be filed within 10 days after the time for filing oppositions has expired.

Subject: Advanced Television Systems and Their Impact Upon the Existing Television Broadcast Service. (MM Docket No. 87–268).

Number of Petitions Filed: 220.

Federal Communications Commission.

William F. Caton,

Acting Secretary. [FR Doc. 97–17406 Filed 7–2–97; 8:45 am] BILLING CODE 6712–01–M

FEDERAL MARITIME COMMISSION

Notice of Agreement(s) Filed

The Commission hereby gives notice of the filing of the following agreement(s) under the Shipping Act of 1984.

Interested parties can review or obtain copies of agreements at the Washington, DC offices of the Commission, 800 North Capitol Street, N.W., Room 962. Interested parties may submit comments on an agreement to the Secretary, Federal Maritime Commission, Washington, DC 20573, within 10 days of the date this notice appears in the **Federal Register**.

Agreement No.: 232–011507–003. Title: Di Gregorio-Tricon Agreement. Parties: Di Gregorio-Navegacao Ltda., DSR-Senator Lines, Cho Yang Shipping Col., Ltd.

Synopsis: The proposed modification extends the minimum term of the Agreement to September 1, 2000, and provides that any party may withdraw from the Agreement upon six months' written notice, but such notice may not commence to run until 30 months following September 1, 1997.

Agreement No.: 224–201028.

Title: Port of Oakland/Stevedoring Services of America Terminal Agreement.

Parties: Port of Oakland ("Port"), Stevedoring Services of America ("SSA").

Synopsis: Under the terms of the Agreement, the Port assigns to SSA a preferential right to manage, operate, and solicit cargo at the Port's Charles B. Howard Terminal. The initial term of the Agreement will be from July 1, 1997, to June 30, 2007.

Dated: June 30, 1997.

By order of the Federal Maritime Commission.

Secretary.

[FR Doc. 97–17482 Filed 7–2–97; 8:45 am] BILLING CODE 6730–01–M

DEPARTMENT OF HEALTH AND HUMAN SERVICES

Office of the Secretary

Agency Information Collection Activities: Submission for OMB Review; Comment request

The Department of Health and Human Services, Office of the Secretary publishes a list of information collections it has submitted to the Office of Management and Budget (OMB) for clearance in compliance with the Paperwork Reduction Act of 1995 (44 U.S.C. Chapter 35) and 5 CFR 1320.5. The following are those information collections recently submitted to OMB.

1. OIG Advisory Opinion Procedures and Preliminary Questions

Section 205 of the Health Insurance Portability and Accountability Act of 1996, Pub. Law 104-191, requires the Department to provide advisory opinions to the public regarding several categories of subject matter, including the requestor's potential liability under sections 1128, 1128A, and 1128B of the Act. The OIG has separately published in the Federal Register (2/19/97) and interim final regulation providing the procedure under which members of the public may request advisory opinions from the OIG. That discussion contains a more thorough discussion of the advisory opinion procedures and the preliminary questions. The procedures in the interim final rule include several provisions for the collection of information. In addition, in order to aid potential requestors and the OIG in providing opinions under this process, the OIG is providing preliminary questions that may be answered in an advisory opinion request. These preliminary questions will be voluntary and will correspond with each sanction provision about which advisory opinions will be rendered. Responsents: Health care providers; Annaul Number of Respondents: 500; Average Burden per Response: 10 hours; Total Annual Burden on Respondents: 5000 hours; Cost Burden: \$1,000,000.

OMB Desk Officer: Allison Eydt. Copies of the information collection packages listed above can be obtained by calling the OS Reports Clearance Officer on (202) 690–6207. Written comments and recommendations for the proposed information collection should be sent directly to the OMB desk officer designated above at the following address:

Human Resources and Housing Branch, Office of Management and Budget, New Executive Office Building, Room 10235, 725 17th Street N.W., Washington, D.C. 20503.

Comments may also be sent to Cynthia Agens Bauer, OS Reports Clearance Officer, Room 503H, Humphrey Building, 200 Independence Avenue S.W., Washington, DC, 20201. Written comments should be received within 30 days of this notice.

