Synopsis: The amendment adds Lykes and MacAndrews as parties to the agreement.

Agreement No.: 011701–007.

Title: Pacific East Coast Express Agreement.

Parties: China Shipping Container Lines Co., Ltd.; CMA CGM, S.A.; P&O Nedlloyd B.V.; and P&O Nedlloyd Limited;

Filing Party: Paul M. Keane, Esq., Cichanowicz, Callan, Keane, Vengrow & Textor, LLP; 61 Broadway, Suite 3000; New York, NY 10006–2802.

Synopsis: The modification reflects that one of the vessels contributed by CMA under the agreement will now be provided by CMA's wholly-owned subsidiary ANL Singapore Pte Ltd. The parties request expedited review.

Agreement No.: 011733–012.

Title: Common Ocean Carrier Platform Agreement

Parties: A.P. Moller-Maersk A/S, P&O Nedlloyd Limited, Hamburg-Süd, Mediterranean Shipping Company S.A., CMA CGM S.A., Hapag Lloyd Container Linie GmbH, and United Arab Shipping Company (SAG), as shareholder parties, and Alianca Navegacao e Logistica Ltda., Safmarine Container Lines N.V., Nippon Yusen Kaisha, CP Ship Limited, Tasman Orient Line C.V., Mitsui O.S.K. lines Ltd., Lykes Lines Limited LLC, Kawasaki Kisen Kaisha Ltd., and FESCO Ocean Management Ltd. as non-shareholder parties.

Filing Party: Wayne R. Rohde, Esq.; Sher & Blackwell; 1850 M Street, NW., Suite 900; Washington, DC 20036.

Synopsis: The amendment adds FESCO Ocean Management Ltd. as a non-shareholder party to the agreement.

Dated: August 6, 2004.

By Order of the Federal Maritime Commission.

Bryant L. VanBrakle,

Secretary.

[FR Doc. 04–18375 Filed 8–10–04; 8:45 am]

BILLING CODE 6730–01–P

FEDERAL MARITIME COMMISSION

[Docket No. 04–08]

Qin's, Incorporated v. Superior Link International, Inc.; Notice of Filing of Complaint and Assignment

Notice is given that a complaint has been filed by Qin's, Incorporated (''Complainant'') against Superior Link International, Inc. (''Respondent''). Complainant contends that Respondent has unreasonably refused to deal or negotiate the proper release of Complainant's two containers in violation of section 10(b)(10)[1] of the Shipping Act of 1984, 46 U.S.C. app. section 1709(b)(10). As a direct result of these allegations, Complainant claims that it has suffered substantial economic damages and injury valued at $23,626.40. Complainant seeks an order directing Respondent to pay reparations, court costs, attorneys fees, and any further relief as the Commission may determine to be warranted.

This proceeding has been assigned to the Office of Administrative Law Judges. A 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 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 August 4, 2005 and a final decision of the Commission shall be issued by December 2, 2005.

Bryant L. VanBrakle,

Secretary.

[FR Doc. 04–18377 Filed 8–10–04; 8:45 am]

BILLING CODE 6730–01–P

FEDERAL MARITIME COMMISSION

Ocean Transportation Intermediary License Reissuances

Notice is hereby given that the following Ocean Transportation Intermediary licenses have 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 part 515.

License No.	Name/address	Date reissued
017958NF	DLM Venturesd, Inc., 1850 NW 84th Avenue, Miami, FL 33126	July 17, 2004.
011157N	The Norton Line Inc., 249 E. Ocean Blvd., Suite 620, Long Beach, CA 90802	July 10, 2004.

Ronald D. Murphy,

Deputy Director, Bureau of Consumer Complaints and Licensing.

[FR Doc. 04–18376 Filed 8–10–04; 8:45 am]

BILLING CODE 6730–01–P

DEPARTMENT OF HEALTH AND HUMAN SERVICES

Centers for Disease Control and Prevention

[60Day–04–04JW]

Proposed Data Collections Submitted for Public Comment and Recommendations

In compliance with the requirement of section 3506(c)(2)(A) of the Paperwork Reduction Act of 1995 for opportunity for public comment on proposed data collection projects, the Centers for Disease Control and Prevention (CDC) will publish periodic summaries of proposed projects. To request more information on the proposed projects or to obtain a copy of the data collection plans and instruments, call the CDC Reports Clearance Officer on (404) 498–1210 or send an e-mail to *omb@cdc.gov*.

CDC is requesting an emergency clearance for this data collection with a two week public comment period. CDC is requesting OMB approval of this

[1] Complainant cites section 10(b)(7) of the Shipping Act of 1984, 46 U.S.C. app. section 1709(b)(7), however, Complainant's narrative statement refers to a 10(b)(10) violation.

package 7 days after the end of the public comment period. While there is currently no SARS outbreak, expedited clearance is needed because the season when SARS outbreaks are most likely to occur is quickly approaching.

