comment and reply comment deadlines by 60 days is in the public interest. Extending the comment period will ensure that parties have sufficient time to consider and address developments in this matter and the extent to which they moot the controversy at issue in the Petition. Therefore, interested parties will now have until June 10, 2011 to file comments and July 11, 2011 to file reply comments as opposed to the April 11, 2011 and May 11, 2011 deadlines set forth in the Public Notice.

Pursuant to §§ 1.415 and 1.419 of the Commission's rules, 47 CFR 1.415, 1.419, interested parties may file comments and reply comments on or before the dates indicated above. Comments may be filed using: (1) The Commission's Electronic Comment Filing System (ECFS), (2) the Federal Government's eRulemaking Portal, or (3) by filing paper copies. *See Electronic Filing of Documents in Rulemaking Proceedings,* 63 FR 24121 (1998).

- *Electronic Filers:* Comments may be filed electronically using the Internet by accessing the ECFS: *http://fjallfoss.fcc.gov/ecfs2/* or the Federal eRulemaking Portal: *http://www.regulations.gov.*
- *Paper Filers:* Parties who choose to file by paper must file an original and four copies of each filing. If more than one docket or rulemaking number appears in the caption of this proceeding, filers must submit two additional copies for each additional docket or rulemaking number.

Filings can be sent by hand or messenger delivery, by commercial overnight courier, or by first-class or overnight U.S. Postal Service mail (although we continue to experience delays in receiving U.S. Postal Service mail). All filings must be addressed to the Commission's Secretary, Office of the Secretary, Federal Communications Commission.

- All hand-delivered or messenger-delivered paper filings for the Commission's Secretary must be delivered to FCC Headquarters at 445 12th St., SW., Room TW–A325, Washington, DC 20554. The filing hours are 8 a.m. to 7 p.m. All hand deliveries must be held together with rubber bands or fasteners. Any envelopes must be disposed of *before* entering the building.
- Commercial overnight mail (other than U.S. Postal Service Express Mail and Priority Mail) must be sent to 9300 East Hampton Drive, Capitol Heights, MD 20743.
- U.S. Postal Service first-class, Express, and Priority mail must be addressed to 445 12th Street, SW., Washington DC 20554.

People with Disabilities: To request materials in accessible formats for people with disabilities (braille, large print, electronic files, audio format), send an e-mail to *fcc504@fcc.gov* or call the Consumer & Governmental Affairs Bureau at 202–418–0530 (voice), 202–418–0432 (tty).

Federal Communications Commission.

Nese Guendelsberger,

Chief, Spectrum and Competition Policy Division, Wireless Telecommunications Bureau.

[FR Doc. 2011–9199 Filed 4–15–11; 8:45 am]

BILLING CODE 6712–01–P

FEDERAL MARITIME COMMISSION

[Docket No. 11–06]

Indigo Logistics, LLC, Liliya Ivanenko, and Leonid Ivanenko—Possible Violations of Section 19 of the Shipping Act of 1984 and the Commission's Regulations; Order of Investigation and Hearing

AGENCY: Federal Maritime Commission.

ACTION: Notice of Order of Investigation and Hearing.

Authority: 46 U.S.C. 41302.

DATES: The Order of Investigation and Hearing was served April 7, 2011.

SUPPLEMENTARY INFORMATION: On April 7, 2011 the Federal Maritime Commission instituted an Order of Investigation and Hearing entitled Indigo Logistics, LLC; Liliya Ivanenko; and Leonid Ivanenko—Possible Violations of Section 19 of the Shipping Act of 1984 and the Commission's Regulations at 46 CFR part 515. Acting pursuant to Section 11 of the Shipping Act, 46 U.S.C. 41302, that investigation is instituted to determine:

(1) Whether Indigo Logistics, LLC, Liliya Ivanenko, and Leonid Ivanenko violated Section 19 of the Shipping Act, 46 U.S.C. 40901, 40902, and the Commission's regulations at 46 CFR part 515, by acting as an ocean freight forwarder without a license or evidence of financial responsibility;

(2) Whether, in the event violations of Section 19 of the Shipping Act of 1984 are found, civil penalties should be assessed against Indigo Logistics, LLC, Liliya Ivanenko, and/or Leonid Ivanenko, and, if so, the amount of penalties to be assessed; and

(3) Whether, in the event violations are found, appropriate cease and desist orders should be issued.