Dated: June 24, 1997.

Dennis P. Williams,

Deputy Assistant Secretary, Budget. [FR Doc. 97–17411 Filed 7–2–97; 8:45 am] BILLING CODE 4150–04–M

DEPARTMENT OF HEALTH AND HUMAN SERVICES

National Committee on Vital and Health Statistics: Meetings

Pursuant to the Federal Advisory Committee Act, the Department of Health and Human Services announces the following advisory committee meetings.

Name: National Committee on Vital and Health Statistics (NCVHS), Subcommittee on Population-Specific Issues.

Times and Dates: 9:00 a.m.–5:00 p.m., July 21, 1997.

Place: Room 337A, Hubert H. Humphrey Building, 200 Independence Avenue, SW, Washington, D.C. 20201.

Status: Open.

Purpose: On July 21, the Subcommittee on Population-Specific Issues of the National Committee on Vital and Health Statistics will meet to formulate its work plan relating to data needed to assess the impact of Medicaid Managed Care on Medicaid beneficiaries. Presentations are tentatively planned on the following topics: the scope of current activities, anticipated work across the agencies within the Department, and plans for monitoring and evaluating the impact of managed care on Medicaid beneficiaries, including child health and mental health. The Subcommittee plans to formulate specific questions to be addressed and resources needed, as well as methods to addressing this priority, including the possibility of public hearings.

Contact Person for More Information: Substantive program information as well as summaries of the meeting and roster of committee members may be obtained from Carolyn Rimes, Lead Staff for the Subcommittee, Health Care Financing Administration, 7500 Security Boulevard, Baltimore Maryland 21244, telephone (410) 786–6620, or Marjorie S. Greenberg, Executive Secretary, NCVHS, NCHS, CDC, Room 1100, Presidential Building, 6525 Belcrest Road, Hyattsville, Maryland 20782, telephone (301) 436–7050. Information also is available on the NCVHS home page of the HHS website: http://aspe.os.dhhs.gov/ncvhs. Dated: June 27, 1997. James Scanlon, Director, Division of Data Policy, Office of the Assistant Secretary for Planning and Evaluation. [FR Doc. 97–17410 Filed 7–2–97; 8:45 am] BILLING CODE 4151–04–M

DEPARTMENT OF HEALTH AND HUMAN SERVICES

Health Care Financing Administration

[HCFA-685, and HCFA-684 A-J]

Agency Information Collection Activities: Proposed Collection; Comment Request

In compliance with the requirement of section 3506(c)(2)(A) of the Paperwork Reduction Act of 1995, the Health Care Financing Administration (HCFA), Department of Health and Human Services, is publishing the following summary of proposed collections for public comment. Interested persons are invited to send comments regarding the burden estimate or any other aspect of this collection of information, including any of the following subjects: (1) The necessity and utility of the proposed information collection for the proper performance of the agency's functions; (2) the accuracy of the estimated burden; (3) ways to enhance the quality, utility, and clarity of the information to be collected; and (4) the use of automated collection techniques or other forms of information technology to minimize the information collection burden.

1. Type of Information Collection *Request:* Extension of a currently approved collection; Title of Information Collection: End Stage Renal Disease (ESRD) Network Semi-Annual Cost Report Forms and Supporting Regulations 42 CFR 405.2110 and 405.2112; Form No.: HCFA-685; Use: The Semi-annual cost report enables HCFA to review specific Network costs, compare costs between Networks, and project future Network costs. The reports are also used as an early warning system to determine if a Network is in danger of exceeding the total cost of its contract. Frequency: Semi-annually; Affected Public: Not-for-profit institutions; Number of Respondents: 18; Total Annual Responses: 36; Total Annual Hours: 108.

2. Type of Information Collection Request: Extension of a currently approved collection; Title of Information Collection: End Stage Renal Disease Network (ESRD) Business Proposal Forms; Form No.: HCFA–684 through 684 A–J; *Use:* Current End Stage Renal Disease (ESRD) Networks and other bidders are required to submit contract proposals to participate as a HCFA sanctioned ESRD Network. The business proposal forms are used to satisfy HCFA's need for consistent, meaningful, and verifiable data to evaluate contract proposals. *Frequency:* Every three years; *Affected Public:* Notfor-profit institutions; *Number of Respondents:* 18; *Total Annual Responses:* 36; *Total Annual Hours:* 1,080.

