License Number: 16848N.
Name: eKKa Forwarding Inc.
Address: 223 Bergen Turnpike, Bldg. 3, Ridgefield Park, NJ 07660.
Date Revoked: December 15, 2001.
Reason: Failed to maintain a valid bond.

License Number: 4098F.
Name: Gaeli, Inc.
Address: 10050 NW 116th Way, Ste. 15, Medley, FL 33178.
Date Revoked: January 10, 2002.
Reason: Failed to maintain a valid bond.

License Number: 3656NF.
Name: Gulfstream Freight Services, Inc. dba Gulfstream Logistics.
Address: 11919 SW 130th Street, Miami, FL 33186.
Date Revoked: December 6, 2001.
Reason: Failed to maintain valid bonds.

License Number: 17090N.
Name: Inter-Connect Transportation, Inc.
Address: 8901 S. La Cienega Blvd., Ste. 210, Inglewood, CA 90301.
Date Revoked: December 8, 2001.
Reason: Failed to maintain a valid bond.

License Number: 3690NF.
Name: Inter-Freight Logistics, Inc.
Address: 5401 W. Kennedy Boulevard, Ste. 999, Tampa, FL 33609.
Date Revoked: December 28, 2001.
Reason: Surrendered license voluntarily.

License Number: 4335N.
Name: International Services, Inc.
Address: 2907 Empress Ct., Valrico, FL 33594.
Date Revoked: December 5, 2001.
Reason: Failed to maintain a valid bond.

License Number: 15712N.
Name: J.H.K. Transportation System, Inc.
Address: 5210 12th Street East, Ste. B, Fife, WA 98424.
Date Revoked: December 30, 2001.
Reason: Failed to maintain a valid bond.

License Number: 12473N.
Name: Jupiter Express, Inc.
Address: 156–19 76th Street, Howard Beach, NY 11414.
Date Revoked: December 8, 2001.
Reason: Failed to maintain a valid bond.

License Number: 4637F.
Name: Lion Cargo Brokers, Inc. dba Polaris Ocean Line.
Address: 8055 NW 77th Court, Ste. 3, Medley, FL 33166.
Date Revoked: January 2, 2002.
Reason: Failed to maintain a valid bond.

License Number: 4215F.
Name: Logistics Management International, Inc.
Address: 816 Thorndale Avenue, Bensenville, IL 60106.
Date Revoked November 7, 2001.
Reason: Failed to maintain a valid bond.

License Number: 4066N.
Name: Maracargo Inc.
Address: 7700 NW 79th Place, Ste. 1, Miami, FL 33166.
Date Revoked: November 4, 2001.
Reason: Failed to maintain a valid bond.

License Number: 3326N.
Name: Modern Cargo Services Inc.
Address: 11265 NW 131st Street, Medley, FL 33178.
Date Revoked: November 1, 2001.
Reason: Failed to maintain a valid bond.

License Number: 16462N.
Name: Multi Transport Inc.
Address: 8422 NW 66th Street, Miami, FL 33166.
Date Revoked: December 29, 2001.
Reason: Failed to maintain a valid bond.

License Number: 4592N.
Name: Natasha International Freight, Inc.
Address: 12912 SW 133 Court, Ste. A, Miami, FL 33186.
Date Revoked: December 8, 2001.
Reason: Failed to maintain a valid bond.

License Number: 12459N.
Name: PSD International, Inc.
Address: 220 W. Ivy Avenue, Inglewood, CA 90302.
Date Revoked: December 15, 2001.
Reason: Failed to maintain a valid bond.

License Number: 16083F.
Name: Palmetto Freight Forwarding Group.
Address: 9695 NW 79th Avenue, Bay 16, Hialeah, FL 33016.
Date Revoked: December 6, 2001.
Reason: Failed to maintain a valid bond.

License Number: 0690F.
Name: Robert E. Landweer & Co., Inc.
Address: 911 Western Avenue, Ste. 208, Seattle, WA 98104.
Date Revoked: December 16 2001.
Reason: Failed to maintain a valid bond.