Comments are invited on: (a) Whether the proposed collection of information is necessary for the proper performance of the functions of the agency, including whether the information shall have practical utility; (b) the accuracy of the agency's estimate of the burden of the proposed collection of information; (c) ways to enhance the quality, utility, and clarity of the information to be collected; and (d) ways to minimize the burden of the collection of information on respondents, including through the use of automated collection techniques or other forms of information technology. Send comments to Seleda M. Perryman, CDC Assistant Reports Clearance Officer, 1600 Clifton Road, MS–E11, Atlanta, GA 30333. Written comments should be received within 14 days of this notice.

Proposed Project

SARS Laboratory Safety Survey—New—National Center for Infectious Diseases (NCID), Centers for Disease Control and Prevention (CDC).

During the past year, incidents of lab-acquired SARS–CoV infection have occurred at research institutions in China, Singapore, and Taiwan. As part of measures taken by CDC to prevent lab-acquired SARS infections in the United States, the Respiratory and Enteric Virus Branch (REVB) would like to collect information on current biosafety practices and use of live SARS–CoV from U.S. institutions that have obtained the virus, either from CDC or from other sources.

Response data will be used to determine to what extent research involving live SARS–CoV and other biosafety level 3 pathogens is being conducted, the safety of the research environment, information about biosafety training, and the level of lab worker experience with BSL–3 pathogens. Expedited collection of this information is necessary to ascertain the safety of environments where live SARS–CoV research may be taking place and to potentially address any deficiencies that may exist. There is no cost to the respondents.

Form	Respondent	No. of respondents	No. of responses/respondent	Avg. burden/response (in hrs.)	Total burden hours
Safety Survey for Laboratories Conducting Research with live SARS–CoV and Other BSL–3 Pathogens.	Laboratorians	120	1	10/60	20
Total	.....	.....	.....	.....	20

Dated: August 4, 2004.

Alvin Hall,

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

[FR Doc. 04–18331 Filed 8–10–04; 8:45 am]

BILLING CODE 4163–18–P

DEPARTMENT OF HEALTH AND HUMAN SERVICES

Centers for Disease Control and Prevention

[60Day–04–0497]

Proposed Data Collections Submitted for Public Comment and Recommendations

In compliance with the requirement of section 3506(c)(2)(A) of the Paperwork Reduction Act of 1995 for opportunity for public comment on proposed data collection projects, the Centers for Disease Control and Prevention (CDC) will publish periodic summaries of proposed projects. To request more information on the proposed projects or to obtain a copy of the data collection plans and instruments, call the CDC Reports Clearance Officer on (404) 498–1210.

Comments are invited on: (a) Whether the proposed collection of information is necessary for the proper performance of the functions of the agency, including whether the information shall have practical utility; (b) the accuracy of the agency's estimate of the burden of the proposed collection of information; (c) ways to enhance the quality, utility, and clarity of the information to be collected; and (d) ways to minimize the burden of the collection of information on respondents, including through the use of automated collection techniques or other forms of information technology. Send comments to Sandi Gambescia, CDC Assistant Reports Clearance Officer, 1600 Clifton Road, MS–E11, Atlanta, GA 30333 or send an e-mail to *omb@cdc.gov.* Written comments should be received within 60 days of this notice.

Proposed Project

Evaluating CDC Funded Health Department HIV Prevention Programs, Partner Counseling and Referral Services, OMB No. 0920–0497—Extension—National Center for HIV, STD, and TB Prevention (NCHSTP), Centers for Disease Control and Prevention (CDC).

Background

CDC is requesting approval for the continued use of two currently approved forms under OMB No. 0920–0497, for collecting HIV partner counseling and referral services (PCRS) program data. The current forms expire October 31, 2004. This request is for a 12-month clearance past this date. The extension of the current forms will allow grantees to continue to collect PCRS data as they transition to the new Program Evaluation and Monitoring System (PEMS) over the next year.

CDC funds HIV prevention projects in 65 public health agencies (50 states, 6 cities, 7 territories, Washington, DC, and Puerto Rico) through cooperative agreements. PCRS is one of a number of public health strategies supported by CDC that is designed to control and prevent the spread of HIV.

A fundamental feature of PCRS is informing current and past partners of an HIV-infected person that they have been identified as a sex or injection-drug-paraphernalia-sharing partner, and advising them to be tested for HIV. Informing partners of their exposure to HIV is confidential, and partners are not told who reported their name, or when the reported exposure occurred. Notified partners, who may not have suspected their risk, can choose whether to have HIV counseling and testing. Those who choose to be tested and are found to be HIV positive can receive a medical evaluation, treatment, and prevention services designed to modify their high risk behavior, thereby possibly reducing the number of new HIV infections.

HIV prevention programs that conduct PCRS interventions can reach significant numbers of persons at very high risk of contracting HIV. The CDC requires aggregate PCRS program data to determine if interventions are being