The Order may be viewed in its entirety at *http://www.fmc.gov.*

Karen V. Gregory,

Secretary.

[FR Doc. 2011–9282 Filed 4–15–11; 8:45 am]

BILLING CODE 6730–01–P

GENERAL SERVICES ADMINISTRATION

[OMB Control No. 3090–0200; Docket 2011–0001; Sequence 1]

General Services Administration Acquisition Regulation; Information Collection; Sealed Bidding

AGENCY: Office of the Chief Acquisition Officer, GSA.

ACTION: Notice of request for comments regarding a renewal to an existing OMB clearance.

SUMMARY: Under the provisions of the Paperwork Reduction Act (44 U.S.C. chapter 35), the Regulatory Secretariat (MVCB) will be submitting to the Office of Management and Budget (OMB) a request to review and approve an extension of a previously approved information collection requirement regarding sealed bidding.

Public comments are particularly invited on: Whether this collection of information is necessary and whether it will have practical utility; whether our estimate of the public burden of this collection of information is accurate and based on valid assumptions and methodology; and ways to enhance the quality, utility, and clarity of the information to be collected.

DATES: Submit comments on or before: June 17, 2011.

FOR FURTHER INFORMATION CONTACT: Michael O. Jackson, Procurement Analyst, Contract Policy Branch, at telephone (202) 208–4949 or *michaelo.jackson@gsa.gov.*

ADDRESSES: Submit comments identified by Information Collection 3090–0200 by any of the following methods:

- *Regulations.gov: http://www.regulations.gov.* Submit comments via the Federal eRulemaking portal by inputting "Information Collection 3090–0200" under the heading "Enter Keyword or ID" and selecting "Search". Select the link "Submit a Comment" that corresponds with "Information Collection 3090–0200". Follow the instructions provided at the "Submit a Comment" screen. Please include your name, company name (if any), and "Information Collection 3090–0200" on your attached document.

- *Fax:* 202–501–4067.
- *Mail:* General Services Administration, Regulatory Secretariat (MVCB), 1275 First Street, NE., Washington, DC 20417. *Attn:* Hada Flowers/IC 3090–0200.

Instructions: Please submit comments only and cite Information Collection 3090–0200, in all correspondence related to this collection. All comments received will be posted without change to *http://www.regulations.gov,* including any personal and/or business confidential information provided.

SUPPLEMENTARY INFORMATION:

A. Purpose

The General Services Administration is requesting that the Office of Management and Budget (OMB) review and approve information collection, 3090–0200, Sealed Bidding. The information requested regarding an offeror's monthly production capability is needed to make progressive awards to ensure coverage of stock items.

B. Annual Reporting Burden

Respondents: 10.
Responses per Respondent: 1.
Hours per Response: .5.
Total Burden Hours: 5.
Obtaining Copies of Proposals: Requesters may obtain a copy of the information collection documents from the General Services Administration, Regulatory Secretariat (MVCB), 1275 First Street, NE., Washington, DC 20417, telephone (202) 501–4755. Please cite OMB Control No. 3090–0200, Sealed Bidding, in all correspondence.

Dated: April 11, 2011.

Millisa Gary,

Acting Director, Office of Governmentwide Acquisition Policy.

[FR Doc. 2011–9264 Filed 4–15–11; 8:45 am]

BILLING CODE 6820–61–P

DEPARTMENT OF HEALTH AND HUMAN SERVICES

Agency for Healthcare Research and Quality

Agency Information Collection Activities: Proposed Collection; Comment Request

AGENCY: Agency for Healthcare Research and Quality, HHS.

ACTION: Notice.

SUMMARY: This notice announces the intention of the Agency for Healthcare Research and Quality (AHRQ) to request that the Office of Management and Budget (OMB) approve the proposed information collection project: "Patient Safety Organization Certification for Initial Listing and Related Forms, Patient Safety Confidentiality Complaint Form, and Common Formats. In accordance with the Paperwork Reduction Act, 44 U.S.C. 3501–3521, AHRQ invites the public to comment on this proposed information collection.