License Number: 15682N.
Name: S/J Americas Service, LLC dba S/J Americas Service.
Address: 11821 I–H 10 East, Ste. 630, Houston, TX 77029.
Date Revoked: December 5, 2001.
Reason: Failed to maintain a valid bond.

License Number: 3406N.
Name: Simmons International Express, Inc.
Address: 101 E. Clarendon Street, Prospect Heights, IL 60070.
Date Revoked: January 4, 2002.
Reason: Failed to maintain a valid bond.

License Number: 4587NF.
Name: Toriello Passarelli, Inc. dba Toriello Freight International.
Address: 8611 NW 72nd Street, Miami, FL 33166.
Date Revoked: August 10, 2001.
Reason: Failed to maintain valid bonds.

License Number: 4511F.
Name: Total Logistic Control, LLC.
Address: 8300 Logistics Drive, Zeeland, MI 49464.
Date Revoked: January 6, 2002.
Reason: Failed to maintain a valid bond.

License Number: 4551F.
Name: Washington World Trading Corp. dba Washington World International Freight Forwarders.
Address: 10411 NW 28th Street, Ste. C–103, Miami, FL 33172.
Date Revoked: December 26, 2001.
Reason: Failed to maintain a valid bond.

Sandra L. Kusumoto,
Director, Bureau of Consumer Complaints and Licensing.

[FR Doc. 02–2273 Filed 1–30–02; 8:45 am]
BILLING CODE 6730–01–P

FEDERAL MARITIME COMMISSION

Ocean Transportation Intermediary License Reissuance

Notice is hereby given that the following Ocean Transportation Intermediary license has been reissued by the Federal Maritime Commission pursuant to section 19 of the Shipping Act of 1984, as amended by the Ocean Shipping Reform Act of 1998 (46 U.S.C. app. 1718) and the regulations of the Commission pertaining to the licensing of Ocean Transportation Intermediaries, 46 CFR 515.

License No.	Name/Address	Date reissuance
16838N	Webtrans Logistics, Inc. dba ANC International, 21136 S. Wilmington Avenue, #110, Carson, CA 90810.	December 19, 2001.

Sandra L. Kusumoto,

Director, Bureau of Consumer Complaints and Licensing.

[FR Doc. 02–2271 Filed 1–30–02; 8:45 am]

BILLING CODE 6730–01–P

FEDERAL MARITIME COMMISSION

Ocean Transportation Intermediary License; Applicant

Notice is hereby given that the following applicant has filed with the Federal Maritime Commission an application for license as Non-Vessel Operating Common Carrier and Ocean Freight Forwarder—Ocean Transportation Intermediary pursuant to section 19 of the Shipping Act of 1984 as amended (46 U.S.C. app. 1718 and 46 CFR 515).

Persons knowing of any reason why the following applicant should not receive a license are requested to contact the Office of Transportation Intermediaries, Federal Maritime Commission, Washington, DC 20573.

Non-Vessel Operating Common Carrier Ocean Transportation Intermediary Applicant:

Security Storage Company of Washington, 1701 Florida Avenue, NW., Washingnton, DC 20009–1697, Officers: Larry DePace, Senior Vice President (Qualifying Individual), Charles R. Lawrence, President/CEO.

Dated: January 25, 2002.

Bryant L. VanBrakle,

Secretary.

[FR Doc. 02–2272 Filed 1–30–02; 8:45 am]

BILLING CODE 6730–01–P

DEPARTMENT OF HEALTH AND HUMAN SERVICES

Food and Drug Administration

[Docket No. 01D–0584]

Draft ''Guidance for Industry: Use of Nucleic Acid Tests on Pooled Samples From Source Plasma Donors to Adequately and Appropriately Reduce the Risk of Transmission of HIV–1 and HCV''; Availability

AGENCY: Food and Drug Administration, HHS.

ACTION: Notice.

