Intership, Inc dba Helm Express (NVO & OFF), 2530 Knoblock, Houston, TX 77023. Officer: Yasser Shaikh, President, (Qualifying Individual), Application Type: Add NVO Service and Trade Name Change.

Jupiter Airline Services, Inc. dba Mercury Logistics (NVO), 5456 McConnell Avenue, Los Angeles, CA 90066. Officers: Zack Vernikovsky, Vice President/Director, (Qualifying Individual), Joseph A. Czyzyk, CEO/Director, Application Type: New NVO License.

Legend Express Co. (OFF), 1506 S. Paloma Street, Los Angeles, CA 90021. Officers: Gila Morad, Chief Executive Officer/Chief Financial Officer, (Qualifying Individual), Natali Morad, Secretary, Application Type: QI Change.

Linsan.Tex Investments, L.L.C. (OFF), 8404 Endicott Lane, Dallas, TX 75227. Officers: Franklin E. Aigbuza, Secretary/Member, (Qualifying Individual), Roseline A. Izedonmwen, CEO/Member, Application Type: New OFF License.

Ocean Air Land Freight, Corp. (OFF), 8600 NW 30th Terrace 2nd Floor, Miami, FL 33122. Officers: Martha Zuluaga, President, (Qualifying Individual), Maria J. Gori, Secretary/Treasurer, Application Type: Trade Name Change.

Ocean Channel Shipping Co., Ltd. (NVO), 13091 Nordland Drive, Corona, CA 92880. Officer: Xiaohua Huo, President, (Qualifying Individual), Application Type: New NVO License. Siman Logistics, Inc. (NVO & OFF), 765 N, IL Route 83, Suite 124, Bensenville, IL 60106. Officers: Wolfgang A. Ries, Senior Vice President, (Qualifying Individual), Christian Ludwig, President, Application Type: New NVO & OFF License.

Top Wise Logistics Inc. (NVO), 654 N. Spring Street, Los Angeles, CA 90012. Officer: George N. Lee, CEO/CFO/Secretary, (Qualifying Individual), Application Type: New NVO License.

Trinity Logistics USA, Inc. (NVO), 10 East Merrick Road, Suite 304, Valley Stream, NY 11580. Officers: Doris McGregory, Treasurer, (Qualifying Individual), David Pereira, President/Secretary, Application Type: New NVO License.

Twenty Two Global Transport, LP (NVO & OFF), 1911 Bagby Street, Houston, TX 77002. Officers: Kevin A. Smoot, Partner/Director, (Qualifying Individual), Robert Crossland, Vice President, Application Type: New NVO & OFF License.

United Marine Lines, L.L.C. (NVO), 201 Sevilla Avenue, Suite 309, Coral Gables, FL 33134. Officers: Eduardo Del Riego, President/Secretary, (Qualifying Individual), Robert Boucek, Vice President/Treasurer, Application Type: New NVO License.

Dated: July 9, 2010.

Karen V. Gregory,

Secretary.

[FR Doc. 2010–17175 Filed 7–13–10; 8:45 am]

BILLING CODE 6730–01–P

DEPARTMENT OF HEALTH AND HUMAN SERVICES

Establishment of the Advisory Board on Elder Abuse, Neglect, and Exploitation

AGENCY: Department of Health and Human Services, Administration on Aging.

ACTION: Notice.

AUTHORITY: The Advisory Board on Elder Abuse, Neglect, and Exploitation is authorized under section 2021, Subtitle H—Elder Justice Act, of the Affordable Care Act, Public Law 111–148. The Advisory Board is governed by provisions of Public Law 92–463, as amended, (5 U.S.C. App. 2), which sets forth standards for the formation and use of advisory committees.

SUMMARY: The U.S. Department of Health and Human Services announces establishment of the Advisory Board on Elder Abuse, Neglect, and Exploitation, as directed by section 2022, Subtitle H—Elder Justice Act, of the Affordable Care Act, Public Law 111–148.

FOR FURTHER INFORMATION CONTACT: Edwin Walker, Deputy Assistant Secretary for Program Operations, Department of Health and Human Services, Administration on Aging, Washington, DC 20201, Telephone: 202–357–3557, Fax: 202–357–3549.

SUPPLEMENTARY INFORMATION: Subtitle H—Elder Justice Act of the Affordable Care Act, Public Law 111–148, establishes the Advisory Board within the Department of Health and Human Services (HHS). To comply with the authorizing directive and guidelines under the Federal Advisory Committee Act (FACA), a charter has been filed with the Committee Management Secretariat in the General Services Administration (GSA), the appropriate committees in the Senate and U.S. House of Representatives, and the Library of Congress to establish the Advisory Board as a non-discretionary Federal advisory committee. The charter was filed on July 8, 2010.

