODE 6730–02–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 June 23, 2023.

A. Federal Reserve Bank of San Francisco: (Joseph Cuenco, Assistant Vice President, Formations & Transactions) 101 Market Street, San Francisco, California 94105–1579. Comments can also be sent electronically to: *sf.fisc.comments.applications@sf.frb.org.*

1. *VB&T Holding Company, LLC;* to become a bank holding company through the formation of its subsidiary bank, Zenith Bank & Trust, both of Scottsdale, Arizona.

Board of Governors of the Federal Reserve System.

Ann E. Misback,

Secretary of the Board.

[FR Doc. 2023–11068 Filed 5–23–23; 8:45 am]

BILLING CODE P

DEPARTMENT OF HEALTH AND HUMAN SERVICES

Food and Drug Administration

[Docket Nos. FDA–2020–E–2361; FDA–2020–E–2362; and FDA–2020–E–2363]

Determination of Regulatory Review Period for Purposes of Patent Extension; ENSPRYNG; Correction

AGENCY: Food and Drug Administration, HHS.

ACTION: Notice; correction.

SUMMARY: The Food and Drug Administration (FDA or the Agency) is correcting a notice that appeared in the **Federal Register** of July 13, 2022. The document determined the regulatory review period for ENSPRYNG. After review of a timely request for reconsideration by the applicant of the calculation of the applicable regulatory review period of the biologic product ENSPRYNG in that notice, FDA has determined that a revision of the supplementary information section is warranted. This notice corrects the applicable regulatory review period language.

FOR FURTHER INFORMATION CONTACT: Beverly Friedman, Office of Regulatory Policy, Food and Drug Administration, 10903 New Hampshire Ave., Bldg. 51, Rm. 6250, Silver Spring, MD 20993, 301–796–3600.

SUPPLEMENTARY INFORMATION: In the **Federal Register** of July 13, 2022 (87 FR 41724), on page 41725, in the second column, under ''II. Determination of Regulatory Review Period,'' the first two sentences should be corrected to read as follows: ''FDA has determined that the applicable regulatory review period for ENSPRYNG is 2,494 days. Of this time, 2,128 days occurred during the testing phase of the regulatory review period, while 366 days occurred during the approval phase.''

Dated: May 18, 2023.

Lauren K. Roth,

Associate Commissioner for Policy.

[FR Doc. 2023–10979 Filed 5–23–23; 8:45 am]

BILLING CODE 4164–01–P

DEPARTMENT OF HEALTH AND HUMAN SERVICES

Health Information Technology Advisory Committee 2023 Schedule of Meetings

AGENCY: Office of the National Coordinator for Health Information Technology (ONC), HHS.

ACTION: Notice of meetings.

SUMMARY: The Health Information Technology Advisory Committee (HITAC) was established in accordance with the 21st Century Cures Act and the Federal Advisory Committee Act. The HITAC, among other things, identifies priorities for standards adoption and makes recommendations to the National Coordinator for Health Information Technology (National Coordinator). The HITAC will hold public meetings throughout 2023. See list of public meetings below.

FOR FURTHER INFORMATION CONTACT: Michael Berry, Designated Federal Officer, at *Michael.Berry@hhs.gov,* (202) 701–0795.

SUPPLEMENTARY INFORMATION: Section 4003(e) of the 21st Century Cures Act (Pub. L. 114–255) establishes the Health Information Technology Advisory Committee (referred to as the ''HITAC''). The HITAC will be governed by the provisions of the Federal Advisory Committee Act (FACA) (Pub. L. 92–463), as amended, (5 U.S.C. app.), which sets forth standards for the formation and use of federal advisory committees.

Composition

The HITAC is comprised of at least 25 members, of which:

- No fewer than 2 members are advocates for patients or consumers of health information technology;
- 3 members are appointed by the HHS Secretary:
  - 1 of whom shall be appointed to represent the Department of Health and Human Services and
  - 1 of whom shall be a public health official;
- 2 members are appointed by the majority leader of the Senate;
- 2 members are appointed by the minority leader of the Senate;
- 2 members are appointed by the Speaker of the House of Representatives;
- 2 members are appointed by the minority leader of the House of Representatives; and
- Other members are appointed by the Comptroller General of the United States.

Members serve for one-, two-, or three-year terms. All members may be reappointed for a subsequent three-year