corresponding changes relating to the services in the expanded scope.

Agreement No.: 012258.

Title: The G6/HSDG Atlantic Space Charter Agreement.

Parties: American President Lines, Ltd. and APL Co. Pte, Ltd. (operating as one party); Hapag-Lloyd AG and Hapag-Lloyd USA LLC (operating as one party); Hyundai Merchant Marine Co., Ltd.; Mitsui O.S.K. Lines, Ltd.; Nippon Yusen Kaisha; Orient Overseas Container Line, Limited; and OOCL (Europe) Limited (all operating as one party); and Hamburg Sud.

Filing Party: David F. Smith, Esq.; Cozen O'Connor; 1627 I Street, NW. Suite 1100; Washington, DC 20006.

Synopsis: The agreement would authorize the G6 lines to charter space to Hamburg Sud in the trade between North Europe and the United Kingdom, on the one hand, and the U.S. Atlantic Coast, on the other hand. The agreement would also authorize the parties to enter into arrangements related to the chartering of such space.

Agreement No.: 012059.

Title: COSCON/ELJSA Vessel Sharing Agreement.

Parties: COSCO Container Lines Company, Limited and Evergreen Line Joint Service Agreement.

Filing Party: Eric. C. Jeffrey, Esq.; Nixon Peabody LLP; 401 9th Street NW., Suite 900; Washington, DC 20004.

Synopsis: The agreement authorizes the parties to charter space on each other's vessels, coordinate sailings, and otherwise cooperate in the carriage of cargo in the trade between China (including Hong Kong), Taiwan, Singapore, Japan, Korea, Vietnam, and Canada, on the one hand, and the U.S. West Coast, on the other hand.

By Order of the Federal Maritime Commission.

Dated: April 4, 2014.

Rachel E. Dickon,

Assistant Secretary.

[FR Doc. 2014–07953 Filed 4–8–14; 8:45 am]

BILLING CODE 6730–01–P

GENERAL SERVICES ADMINISTRATION

[Notice–PBS–2014–02; Docket 2014–0002; Sequence 15]

Announcement of the Publication of GSA PBS–P100, Facilities Standards for the Public Buildings Service

AGENCY: Office of Design and Construction Office of Public Buildings Service, General Services Administration.

ACTION: Notice of Publication of the GSA PBS–P100, Facilities Standards for the Public Buildings Service.

SUMMARY: On March 12, 2014, the PBS–P100, Facilities Standards for the Public Buildings Service (P100), was issued. The Facilities Standards for the Public Buildings Service establishes design standards and criteria for new buildings, repairs and alterations, modernizations, lease construction buildings with government option to purchase, and work in historic structures for the Public Buildings Service (PBS) of the U.S. General Services Administration (GSA). This lastest update to the document contains both performance based standards and prescriptive requirements to be used in the programming, design, and documentation of GSA buildings. The PBS–P100 2014 version is available at *http://www.gsa.gov/portal/category/106319* and at *http://gsap100.wbdg.org.*

DATES: April 9, 2014.

FOR FURTHER INFORMATION CONTACT: Mr. Martin Weiland, 202–219–0634, U.S. General Services Administration, 1800 F Street NW., Suite 5400, Washington, DC 20405, or via email *pbsp100@gsa.gov.*

Dated: April 3, 2014.

David Insinga,

Acting Assistant Commissioner, Office of Design and Construction, General Services Administration.

[FR Doc. 2014–07901 Filed 4–8–14; 8:45 am]

BILLING CODE 6820–23–P

DEPARTMENT OF HEALTH AND HUMAN SERVICES

Designation of a Class of Employees for Addition to the Special Exposure Cohort

AGENCY: National Institute for Occupational Safety and Health (NIOSH), Centers for Disease Control and Prevention, Department of Health and Human Services (HHS).

ACTION: Notice.

