decision of the Commission shall be issued by November 14, 2011.

Karen V. Gregory,

Secretary.

[FR Doc. 2010–17786 Filed 7–20–10; 8:45 am]

BILLING CODE P

FEDERAL MARITIME COMMISSION

Ocean Transportation Intermediary License Applicants

Notice is hereby given that the following applicants have filed with the Federal Maritime Commission an application for a license as a Non-Vessel-Operating Common Carrier (NVO) and/or Ocean Freight Forwarder (OFF)—Ocean Transportation Intermediary (OTI) pursuant to section 19 of the Shipping Act of 1984 as amended (46 U.S.C. Chapter 409 and 46 CFR 515). Notice is also hereby given of the filing of applications to amend an existing OTI license or the Qualifying Individual (QI) for a license.

Interested persons may contact the Office of Transportation Intermediaries, Federal Maritime Commission, Washington, DC 20573.

American Lamprecht Transport, Inc. (NVO & OFF), 700 Rockaway Turnpike, Lawrence, NY 11559. *Officers:* Alain Tiercy, CFO/Secretary/Treasurer (Qualifying Individual), Hans-Peter Widmer, President.
Application Type: QI Change.

CACC Global Logistics, Inc. (NVO & OFF), 151 E. 220th Street, Carson, CA 90754. *Officers:* Annie Sun, President/CEO (Qualifying Individual), Chuck Sun, Vice President/Secretary.
Application Type: New NVO & OFF License.

E-Freight Solutions Inc. dba E-Lines Shipping and Logistics, and Ocean Champ Shipping Limited (NVO), 1000 Corporate Center Drive, Suite 320, Monterey Park, CA 91754. *Officers:* Joey Tam, President/CEO (Qualifying Individual), Yu C. Lee, Secretary/Treasurer.
Application Type: Name Change.

Ever-Swift Worldwide Inc. (NVO & OFF), Cargo Bldg. 151, Room 377, Jamaica, NY 11430. *Officer:* Chiang Yu-Chen, President (Qualifying Individual).
Application Type: Add OFF Service.

Limitless International, Inc. (NVO & OFF), 8750 Exchange Drive, #3, Orlando, FL 32809. *Officer:* Cheryl A. Stockstad, President (Qualifying Individual).
Application Type: Add NVO Service.

Meadwestvaco Corporation (NVO & OFF), 501 South 5th Street, Richmond, VA 23219. *Officers:* Christopher L. Osen, Vice President Supply Management (Qualifying Individual), Susan J. Kropf, Director.
Application Type: New NVO & OFF License.

Mutual Pacific Logistics, Inc. (NVO), 12801 South Figueroa Street, Los Angeles, CA 90061. *Officer:* Chee (CT) T. Tsui, President/Secretary/Treasurer (Qualifying Individual).
Application Type: New NVO License.

Unity Container Line, Inc. (NVO & OFF), 12552 SW. 143 Lane, Miami, FL 33186. *Officer:* Pedro Streb, President/Secretary/Treasurer (Qualifying Individual).
Application Type: New NVO & OFF License.

Dated: July 16, 2010.

Karen V. Gregory,

Secretary.

[FR Doc. 2010–17784 Filed 7–20–10; 8:45 am]

BILLING CODE 6730–01–P

FEDERAL TRADE COMMISSION

Sunshine Act Meeting Notice

AGENCY: Federal Trade Commission.

TIME AND DATE: 2 p.m., Wednesday, July 28, 2010.

PLACE: Federal Trade Commission Building, Room 532, 600 Pennsylvania Avenue, NW., Washington, DC 20580.

STATUS: Part of this meeting will be open to the public. The rest of the meeting will be closed to the public.

Matters To Be Considered

Portion Open to the Public

(1) Oral Argument in Polypore International, Inc., Docket 9327.

Portion Closed to the Public

(2) Executive Session to follow Oral Argument in Polypore International, Inc., Docket 9327.

