Name of harbor or creek	Number of vessels
Lake Montauk	883
Total	1711

*No data available.

Information regarding vessel population based on length shows that 63% of the boats are less than 40 feet and 37% of the vessels are 40 feet or greater in length. These percentages are based on a survey of overnight and long term occupancy and omitted marinas with recreational small crafts. Based on the number and size of boats, and using various methods to estimate the number of holding tanks, it is estimated that 5 to 8 pumpouts are needed to service the vessel population in the NDAs. Currently, ten pumpouts and three pumpout boats exist in the NDAs.

The EPA hereby makes a final affirmative determination that adequate facilities for the safe and sanitary removal and treatment of sewage from all vessels are reasonably available for Northwest Creek, Three Mile Harbor, Hog Creek, Accabonac Harbor, Napeague Harbor and Lake Montauk in the Town of East Hampton, New York. This final affirmative determination will result in a New York State prohibition of any sewage discharges from vessels in Northwest Creek, Three Mile Harbor, Hog Creek, Accabonac Harbor, Napeague Harbor and Lake Montauk.

Dated: January 28, 1999.

William J. Muszynski,

Acting Regional Administrator, Region 2.

[FR Doc. 99–3518 Filed 2–11–99; 8:45 am]

BILLING CODE 6560–50–P

FARM CREDIT ADMINISTRATION

Sunshine Act Meeting

Farm Credit Administration Board; Regular Meeting

AGENCY: Farm Credit Administration.

SUMMARY: Notice is hereby given, pursuant to the Government in the Sunshine Act (5 U.S.C. 552b(e)(3)), that the March 11, 1999 regular meeting of the Farm Credit Administration Board (Board) will not be held. The Board will hold a special meeting at 2:00 p.m. on Tuesday, March 23, 1999. An agenda for that meeting will be forthcoming.

FOR FURTHER INFORMATION CONTACT: Vivian L. Portis, Secretary to the Farm Credit Administration Board, (703) 883–4025, TDD (703) 883–4444.

ADDRESSES: Farm Credit Administration, 1501 Farm Credit Drive, McLean, Virginia 22102–5090.

Dated: February 9, 1999.

Vivian L. Portis,

Secretary,

Farm Credit Administration Board.

[FR Doc. 99–3697 Filed 2–10–99; 3:22 pm]

BILLING CODE 6705–01–P

FEDERAL MARITIME COMMISSION

Notice of Request for Public Comments Regarding Extensions to Existing OMB Clearances

AGENCY: Federal Maritime Commission.

ACTION: Notice.

SUMMARY: The FMC is preparing a submission to the Office of Management and Budget (OMB) for continued approval of the following information collection (extension with no changes) under the provisions of the Paperwork Reduction Act of 1995, as amended (44 U.S.C. Chapter 35): OMB No. 3072–0012 (Security for the Protection of the Public and Related Application Form FMC–131). Comments submitted in response to this notice will be summarized and/or included in the request for OMB approval and will become a matter of public record.

DATES: Comments must be submitted on or before April 13, 1999.

ADDRESSES: Send comments to: Edward P. Walsh, Managing Director, Federal Maritime Commission, 800 North Capitol Street, N.W., Washington, D.C. 20573, (Telephone: (202) 523–5800).

FOR FURTHER INFORMATION CONTACT: Send requests for copies of the current OMB clearances to: George D. Bowers, Director, Office of Information Resources Management, Federal Maritime Commission, 800 North Capitol Street, N.W., Washington, D.C. 20573, (Telephone: (202) 523–5834).

SUPPLEMENTARY INFORMATION:

Security for the Protection of the Public and Application Form FMC–131—OMB Approval Number 3072–0012 Expires July 31, 1999

Abstract: Sections 2 and 3 of Public Law 89–777 (46 U.S.C. app. 817(d) and (e)) require owners or charterers of vessels with 50 or more passenger berths or stateroom accommodations and embarking passengers at United States ports and territories to establish their financial responsibility to meet liability incurred for death or injury and to indemnify passengers in the event of nonperformance of transportation. 46 CFR Part 540 implements Public Law 89–777 and specifies the amount of financial responsibility coverages required of such owners or charterers.

Needs and Uses: The information will be used by the Commission's staff to ensure that passenger vessel owners and charterers have evidenced financial responsibility to indemnify passengers and others in the event of nonperformance or casualty.

Frequency: Financial information is furnished quarterly, semi-annually or annually. Other information is submitted as circumstances may warrant.

Type of Respondents: The types of respondents are owners, charterers and operators of passenger vessels with 50 or more passenger berths that embark passengers from U.S. ports or territories.

Number of annual respondents: The Commission estimates an annual respondent universe of 60.

Estimated time per response: The time per response ranges from .5 to 6 hours for complying with the regulations and 8 hours for completing Application Form FMC–131. The total average time for both requirements for each respondent is 34.66 manhours.

