FEMA system of records may be disclosed as a routine use to a Federal, State, or local agency maintaining civil, criminal, regulatory, licensing or other enforcement information or other pertinent information, such as current licenses, if necessary, to obtain information relevant to an agency decision concerning hiring or retention of an employee, issuance of a security clearance, letting of a contract, or issuance of a license, grant, or other benefit.

3. *Routine Use—Disclosure of Requested Information:* A record from a FEMA system of records may be disclosed to a Federal agency, in response to a written request in connection with hiring or retention of an employee, issuance of an investigation of an employee, letting of a contract, or issuance of a license, grant, or other benefit by the requesting agency, to the extent that the information is relevant and necessary to the requesting agency's decision on the matter.

4. *Routine Use—Grievance, Complaint, Appeal:* A record from a FEMA system of records may be disclosed to an authorized appeal or grievance examiner, formal complaints examiner, equal employment opportunity investigator, arbitrator, or other duly authorized official engaged in investigation or settlement of a grievance, complaint, or appeal filed by an employee. A record from this system of records may be disclosed to the Office of Personnel Management in accordance with that agency's responsibility for evaluation of Federal personnel management.

To the extent that official personnel records in the custody of FEMA are covered within systems of records published by the Office of Personnel Management as governmentwide records, those records will be considered as a part of that governmentwide system. Other official personnel records covered by notices published by FEMA and considered to be separate systems of records may be transferred to the Office of Personnel Management in accordance with official personnel programs and activities as a routine use.

5. *Routine Use—Congressional Inquiries:* A record from a FEMA system of records may be disclosed as a routine use to a Member of Congress or to a Congressional staff member in response to an inquiry of the Congressional office made at the direct, written request of the individual about whom the record is maintained.

6. *Routine Use—Private Relief Legislation:* The information contained in a FEMA system of records may be disclosed as a routine use to the Office of Management and Budget in connection with the review of private relief legislation as set forth in OMB Circular No. A-19 at any stage of the legislative coordination and clearance process as set forth in that circular.

7. *Routine Use—Disclosure to the Office of Personnel Management:* A record from a FEMA system of records may be disclosed to the Office of Personnel Management concerning information on pay and leave benefits, retirement deductions, and any other information concerning personnel actions.

8. *Routine Use—Disclosure to National Archives and Records Administration:* A record from a FEMA system of records may be disclosed as a routine use to the National Archives and Records Administration in records management inspections conducted under authority of 44 U.S.C. 2904 and 2906.

9. *Routine Use—Grand Jury:* A record from any system of records may be disclosed, as a routine use, to a grand jury agent pursuant to a Federal or State grand jury subpoena or to a prosecution request that such record be released for the purpose of its introduction to a grand jury.

Appendix AA

Addresses for FEMA Regional Offices:

Region I—Regional Director, Federal Emergency Management Agency, room 442, J.W. McCormack Post Office and Courthouse Building, Boston, MA 02109–4595;

Region II—Regional Director, Federal Emergency Management Agency, 26 Federal Plaza, room 1338, New York, NY 10278–0002;

Region III—Regional Director, Federal Emergency Management Agency, Liberty Square Building (Second Floor), 105 South Seventh Street, Philadelphia, PA 19106–3316;

Region IV—Regional Director, Federal Emergency Management Agency, 1371 Peachtree Street, NE., suite 700, Atlanta, GA 30309–3108;

Region V—Regional Director, Federal Emergency Management Agency, 175 West Jackson Blvd., 4th Floor, Chicago, IL 60604–2698;

Region VI—Regional Director, Federal Emergency Management Agency, Federal Regional Center, 800 North Loop 288, Denton, TX 76201-3698;

Region VII—Regional Director, Federal Emergency Management Agency, 2323 Grand Boulevard, room 900, Kansas City, MO 64108–2670;

Region VIII—Regional Director, Federal Emergency Management Agency, Denver Federal Center, Building 710, Box 25267, Denver, CO 80225–0267;

Region IX—Regional Director, Federal Emergency Management Agency, Building 105, Presidio of San Francisco, CA 94129–1250;

Region X—Regional Director, Federal Emergency Management Agency, Federal Regional Center, 130 228th Street SW., Bothell, WA 98021–9796.

Dated: September 10, 1996.

John P. Carey,

General Counsel.

[FR Doc. 96–24320 Filed 9–20–96; 8:45 am]

BILLING CODE 6718–01–P

FEDERAL MARITIME COMMISSION

Notice of Agreement(s) Filed

The Federal Maritime Commission hereby gives notice of the filing of the following agreement(s) pursuant to section 5 of the Shipping Act of 1984.

Interested parties may inspect and obtain a copy of each agreement at the Washington, D.C. Office of the Federal Maritime Commission, 800 North Capitol Street, N.W., 9th Floor. Interested parties may submit comments on each agreement to the Secretary, Federal Maritime Commission, Washington, D.C. 20573, within 10 days after the date of the Federal Register in which this notice appears. The requirements for comments are found in § 572.603 of Title 46 of the Code of Federal Regulations. Interested persons should consult this section before communicating with the Commission regarding a pending agreement.

Agreement No.: 224–200005–007, 008, 009 & 010.

Title: Port Authority of New York & New Jersey/Maher Terminals Lease Agreement.

Parties: Port Authority of New York & New Jersey Maher Terminals, Inc.

Synopsis: The modifications provide for the substitution of certain open areas for Maher's Fleet Street Terminal, the change of definitions regarding certain ''qualified containers'', the construction of certain berth areas, and the change of definitions regarding certain charges and disposal costs.

Dated: September 17, 1996.

By Order of the Federal Maritime Commission.