CONTACT PERSON FOR MORE INFORMATION: Mitch Katz, Office of Public Affairs, (202) 326–2180. *Recorded Message:* (202) 326–2711.

Donald S. Clark,

Secretary.

[FR Doc. 2010–17651 Filed 7–20–10; 8:45 am]

BILLING CODE 6750–01–M

DEPARTMENT OF HEALTH AND HUMAN SERVICES

Food and Drug Administration

[Docket No. FDA–2010–N–0373]

Agency Information Collection Activities; Proposed Collection; Comment Request; Preparing a Claim of Categorical Exclusion or an Environmental Assessment for Submission to the Center for Food Safety and Applied Nutrition

AGENCY: Food and Drug Administration, HHS.

ACTION: Notice.

SUMMARY: The Food and Drug Administration (FDA) is announcing an opportunity for public comment on the proposed collection of certain information by the agency. Under the Paperwork Reduction Act of 1995 (the PRA), Federal agencies are required to publish notice in the **Federal Register** concerning each proposed collection of information, including each proposed extension of an existing collection of information, and to allow 60 days for public comment in response to the notice. This notice solicits comments on the information collection provisions in the guidance document entitled "Preparing a Claim of Categorical Exclusion or an Environmental Assessment for Submission to the Center for Food Safety and Applied Nutrition."

DATES: Submit either electronic or written comments on the collection of information by September 20, 2010.

ADDRESSES: Submit electronic comments on the collection of information to *http://www.regulations.gov*. Submit written comments on the collection of information to the Division of Dockets Management (HFA–305), Food and Drug Administration, 5630 Fishers Lane, rm. 1061, Rockville, MD 20852. All comments should be identified with the docket number found in brackets in the heading of this document.

FOR FURTHER INFORMATION CONTACT: Denver Presley, Jr., Office of Information Management, Food and Drug Administration, 1350 Piccard Dr., PI50–400B, Rockville, MD 20850, 301–796–3793.

SUPPLEMENTARY INFORMATION: Under the 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. "Collection of information" is defined in 44 U.S.C. 3502(3) and 5 CFR 1320.3(c)

and includes agency requests or requirements that members of the public submit reports, keep records, or provide information to a third party. Section 3506(c)(2)(A) of the PRA (44 U.S.C. 3506(c)(2)(A)) requires Federal agencies to provide a 60-day notice in the **Federal Register** concerning each proposed collection of information, including each proposed extension of an existing collection of information, before submitting the collection to OMB for approval. To comply with this requirement, FDA is publishing notice of the proposed collection of information set forth in this document.

With respect to the following collection of information, FDA invites comments on these topics: (1) Whether the proposed collection of information is necessary for the proper performance of FDA's functions, including whether the information will have practical utility; (2) the accuracy of FDA'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 respondents, including through the use of automated collection techniques, when appropriate, and other forms of information technology.

Preparing a Claim of Categorical Exclusion or an Environmental Assessment for Submission to the Center for Food Safety and Applied Nutrition (OMB Control Number 0910–0541)—Extension

As an integral part of its decisionmaking process, FDA is obligated under the National Environmental Policy Act of 1969 (NEPA) to consider the environmental impact of its actions, including allowing notifications for food contact substances to become effective and approving food additive petitions, color additive petitions, GRAS affirmation petitions, requests for exemption from regulation as a food additive, and actions on certain food labeling citizen petitions, nutrient content claims petitions, and health claims petitions. In 1997, FDA amended its regulations in part 25 (21 CFR part 25) to provide for categorical exclusions for additional classes of actions that do not individually or cumulatively have a significant effect on the human environment (62 FR 40570, July 29, 1997). As a result of that rulemaking, FDA no longer routinely requires submission of information about the manufacturing and production of FDA-regulated articles. FDA also has eliminated the previously required Environmental Assessment (EA) and abbreviated EA formats from the amended regulations. Instead, FDA has provided guidance that contains sample formats to help industry submit a claim of categorical exclusion or an EA to FDA's Center for Food Safety and Applied Nutrition (CFSAN). The guidance document entitled "Preparing a Claim of Categorical Exclusion or an Environmental Assessment for Submission to the Center for Food Safety and Applied Nutrition" identifies, interprets, and clarifies existing requirements imposed by statute and regulation, consistent with the Council on Environmental Quality regulations (40 CFR 1507.3). It consists of recommendations that do not themselves create requirements; rather, they are explanatory guidance for FDA's own procedures in order to ensure full compliance with the purposes and provisions of NEPA.