SUMMARY: HHS gives notice of a decision to designate a class of employees from the Joslyn Manufacturing and Supply Company in Fort Wayne, Indiana as an addition to the Special Exposure Cohort (SEC) under the Energy Employees Occupational Illness Compensation Program Act of 2000. On March 27, 2014, the Secretary of HHS designated the following class of employees as an addition to the SEC:

All Atomic Weapons Employees who worked for Joslyn Manufacturing and Supply Co. at the covered facility in Fort Wayne, Indiana, from March 1, 1943, through July 31, 1948, for a number of work days aggregating at least 250 work days, occurring either solely under this employment, or in combination with work days within the parameters established for one or more other classes of employees included in the Special Exposure Cohort.

This designation will become effective on April 26, 2014, unless Congress provides otherwise prior to the effective date. After this effective date, HHS will publish a notice in the **Federal Register** reporting the addition of this class to the SEC or the result of any provision by Congress regarding the decision by HHS to add the class to the SEC.

FOR FURTHER INFORMATION CONTACT: Stuart L. Hinnefeld, Director, Division of Compensation Analysis and Support, NIOSH, 4676 Columbia Parkway, MS C–46, Cincinnati, OH 45226, Telephone 1–877–222–7570. Information requests can also be submitted by email to *DCAS@CDC.GOV*.

John Howard,

Director, National Institute for Occupational Safety and Health.

[FR Doc. 2014–07935 Filed 4–8–14; 8:45 am]

BILLING CODE 4163–19–P

DEPARTMENT OF HEALTH AND HUMAN SERVICES

Meeting of the Advisory Group on Prevention, Health Promotion, and Integrative and Public Health

AGENCY: Office of the Surgeon General of the United States Public Health Service, Office of the Assistant Secretary for Health, Office of the Secretary, Department of Health and Human Services.

ACTION: Notice.

SUMMARY: In accordance with Section 10(a) of the Federal Advisory Committee Act, Public Law 92–463, as amended (5 U.S.C. App.), notice is hereby given that a meeting is scheduled to be held for the Advisory Group on Prevention, Health Promotion, and Integrative and Public Health (the ''Advisory Group''). The meeting will be open to the public. Information about the Advisory Group and the agenda for this meeting can be obtained by accessing the following Web site: *http://www.surgeongeneral.gov/initiatives/prevention/advisorygrp/index.html.*

DATES: The meeting will be held on April 28 and 29, 2014. Exact start and end times will be published closer to the meeting date at: *http://www.surgeongeneral.gov/initiatives/prevention/advisorygrp/index.html.*

ADDRESSES: The meeting will be held at 200 Independence Ave. SW., Room 505A, Washington, DC 20201 on April 28, 2014. The meeting will take place via teleconference on April 29, 2014.

FOR FURTHER INFORMATION CONTACT: Office of the Surgeon General, 200 Independence Ave. SW., Washington, DC 20201; 202–205–9517; *prevention.council@hhs.gov.*

SUPPLEMENTARY INFORMATION: The Advisory Group is a non-discretionary federal advisory committee that was initially established under Executive Order 13544, dated June 10, 2010, to comply with the statutes under Section 4001 of the Patient Protection and Affordable Care Act, Public Law 111–148. The Advisory Group was established to assist in carrying out the mission of the National Prevention, Health Promotion, and Public Health Council (the Council). The Advisory Group provides recommendations and advice to the Council.

The Advisory Group was terminated on September 30, 2012, by Executive Order 13591, dated November 23, 2011. Authority for the Advisory Group to be re-established was given under Executive Order 13631, dated December 7, 2012. Authority for the Advisory Group to continue to operate until September 30, 2015 was given under Executive Order 13652, dated September 30, 2013.

It is authorized for the Advisory Group to consist of not more than 25 non-federal members. The Advisory Group currently has 22 members who were appointed by the President. The membership includes a diverse group of licensed health professionals, including integrative health practitioners who have expertise in (1) worksite health promotion; (2) community services, including community health centers; (3) preventive medicine; (4) health coaching; (5) public health education; (6) geriatrics; and (7) rehabilitation medicine.

During this meeting, the Advisory Group will have round table discussions with representatives of Council member departments and develop recommendations for the Council for the upcoming year.