Joseph C. Polking,

Secretary.

[FR Doc. 96–24260 Filed 9–20–96; 8:45 am]

BILLING CODE 6730–01–M

DEPARTMENT OF HEALTH AND HUMAN SERVICES

Administration for Children and Families

Proposed Information Collection Activity; Comment Request Proposed Projects

Title: Runaway and Homeless Youth Management Information System.

OMB No.: 0970–0123.

Description: In the runaway and home less Youth Act (42 U.S.C. 5701 *et seq.*) Congress mandated that the Department of Health and Human Services (HHS) report regularly on the status of HHS-funded programs serving runaway and homeless youth. In the Anti-Drug Abuse Act of 1988 (42 U.S.C. 11801 *et seq.*) Congress mandated that HHS report regularly on the status of HHS-funded Drug Abuse and Prevention Programs (DAPP) serving runaway and homeless youth. Organizations funded under the Runaway and Homeless Youth Program and/or Drug Abuse and Prevention Program are required by statute (42 U.S.C. 5712, 42 U.S.C. 5714–2 and/or 42

U.S.C. 11824) to meet several data collection and reporting requirements, including maintaining client statistical records and submitting annual program reports with regard to the profile of youth and families served and the services provided to them. The RHY MIS data support these organizations as they carry out a variety of integrated, ongoing responsibilities and projects, including legislative reporting requirements, planning and public policy development for runaway and homeless youth programs, accountability monitoring, program management, research, and evaluation.

Respondents: Runaway and Homeless Youth Grantees and Drug Abuse and Prevention Program Grantees.

Annual Burden Estimates:

Instrument	No. of respondent	No. of responses per respondent	Average burden hours per response	Total burden hours
Youth Program status	400	4	2.2	3,466.67
Youth profile	400	4	29.1	46,501.00
Agency profile	400	1	0.17	66.67
Program profile	400	1	1.0	400.00
Staff profile	400	1	1.2	466.67
Coordinating agency	400	1	0.3	133.33
community education	400	1	0.4	166.67
Promotional/instructional materials	400	1	0.2	66.67
Estimated total annual burden hours	.....	.....	.....	51,267.67

In compliance with the requirements of Section 3506(c)(2)(A) of the Paperwork Reduction act of 1995, the Administration for Children and Families is soliciting public comment on the specific aspects of the information collection described above. Copies of the proposed collection of information can be obtained and comments may be forwarded by writing to the Administration for Children and Families, Office of Information Services, Division of Information Resource management Services, 370 L'Enfant Promenade, S.W., Washington, DC 20447, Attn: ACF Reports Clearance Officer. All requests should be identified by the title of the information collection.

The Department specifically requests comments 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 propose 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. Consideration will be given to comments and suggestions submitted within 60 days of this publication.

Dated: September 17, 1996.

Bob Sargis,

Acting Reports Clearance Officer.

[FR Doc. 96–24226 Filed 9–20–96; 8:45 am]

BILLING CODE 4184–01–M

Food and Drug Administration

[Docket No. 96M–0332]

Neopath, Inc.; Premarket Approval of the AutoPap® 300 QC System

AGENCY: Food and Drug Administration, HHS.

ACTION: Notice.

SUMMARY: The Food and Drug Administration (FDA) is announcing its approval of the application by Neopath, Inc., Redmond, WA, for premarket approval, under the Federal Food, Drug, and Cosmetic Act (the act), of the AutoPap® 300 QC System. After reviewing the recommendation of the Hematology and Pathology Devices Panel, FDA's Center for Devices and Radiological Health (CDRH) notified the applicant, by letter of September 29, 1995, of the approval of the application.

DATES: Petitions for administrative review by October 23, 1996.

ADDRESSES: Written requests for copies of the summary of safety and effectiveness data and petitions for administrative review to the Dockets Management Branch (HFA–305), Food and Drug Administration, 12420 Parklawn Dr., rm. 1–23, Rockville, MD 20857.

FOR FURTHER INFORMATION CONTACT: Peter E. Maxim, Center for Devices and Radiological Health (HFZ–440), Food and Drug Administration, 2098 Gaither Rd., Rockville, MD 20850, 301–594–1293.

SUPPLEMENTARY INFORMATION: On February 24, 1995, Neopath, Inc., Redmond, WA 98052, submitted to CDRH an application for premarket approval of the AutoPap® 300 QC System. The device is an automated cervical cytology screening device intended for use in the quality control and rescreening of previously screened Papanicolaou (Pap) smear slides. The AutoPap® 300 QC System is to be used only on conventionally prepared Pap smear slides that have been previously classified as within normal limits (WNL) and satisfactory for interpretation by a screening cytologist. The AutoPap® 300 QC System is not intended to replace the current laboratory slide review processes referred to as ''high risk rescreen.''

On August 8, 1995, the Hematology and Pathology Devices Panel of the Medical Devices Advisory Committee, an FDA advisory committee, reviewed and recommended approval of the application. On September 29, 1995, CDRH approved the application by a letter to the applicant from the Director of the Office of Device Evaluation, CDRH.

A summary of the safety and effectiveness data on which CDRH based its approval is on file in the Dockets Management Branch (address above) and is available from that office upon written request. Requests should be identified with the name of the device and the docket number found in brackets in the heading of this document.

Opportunity for Administrative Review

Section 515(d)(3) of the act (21 U.S.C. 360e(d)(3)) authorizes any interested person to petition, under section 515(g) of the act, for administrative review of CDRH's decision to approve this application. A petitioner may request either a formal hearing under part 12 (21 CFR part 12) of FDA's administrative practices and procedures regulations or a review of the application and CDRH's action by an independent advisory