To obtain copies of the supporting statement for the proposed paperwork collections referenced above, access HCFA's Web Site Address at http:// www.hcfa.gov/regs/prdact95.htm, or to obtain the supporting statement and any related forms, E-mail your request, including your address and phone number, to Paperwork@hcfa.gov, or call the Reports Clearance Office on (410) 786-1326. Written comments and recommendations for the proposed information collections must be mailed within 60 days of this notice directly to the HCFA Paperwork Clearance Officer designated at the following address: HCFA, Office of Financial and Human Resources, Management Analysis and Planning Staff, Attention: John Rudolph, Room C2-26-17, 7500 Security Boulevard, Baltimore, Maryland 21244-1850.

Dated: June 26, 1997.

Edwin J. Glatzel,

Director, Management Analysis and Planning Staff, Office of Financial and Human Resources.

[FR Doc. 97–17434 Filed 7–2–97; 8:45 am] BILLING CODE 4120–03–P

DEPARTMENT OF HEALTH AND HUMAN SERVICES

Substance Abuse and Mental Health Services Administration

Current List of Laboratories Which Meet Minimum Standards, To Engage in Urine Drug Testing for Federal Agencies, and Laboratories That Have Withdrawn From the Program

AGENCY: Substance Abuse and Mental Health Services Administration, HHS (Formerly: National Institute on Drug Abuse, ADAMHA, HHS). **ACTION:** Notice.

SUMMARY: The Department of Health and Human Services notifies Federal agencies of the laboratories currently certified to meet standards of Subpart C of Mandatory Guidelines for Federal Workplace Drug Testing Programs (59 FR 29916, 29925). A similar notice listing all currently certified laboratories will be published during the first week of each month, and updated to include laboratories which subsequently apply for and complete the certification process. If any listed laboratory's certification is totally suspended or revoked, the laboratory will be omitted from updated lists until such time as it is restored to full certification under the Guidelines.

If any laboratory has withdrawn from the National Laboratory Certification Program during the past month, it will be identified as such at the end of the current list of certified laboratories, and will be omitted from the monthly listing thereafter.

This Notice is now available on the internet at the following website: http://www.health.org

FOR FURTHER INFORMATION CONTACT: Mrs. Giselle Hersh or Dr. Walter Vogl, Division of Workplace Programs, Room 13A–54, 5600 Fishers Lane, Rockville, Maryland 20857; Tel.: (301) 443–6014.

SUPPLEMENTARY INFORMATION:

Mandatory Guidelines for Federal Workplace Drug Testing were developed in accordance with Executive Order 12564 and section 503 of Pub. L. 100-71. Subpart C of the Guidelines, "Certification of Laboratories Engaged in Urine Drug Testing for Federal Agencies," sets strict standards which laboratories must meet in order to conduct urine drug testing for Federal agencies. To become certified an applicant laboratory must undergo three rounds of performance testing plus an on-site inspection. To maintain that certification a laboratory must participate in a quarterly performance testing program plus periodic, on-site inspections.

Laboratories which claim to be in the applicant stage of certification are not to be considered as meeting the minimum requirements expressed in the HHS Guidelines. A laboratory must have its letter of certification from SAMHSA, HHS (formerly: HHS/NIDA) which attests that it has met minimum standards.

In accordance with Subpart C of the Guidelines, the following laboratories meet the minimum standards set forth in the Guidelines:

- ACL Laboratory, 8901 W. Lincoln Ave., West Allis, WI 53227, 414–328–7875 (formerly: Bayshore Clinical Laboratory)
- Aegis Analytical Laboratories, Inc., 345 Hill Ave., Nashville, TN 37210, 615–255–2400
- Alabama Reference Laboratories, Inc., 543 South Hull St., Montgomery, AL 36103, 800–541–4931/334–263–5745