SUMMARY: The Food and Drug Administration (FDA) is announcing the availability of a draft document entitled ''Guidance for Industry: Use of Nucleic Acid Tests on Pooled Samples From Source Plasma Donors to Adequately and Appropriately Reduce the Risk of Transmission of HIV–1 and HCV'' dated December 2001. The draft guidance document, when finalized, would inform all establishments that manufacture Source Plasma that FDA has approved nucleic acid tests (NAT) to identify human immunodeficiency virus type 1 (HIV–1) and hepatitis C virus (HCV) in Source Plasma donations. The draft document recommends that manufacturers submit a prior approval supplement to a biologics license application (BLA) to implement HIV–1 and HCV NAT by a specified date.

DATES: Submit written or electronic comments on the draft guidance to ensure their adequate consideration in preparation of the final document by May 1, 2002. General comments on agency guidance documents are welcome at any time.

ADDRESSES: Submit written requests for single copies of the draft guidance to the Office of Communication, Training, and Manufacturers Assistance (HFM–40), Center for Biologics Evaluation and Research (CBER), Food and Drug Administration, 1401 Rockville Pike, Rockville, MD 20852–1448. Send one self-addressed adhesive label to assist the office in processing your requests. The document may also be obtained by mail by calling the CBER Voice Information System at 1–800–835–4709 or 301–827–1800, or by fax by calling the FAX Information System at 1–888–CBER–FAX or 301–827–3844. See the **SUPPLEMENTARY INFORMATION** section for electronic access to the draft guidance document.

Submit written comments on the document to the Dockets Management Branch (HFA–305), Food and Drug Administration, 5630 Fishers Lane, rm. 1061, Rockville, MD 20852. Submit electronic comments to http://www.fda.gov/dockets/ecomments.

FOR FURTHER INFORMATION CONTACT: Nathaniel L. Geary, Center for Biologics Evaluation and Research (HFM–17), Food and Drug Administration, 1401 Rockville Pike, Rockville, MD 20852–1448, 301–827–6210.

SUPPLEMENTARY INFORMATION:

I. Background

FDA is announcing the availability of a draft document entitled ''Guidance for Industry: Use of Nucleic Acid Tests on Pooled Samples From Source Plasma Donors to Adequately and Appropriately Reduce the Risk of Transmission of HIV–1 and HCV'' dated December 2001. FDA's final rule (66 FR 31146, June 11, 2001) entitled ''Requirements for Testing Human Blood Donors for Evidence of Infection Due to Communicable Diseases'' became effective on December 10, 2001. The provision in 21 CFR 610.40(b) of the rule provides that manufacturers ''must perform one or more screening tests to adequately and appropriately reduce the risk of transmission of communicable disease agents'' (66 FR 31146 at 31162). As we noted in the preamble to the final rule, the standard for adequate and appropriate testing will change as new testing technology is approved by FDA. We explained, ''we intend to regularly issue guidance describing those tests that we believe would adequately and appropriately reduce the risk of transmission of communicable disease agents'' (66 FR 31146 at 31149).

The availability of NAT to identify HIV–1 and HCV will change the testing protocol that should be used to adequately and appropriately reduce the risk of transmission of those diseases. The draft document recommends that manufacturers submit a prior approval supplement to a BLA to implement HIV–1 and HCV NAT by a specified date.

This draft guidance is being issued consistent with FDA's good guidance practices regulation (21 CFR 10.115). This draft guidance document represents the agency's current thinking on this topic. It does not create or confer any rights for or on any person and does not operate to bind FDA or the public. An alternative approach may be used if such approach satisfies the requirement of the applicable statutes and regulations.

II. Comments

This draft document is being distributed for comment purposes only and is not intended for implementation at this time. Interested persons may submit to the Dockets Management Branch (address above) written or electronic comments regarding this draft guidance document. Submit written or electronic comments to ensure adequate consideration in preparation of the final document by May 1, 2002. Two copies of any comments are to be submitted, except individuals may submit one copy. Comments should be identified with the docket number found in the brackets in the heading of this document. A copy of the document and received comments are available for public examination in the Dockets Management Branch between 9 a.m. and 4 p.m., Monday through Friday.

III. Electronic Access

Persons with access to the Internet may obtain the document at either http://www.fda.gov/cber/guidelines.htm or http://www.fda.gov/ohrms/dockets/default.htm.