Total Annual Burden: The Commission estimates the total manhour burden at 2,080 manhours.

Bryant L. VanBrakle,

Secretary.

[FR Doc. 99–3436 Filed 2–11–99; 8:45 am]

BILLING CODE 6730–01–M

DEPARTMENT OF HEALTH AND HUMAN SERVICES

Administration for Children and Families

Submission for OMB Review; Comment Request

Title: Interim Tribal TANF Data Report.

OMB No.: 0970–0176.

Description: This information is being collected to meet the statutory requirements of section 411 of the Social Security Act and section 116 of the Personal Responsibility and Work Opportunity Reconciliation Act of 1996. It consists of disaggregated demographic and program information that will be used to determine participation rates and other statutorily required indicators for the Tribal Temporary Assistance for Needy Families (Tribal TANF) program.

Respondents: State, Local or Tribal Government.

ANNUAL BURDEN ESTIMATES

Instrument	Number of respondents	Number of responses per respondent	Average burden hours per response	Total burden hours
Interim Tribal TANF Data Report	18	4	451	32,472

Estimated Total Annual Burden Hours: 32,472.

Additional Information:

Copies of the proposed collection may be obtained by writing to the Administration for Children and Families, Office of Information Services, 370 L'Enfant Promenade, S.W., Washington, D.C. 20447, Attn: ACF Reports Clearance Officer.

OMB Comment:

OMB is required to make a decision concerning the collection of information between 30 to 60 days after publication of this document in the **Federal Register.** Therefore, a comment is best assured of having its full effect if OMB receives it within 30 days of publication. Written comments and recommendations for the proposed information collection should be sent directly to the following: Office of Management and Budget, Paperwork Reduction Project, 725 17th Street, N.W., Washington, D.C. 20503, Attn: Lori Schack.

Dated: February 8, 1999.

Bob Sargis,

Acting Reports Clearance Officer.

[FR Doc. 99–3475 Filed 2–11–99; 8:45 am]

BILLING CODE 4184–01–M

DEPARTMENT OF HEALTH AND HUMAN SERVICES

Food and Drug Administration

[Docket No. 98N–0364]

Agency Information Collection Activities; Announcement of OMB Approval; Recordkeeping for Electronic Products, Specific Product Requirements; Correction

AGENCY: Food and Drug Administration, HHS.

ACTION: Notice; correction.

SUMMARY: The Food and Drug Administration (FDA) is correcting a notice that appeared in the **Federal Register** of January 5, 1999 (64 FR 516). The document announced that a collection of information entitled ''Reporting and Recordkeeping for Electronic Products: Specific Product Requirements'' has been approved by the Office of Management and Budget under the Paperwork Reduction Act of 1995. The document was inadvertently published with an incorrect docket number. This document corrects that error.

FOR FURTHER INFORMATION CONTACT: Silvia R. Fasce, Office of Policy (HF–27), Food and Drug Administration, 5600 Fishers Lane, Rockville, MD 20857, 301–443–2994.

In FR Doc. 99–71, appearing on page 516 in the **Federal Register** of Tuesday, January 5, 1999, the following correction is made:

On page 516, in the first column, ''[Docket No. 98N–0213]'' is corrected to read ''[Docket No. 98N–0364]''.

Dated: February 5, 1999.

William K. Hubbard,

Associate Commissioner for Policy Coordination.

[FR Doc. 99–3438 Filed 2–11–99; 8:45 am]

BILLING CODE 4160–01–F

DEPARTMENT OF HEALTH AND HUMAN SERVICES

Food and Drug Administration

[Docket No. 98D–0132]

FDA Modernization Act of 1997: Guidance on Medical Device Tracking; Availability

AGENCY: Food and Drug Administration, HHS.

ACTION: Notice.

SUMMARY: The Food and Drug Administration (FDA) is announcing the availability of the revised final guidance entitled ''Guidance on Medical Device Tracking.'' It replaces the previous final guidance issued on March 4, 1998. This revised final guidance provides guidelines to manufacturers and distributors concerning their responsibilities for medical device tracking under the Federal Food, Drug, and Cosmetic Act (the act) as amended by the Food and Drug Administration Modernization Act of 1997 (FDAMA).

DATES: Written comments may be submitted at any time.

ADDRESSES: Submit written requests for single copies on a 3.5″ diskette of the revised final guidance entitled ''Guidance on Medical Device Tracking'' to the Division of Small Manufacturers Assistance (HFZ–220), Center for Devices and Radiological Health, Food and Drug Administration, 1350 Piccard Dr., Rockville, MD 20850. Send two self-addressed adhesive labels to assist that office in processing your request, or fax your request to 301–443–8818. Submit written comments on ''Guidance on Medical Device Tracking'' to the contact person (address below). See the **SUPPLEMENTARY INFORMATION** section for information on electronic access to the guidance.

FOR FURTHER INFORMATION CONTACT: Chester T. Reynolds, Center for Devices and Radiological Health (HFZ–300), