OEF; that Respondents Louis J. Mischianti, David Cardenas, and Keith Heffernan, are the partners in Olympus Partners, and were officers and directors of Global Link; that Respondent CJR World Enterprises, Inc. (''CJR''), is a Florida corporation that was an owner of Global Link; and that Respondent Chad Rosenberg is the owner of CJR and was an officer and director of Global Link.

Complainant alleges that Respondents violated the Shipping Act of 1984, as amended (''Shipping Act''), by: (1) Engaging in a deliberate scheme to obtain ocean transportation of property at rates lower than the applicable service contract or tariff rates; and (2) failing to establish, observe and enforce just and reasonable practices relating to or connected with receiving, handling, and delivering property. 46 U.S.C. 41102(a), (c). Complainant also asserts that Respondents violated the Commission's regulations at 46 CFR 515.31(e) which prohibits preparation or filing of false or fraudulent claims or false information relative to an Ocean Transportation Intermediary transaction. Complainant claims that, as a direct result of Respondents' actions, Complainant suffered damages of no less than $4.5 million.

Specifically, Complainant MOL asserts that it provided transportation to Global Link subject to MOL's tariff rules; including rules related to the diversion of cargo, defined as a change in the original billed destination. Complainant maintains that its tariff rules require shippers to request diversion of cargo in writing and require payment of a diversion charge, as well as the difference in price between the original and new destination. Complainant alleges that Respondent Global Link booked cargo to false inland destinations while intending to deliver the cargo to different inland destinations, and diverted cargo without submitting a request to Complainant or paying Complainant the difference in rate and the applicable diversion changes. Complainant claims that Respondents referred to this practice as ''split routing,'' ''mis-booking,'' and re-routing.'' This practice, Complainant contends, resulted in lower rates paid to Complainant than the rates applicable to the actual destinations.

Complainant requests that the Commission: (1) Require Respondents to answer the charges in this Complaint; (2) order Respondents to cease and desist from the violations of the Shipping Act; (3) establish and put in force such practices as the Commission determines lawful and reasonable; (3) order Respondents to pay to the Complainant reparations plus interests, costs and attorney's fees, and any other damages to be determined; and (4) take any other action or provide any other relief as the Commission determines to be proper, fair and just under the circumstances. Complainant also requests that a hearing be held in Washington, DC.

This proceeding has been assigned to the Office of Administrative Law Judges. Hearing in this matter, if any is held, shall commence within the time limitations prescribed in 46 CFR 502.61, and only after consideration has been given by the parties and the presiding officer to the use of alternative forms of dispute resolution. The hearing shall include oral testimony and cross-examination in the discretion of the presiding officer only upon proper showing that there are genuine issues of material fact that cannot be resolved on the basis of sworn statements, affidavits, depositions, or other documents or that the nature of the matter in issue is such that an oral hearing and cross-examination are necessary for the development of an adequate record. Pursuant to the further terms of 46 CFR 502.61, the initial decision of the presiding officer in this proceeding shall be issued by May 14, 2010, and the final decision of the Commission shall be issued by September 13, 2010.

Karen V. Gregory,

Secretary.

[FR Doc. E9–11755 Filed 5–20–09; 8:45 am]

BILLING CODE 6730–01–P

DEPARTMENT OF HEALTH AND HUMAN SERVICES

CBRN Medical Countermeasures Workshop 2009

AGENCY: Department of Health and Human Services, Assistant Secretary for Preparedness and Response.

ACTION: Notice.

SUMMARY: As stipulated by the Federal Advisory Committee Act, the U.S. Department of Health and Human Services is hereby giving notice that the Biomedical Advanced Development Authority (BARDA), Chemical Biological, Radiological & Nuclear (CBRN) Medical Countermeasures and Acquisitions Management Systems (AMS) will be holding a public workshop. The workshop is open to the public.

DATES: The BARDA Divisions of CBRN and AMS will hold a public workshop on June 25 and 26, 2009 from 8:30 a.m. to 5 p.m. EDT daily. This agenda is subject to change as priorities dictate.

ADDRESSES: Hyatt Regency Bethesda, One Bethesda Metro Center, Bethesda, MD 20814. Phone: 301–657–1234.

FOR FURTHER INFORMATION, CONTACT: *CBRN-Workshop@hhs.gov.*

SUPPLEMENTARY INFORMATION: The U.S. Department of Health and Human Services, Office of the Secretary, Office of the Assistant Secretary for Preparedness and Response, Biomedical Advanced Research and Development Authority, Division of Chemical, Biological, Radiological and Nuclear and the Division of Acquisitions Management Systems will host a 2-day Medical Countermeasures Workshop.

The purpose of this two-day event is to engage industry and academic stakeholders in a discussion of how to engage and work with BARDA. Informational sessions include:

- The strategic vision for CBRN MCM development.
- The generation of MCM requirements.
- Responding to a Request for Proposal (RFP) or Broad Agency Announcement (BAA).
- The proposal review process.
- Earned Value Management.
- The FDA Animal Rule.

This Workshop is open to the public. There is no fee to attend; however, seating is limited and registration is required. Online registration is available at *http://www.medicalcountermeasures.gov.* Stakeholder Registration will be open from April 14–June 1, 2009. Federal Government Employee Registration will be open from June 2–12, 2009.

Availability of Materials: The workshop agenda and other materials will be available on site on the workshop dates.

Dated: May 6, 2009.

RADM William C. Vanderwagen,

Assistant Secretary for Preparedness and Response, U.S. Department of Health and Human Services.

[FR Doc. E9–11948 Filed 5–20–09; 8:45 am]

BILLING CODE P

DEPARTMENT OF HEALTH AND HUMAN SERVICES

National Institutes of Health

Submission to OMB, Comment Request; A Process Evaluation of the NIH Director's New Innovator Award (NIA) Program

SUMMARY: Under the provisions of Section 3507(a)(1)(D) of the Paperwork Reduction Act of 1995, the Office of the

Director, National Institutes of Health (NIH), has submitted to the Office of Management and Budget (OMB) a request for review and approval of the information collection listed below. This proposed information collection was previously published in the **Federal Register** on February 20, 2009, pages 7908–7909 and allowed 60 days for public comment. No public comments were received. The National Institutes of Health may not conduct or sponsor, and the respondent is not required to respond to, an information collection that has been extended, revised, or implemented on or after October 1, 1995, unless it displays a currently valid OMB control number.

Proposed Collection: Title: A Process Evaluation of the NIH Director's New Innovator Award (NIA) Program. *Type of Information Collection Request:* New collection. *Need and Use of Information Collection:* This study will assess the NIA Program operations and the outputs of the identification, evaluation and selection process. The primary objectives of the study are to: (1) Assess the NIA award selection process; (2) determine if the program was implemented as planned; and (3) determine if the process was conducted in accordance with the overall mission of the NIA program. The findings will provide valuable information concerning: (1) The characteristics of applicants and reviewers; (2) the criteria used to evaluate and select awardees; and (3) aspects of the process that could be revised or improved.

Frequency of Response: Once. *Affected Public:* None. *Type of Respondents:* Applicants, Reviewers. There are no Capital Costs to report. There are no Operating or Maintenance Costs to report. *Estimated Number of Respondents:* 662; *Estimated Number of Responses per Respondent:* 1; *Average Burden Hours per Response:* .28 (15 minutes for applicants and 30 minutes for Extramural Reviewers), and *Estimated Total Annual Burden Hours Requested:* 188.5 and the annualized cost to respondents is estimated at $12,199.72. Table 1 and Table 2 respectively present data concerning the burden hours and cost burdens for this data collection.

TABLE 1—ANNUALIZED ESTIMATE OF HOUR BURDEN

Type of respondents	Number of respondents	Frequency of response	Average time for response (hr)	Total hour burden *
Applicants	570	1	.25	142.5
Extramural Reviewers	92	1	.50	46
Total	662	1	.28	188.5

* Total Burden = N Respondents * Response Frequency * (minutes to complete/60).

TABLE 2—ANNUALIZED COST TO RESPONDENTS

Type of respondents	Number of respondents	Response frequency	Approx. hourly wage rate	Total respondent cost **
Applicants	570	1	$64.72	$9,226.60
Extramural Reviewers	92	1	64.72	2977.12
Total	662	1	64.72	12,199.72

**Total Respondent Cost = Total Hour Burden * Hourly Wage Rate.

Request for Comments: Written comments and/or suggestions from the public and affected agencies are invited on one or more of the following points: (1) Whether the proposed collection of information is necessary for the proper performance of the function of the agency, including whether the information will have practical utility; (2) 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) ways to enhance the quality, utility, and clarity of the information to be collected; and (4) ways to minimize the burden of the collection of information on those who are to respond, including the use of appropriate automated, electronic, mechanical, or other technological collection techniques or other forms of information technology.

Direct Comments to OMB: Written comments and/or suggestions regarding the item(s) contained in this notice, especially regarding the estimated public burden and associated response time, should be directed to the Office of Management and Budget, Office of Regulatory Affairs (OIRA). All comments should be sent via e-mail to *OIRA_submission@omb.eop.gov* or by fax to 202–395–6974. Attention: Desk Office for NIH. To request more information on the project or to obtain a copy of the data collection plans and instruments contact G. Stephane Philogene, PhD, Assistant Director for Policy and Planning, Office of Behavioral and Social Sciences Research, National Institutes of Health, 31 Center Drive, Building 31, Room B2–B37, Bethesda, MD 20892, or call non-toll-free number 301–402–3902 or e-mail your request, including your address, to: *philoges@od.nih.gov*.

Comments Due Date: Comments regarding this information collection are best assured of having their full effect if received within 30 days of the date of this publication.

Dated: May 13, 2009.

G. Stephane Philogene,

Assistant Director for Policy and Planning, Office of Behavioral and Social Sciences Research, National Institutes of Health.

[FR Doc. E9–11817 Filed 5–20–09; 8:45 am]

BILLING CODE 4140–01–P

DEPARTMENT OF HEALTH AND HUMAN SERVICES

Centers for Disease Control and Prevention

[30Day–09–09AK]

Agency Forms Undergoing Paperwork Reduction Act Review

The Centers for Disease Control and Prevention (CDC) publishes a list of information collection requests under review by the Office of Management and