The guidance provides information to assist in the preparation of claims of categorical exclusion and EAs for submission to CFSAN. The following questions are covered in this guidance: (1) What types of industry-initiated actions are subject to a claim of categorical exclusion? (2) what must a claim of categorical exclusion include by regulation? (3) what is an EA? (4) when is an EA required by regulation and what format should be used? (5) what are extraordinary circumstances? and (6) what suggestions does CFSAN have for preparing an EA? Although CFSAN encourages industry to use the EA formats described in the guidance because standardized documentation submitted by industry increases the efficiency of the review process, alternative approaches may be used if these approaches satisfy the requirements of the applicable statutes and regulations. FDA is requesting the extension of OMB approval for the information collection provisions in the guidance.

Description of Respondents: The likely respondents include businesses engaged in the manufacture or sale of food, food ingredients, and substances used in materials that come into contact with food.

FDA estimates the burden of this collection of information as follows:

TABLE 1.—ESTIMATED ANNUAL REPORTING BURDEN[1]

21 CFR Section	No. of Respondents	Annual Frequency per Response	Total Annual Responses	Hours per Response	Total Hours
25.32(i)	34	1	34	1	34
25.32(o)	1	1	1	1	1
25.32(q)	2	1	2	1	2
Total					37

[1] There are no capital costs or operating and maintenance costs associated with this collection of information.

The estimates for respondents and numbers of responses are based on the annualized numbers of petitions and notifications qualifying for § 25.32(i) and (q) that the agency has received in the past 3 years. Please note that, in the past 3 years, there have been no submissions that requested an action that would have been subject to the categorical exclusion in § 25.32(o). To avoid counting this burden as zero, FDA has estimated the burden for this categorical exclusion at one respondent making one submission a year for a total of one annual submission.

To calculate the estimate for the hours per response values, we assumed that the information requested in this guidance for each of these three categorical exclusions is readily available to the submitter. For the information requested for the exclusion in § 25.32(i), we expect that the submitter will need to gather information from appropriate persons in the submitter's company and to prepare this information for attachment to the

claim for categorical exclusion. We believe that this effort should take no longer than 1 hour per submission. For the information requested for the exclusions in § 25.32(o) and (q), the submitters will almost always merely need to copy existing documentation and attach it to the claim for categorical exclusion. We believe that collecting this information should also take no longer than 1 hour per submission.

Dated: July 12, 2010.

Leslie Kux,

Acting Assistant Commissioner for Policy.

[FR Doc. 2010–17751 Filed 7–20–10; 8:45 am]

BILLING CODE 4160–01–S

DEPARTMENT OF HEALTH AND HUMAN SERVICES

Administration on Aging

Agency Information Collection Activities; Proposed Collection; Comment Request; Extension of Supplemental Form to the Financial Status Report for all AoA Title III Grantees

AGENCY: Administration on Aging, HHS.

ACTION: Notice.

SUMMARY: The Administration on Aging (AoA) is announcing an opportunity for public comment on the proposed collection of certain information by the agency. Under the Paperwork Reduction Act of 1995 (the PRA), Federal agencies are required to publish notice in the **Federal Register** concerning each proposed collection of information, including each proposed extension of an existing collection of information, and to allow 60 days for public comment in response to the notice. This notice solicits comments on the information collection requirements relating to the Supplemental Form to the Financial Status Report for all AoA Title III Grantees.

DATES: Submit written or electronic comments on the collection of information by September 20, 2010.