Members of the public who wish to attend the meeting on April 28, 2014 or to participate by phone in the April 29, 2014 meeting must register by 12:00 p.m. EST on April 21, 2014. Individuals should register for public attendance at *prevention.council@hhs.gov* by providing your full name and affiliation. Individuals who plan to attend the meeting and need special assistance and/or accommodations, i.e., sign language interpretation or other reasonable accommodations, should indicate so when they register. The public will have the opportunity to provide comments to the Advisory Group on April 28, 2014; public comment will be limited to 3 minutes per speaker. Registration via email (*prevention.council@hhs.gov*) is also required for the public comment session. Any member of the public who wishes to have printed materials distributed to the Advisory Group for this scheduled meeting should submit material to *prevention.council@hhs.gov* no later than 12:00 p.m. EST on April 21, 2014.

Dated: March 25, 2014.

Corinne M. Graffunder,

Designated Federal Officer, Advisory Group on Prevention, Health Promotion, and Integrative and Public Health, Office of the Surgeon General.

[FR Doc. 2014–07848 Filed 4–8–14; 8:45 am]

BILLING CODE 4163–18–P

DEPARTMENT OF HEALTH AND HUMAN SERVICES

Food and Drug Administration

[Docket No. FDA–2010–D–0194]

Agency Information Collection Activities; Submission for Office of Management and Budget Review; Comment Request; Draft Guidance for Industry and FDA Staff; Total Product Life Cycle: Infusion Pump—Premarket Notification Submissions

AGENCY: Food and Drug Administration, HHS.

ACTION: Notice.

SUMMARY: The Food and Drug Administration (FDA) is announcing that a proposed collection of information has been submitted to the Office of Management and Budget (OMB) for review and clearance under the Paperwork Reduction Act of 1995.

DATES: Fax written comments on the collection of information by May 9, 2014.

ADDRESSES: To ensure that comments on the information collection are received, OMB recommends that written comments be faxed to the Office of Information and Regulatory Affairs, OMB, Attn: FDA Desk Officer, FAX: 202–395–7285, or emailed to *oira_submission@omb.eop.gov.* All comments should be identified with the OMB control number 0910–NEW and the title ''Draft Guidance for Industry and FDA Staff; Total Product Life Cycle: Infusion Pump—Premarket Notification [510(k)] Submissions.'' Also include the FDA docket number found in brackets in the heading of this document.

FOR FURTHER INFORMATION CONTACT: FDA PRA Staff, Office of Operations, Food and Drug Administration, 1350 Piccard Dr., PI50–400B, Rockville, MD 20850, *PRAStaff@fda.hhs.gov.*

SUPPLEMENTARY INFORMATION: In compliance with 44 U.S.C. 3507, FDA has submitted the following proposed collection of information to OMB for review and clearance.

Draft Guidance for Industry and FDA Staff; Total Product Life Cycle: Infusion Pump—Premarket Notification [510(k)] Submissions—(OMB Control Number 0910–NEW)

This draft guidance is intended to assist industry in preparing premarket notification submissions for infusion pumps and to identify device features that manufactures should address throughout the total product life cycle. The draft guidance is available at (*http://www.fda.gov/MedicalDevices/DeviceRegulationandGuidance/GuidanceDocuments/ucm206153.htm*).

In the **Federal Register** of April 26, 2010 (75 FR 21632), FDA published a notice seeking comment on the proposed information collection activity. Given the lapse in time since its publication, FDA is reissuing this notice, responding to a single comment and providing the public an additional opportunity to comment on this proposed information collection activity, prior to the issuance of the final guidance document.

In the April 26, 2010, notice, FDA estimated it will receive 31 infusion pump submissions annually. The Agency reached this estimate by averaging the number of premarket notifications for infusion pumps submitted to FDA over the past 5 years. The draft guidance identifies 56 potential hazards FDA recommends addressing if applicable to a particular device. Although there may be additional hazards identified by a manufacturer, the Agency believes these hazards may offset FDA identified hazards not applicable to a particular device. FDA estimates it will take infusion pump manufactures approximately 56 hours (approximately 1 hour per hazard) to complete the case assurance report described in section 6 of the draft guidance. FDA reached this estimate based on its expectation of the amount of information that will be contained in the report.

However, based on a single public comment provided to FDA, related to the FDA burden estimate, we are