DATES: Comments on this notice must be received by June 17, 2011.

ADDRESSES: Written comments should be submitted to: Doris Lefkowitz, Reports Clearance Officer, AHRQ, by e-mail at *doris.lefkowitz@AHRQ.hhs.gov.* Copies of the proposed collection plans, data collection instruments, and specific details on the estimated burden can be obtained from the AHRQ Reports Clearance Officer.

FOR FURTHER INFORMATION CONTACT: Doris Lefkowitz, AHRQ Reports Clearance Officer, (301) 427–1477, or by e-mail at *dorislefkowitz@AHRQ.hhs.gov.*

SUPPLEMENTARY INFORMATION:

Proposed Project

Patient Safety Organization Certification for Initial Listing and Related Forms, Patient Safety Confidentiality Complaint Form, and Common Formats

The Patient Safety and Quality Improvement Act of 2005 (hereafter the Patient Safety Act), 42 U.S.C. 299b–21 to 299b–26, was enacted in response to growing concern about patient safety in the United States and the Institute of Medicine's 1999 report, To Err is Human: Building a Safer Health System. The goal of the statute is to improve patient safety by providing an incentive for health care providers to work voluntarily with experts in patient safety to reduce risks and hazards to the safety and quality of patient care. The Patient Safety Act signifies the Federal Government's commitment to fostering a culture of patient safety among health care providers; it offers a mechanism for creating an environment in which the causes of risks and hazards to patient safety can be thoroughly and honestly examined and discussed without fear of penalties and liabilities. It provides for the voluntary formation of Patient Safety Organizations (PSOs) that can collect, aggregate, and analyze confidential information reported voluntarily by health care providers. By analyzing substantial amounts of patient safety event information across multiple institutions, PSOs will be able to identify patterns of failures and propose measures to eliminate or reduce patient safety risks and hazards.

In order to implement the Patient Safety Act, the Department of Health and Human Services (HHS) issued the Patient Safety and Quality Improvement Final Rule (hereafter the Patient Safety Rule), 42 CFR part 3, which became effective on January 19, 2009. The Patient Safety Rule establishes a framework by which hospitals, doctors, and other health care providers may voluntarily report information to PSOs, on a privileged and confidential basis, for the aggregation and analysis of patient safety events. In addition, the Patient Safety Rule outlines the requirements that entities must meet to become PSOs and the process by which the Secretary of HHS (hereafter the Secretary) will review and accept certifications and list PSOs.

In addition to the Patient Safety Act and the Patient Safety Rule, HHS issued Guidance Regarding Patient Safety Organizations' Reporting Obligations and the Patient Safety and Quality Improvement Act of 2005 (hereafter Guidance) on December 30, 2010. The Guidance addresses questions that have arisen regarding the obligations of PSOs where they or the organization of which they are a part are legally obligated under the Federal Food, Drug, and Cosmetic Act (FDCA) and its implementing regulations to report certain information to the Food and Drug Administration (FDA) and to provide FDA with access to its records, including access during an inspection of its facilities. This Guidance applies to all entities that seek to be or are PSOs or component PSOs that have mandatory FDA-reporting obligations under the FDCA and its implementing regulations ("FDA-regulated reporting entities") or are organizationally related to such FDA-regulated reporting entities (*e.g.,* parent organizations, subsidiaries, sibling organizations).

When PSOs meet the requirements of the Patient Safety Act, the information collected and the analyses and deliberations regarding the information receive Federal confidentiality and privilege protections under this legislation. The Secretary delegated authority to the Director of the Office for Civil Rights (OCR) to enforce the confidentiality protections of the Patient Safety Act. 71 **Federal Register** 28701–28702 (May 17, 2006). OCR is responsible for enforcing protections regarding patient safety work product (PSWP), which generally includes information that could improve patient safety, health care quality, or health care outcomes and (1) is assembled or developed by a provider for reporting to a PSO and is reported to a PSO or (2) is developed by a PSO for the conduct of patient safety activities. Civil money penalties may be imposed for knowing or reckless impermissible disclosures of