ADDRESSES: Submit electronic comments on the collection of information to: *Rimas.Liogys@aoa.hhs.gov.* Submit written comments on the collection of information to Administration on Aging, Washington, DC 20201.

FOR FURTHER INFORMATION CONTACT: Rimas Liogys, Director of Grants Management, Administration on Aging, Washington, DC 20201.

SUPPLEMENTARY INFORMATION: Under the 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. "Collection of information" is defined in 44 U.S.C. 3502(3) and 5 CFR 1320.3(c) and includes agency request or requirements that members of the public submit reports, keep records, or provide information to a third party. Section 3506(c)(2)(A) of the PRA (44 U.S.C. 3506(c)(2)(A)) requires Federal agencies to provide a 60-day notice in the **Federal Register** concerning each proposed collection of information, including each proposed extension of an existing collection of information, before submitting the collection to OMB for approval. To comply with this requirement, AoA is publishing notice of the proposed collection of information set forth in this document. With respect to the following collection of information, AoA invites comments on: (1) Whether the proposed collection of information is necessary for the proper performance of AoA's functions, including whether the information will have practical utility; (2) the accuracy of AoA'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 respondents, including through the use of automated collection techniques when appropriate, and other forms of information technology. The template may be found on the AoA Web site at *http://www.aoa.gov/AoARoot/Grants/Reporting_Requirements/Formula_269.aspx.*

The Supplemental form to the Financial Status Report for all AoA Title III Grantees provides an understanding of how projects funded by the Older Americans Act are being administered by grantees, in conformance with legislative requirements, pertinent Federal regulations and other applicable instructions and guidelines issued by Administration on Aging (AoA). This information will be used for Federal oversight of Title III Projects. AoA estimates the burden of this collection of information as follows: 56 State Agencies on Aging respond semiannually which should be an average burden of 1 hour per State agency per submission for a total of 56 hours.

Kathy Greenlee,

Assistant Secretary for Aging.

[FR Doc. 2010–17822 Filed 7–20–10; 8:45 am]

BILLING CODE 4154–01–P

DEPARTMENT OF HEALTH AND HUMAN SERVICES

Centers for Disease Control and Prevention

Board of Scientific Counselors, Coordinating Office for Terrorism Preparedness and Emergency Response; Notice of Charter Amendment

This gives notice under the Federal Advisory Committee Act (Pub. L. 92–463) of October 6, 1972, that the Board of Scientific Counselors, Coordinating Office for Terrorism Preparedness and Emergency Response, Department of Health and Human Services, has amended their charter to reflect the change in the name of the board to the Board of Scientific Counselors, Office of Public Health Preparedness and Response.

For information, contact Barbara Ellis, Ph.D, Executive Secretary, Board of Scientific Counselors, Office of Public Health Preparedness and Response, Department of Health and Human Services, 1600 Clifton Road, M/S D44, Atlanta, Georgia 30341, telephone (404)639–0637, or fax (404)639–7977.

The Director, Management Analysis and Services Office, has been delegated the authority to sign **Federal Register** notices pertaining to announcements of meetings and other committee management activities, for both the CDC and the Agency for Toxic Substances and Disease Registry.

Dated: July 13, 2010.

Elaine L. Baker,

Director, Management Analysis and Services Office, Centers for Disease Control and Prevention.

[FR Doc. 2010–17761 Filed 7–20–10; 8:45 am]

BILLING CODE 4163–18–P

DEPARTMENT OF HEALTH AND HUMAN SERVICES

Centers for Disease Control and Prevention (CDC)

Board of Scientific Counselors (BSC), Coordinating Center for Health Promotion (CCHP): Notice of Charter Amendment

This gives notice under the Federal Advisory Committee Act (Pub. L. 92–463) of October 6, 1972, that the BSC, CCHP, has amended their charter to reflect the change in the name of the board to the BSC, National Center on Birth Defects and Developmental Disabilities (NCBDDD) and National Center for Chronic Disease Prevention and Health Promotion (NCCDPHP).