Objectives and Scope of Activities

The Advisory Board on Elder Abuse, Neglect, and Exploitation (Advisory Board) is the Department's statutory public advisory body in the Elder Justice Act on creating short- and long-term multidisciplinary strategic plans for the development of the field of elder justice in the U.S. The Advisory Board will examine relevant research and identify best practices and make recommendations to the Elder Justice Coordinating Council and Congress regarding improving and enhancing Federal, State, and local elder justice programs, research, training, and coordination.

Membership and Designation

The Secretary is soliciting nominations for appointment to the 27-member Advisory Board from among members of the general public who are individuals with experience and expertise in elder abuse, neglect, and exploitation prevention, detection, treatment, intervention, or prosecution. Each member of the Advisory Board shall be appointed for a term of 3 years except that, of the members first appointed, 9 shall be appointed for a term of 3 years; 9 shall be appointed for a term of 2 years; and 9 shall be appointed for a term of 1 year. Nominations shall be submitted to: Edwin Walker, Deputy Assistant Secretary for Program Operations, Department of Health and Human Services, Administration on Aging, Washington, DC 20201, no later than August 15, 2010. Any vacancy on the Advisory Board shall not affect its powers, but shall be filled in the same manner as the original appointment was made. An individual chosen to fill a vacancy shall be appointed for the unexpired term of the member replaced. The Advisory Board shall elect a Chair and Vice Chair from among its members.

Administrative Management and Support

HHS will provide funding and administrative support for the Advisory Board to the extent permitted by law within existing appropriations. Management and oversight for support services provided to the Advisory Board will be the responsibility of the Administration on Aging, which is an operating division within HHS. Staff will be assigned to support the activities of the Advisory Board. All executive departments and agencies shall provide information to the Advisory Board as the Chair may request for purposes of carrying out the Advisory Board's

functions, to the extent permitted by law. A copy of the Commission charter can be obtained from the designated contacts or by accessing the FACA database that is maintained by the GSA Committee Management Secretariat. The website for the FACA database is *http://fido.gov/facadatabase/*.

Dated: July 9, 2010.

Kathy Greenlee,

Assistant Secretary for Aging.

[FR Doc. 2010–17197 Filed 7–13–10; 8:45 am]

BILLING CODE 4154–01–P

DEPARTMENT OF HEALTH AND HUMAN SERVICES

Food and Drug Administration

[Docket No. FDA–2010–N–0357]

Agency Information Collection Activities; Proposed Collection; Comment Request; Hazard Analysis and Critical Control Point Procedures for the Safe and Sanitary Processing and Importing of Juice

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 recordkeeping requirements for applying hazard analysis and critical control point (HAACP) procedures for safe and sanitary processing for processors of fruit and vegetable juice.

DATES: Submit either electronic or written comments on the collection of information by September 13, 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.

Hazard Analysis and Critical Control Point (HACCP) Procedures for the Safe and Sanitary Processing and Importing of Juice—21 CFR Part 120 (OMB Control Number 0910–0466)—Extension

FDA's regulations in part 120 (21 CFR part 120) mandate the application of HACCP procedures to fruit and vegetable juice processing. HACCP is a preventative system of hazard control that can be used by all food processors to ensure the safety of their products to consumers. A HACCP system of preventive controls is the most effective and efficient way to ensure that these food products are safe. FDA's mandate to ensure the safety of the Nation's food supply is derived principally from the Federal Food, Drug, and Cosmetic Act (the act) (21 U.S.C. 321 *et seq.*). Under the act, FDA has authority to ensure that all foods in interstate commerce, or that have been shipped in interstate commerce, are not contaminated or otherwise adulterated, are produced and held under sanitary conditions, and are not misbranded or deceptively packaged; under section 701 (21 U.S.C. 371), the act authorizes the agency to issue regulations for its efficient enforcement. The agency also has authority under section 361 of the Public Health Service Act (42 U.S.C. 264) to issue and enforce regulations to prevent the introduction, transmission, or spread of communicable diseases from one State to another other State. Information development and recordkeeping are essential parts of any HACCP system. The information collection requirements are narrowly tailored to focus on the development of appropriate controls and document those aspects of processing that are critical to food safety. Through these regulations, FDA is implementing its authority under section 402(a)(4) of the act (21 U.S.C. 342(a)(4)).

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

TABLE 1.—ESTIMATED ANNUAL REPORTING BURDEN[1]

21 CFR Section	No of Recordkeepers	Annual Frequency per Recordkeeping	Total Annual Records	Hours Per Record	Total Hours
120.6(c) and 120.12(a)(1) and (b)	1,875	365	684,375	0.1	68,437.5
120.7; 120.10(a); and 120.12(a)(2), (b), and (c)	2,300	1.1	2,530	20	50,600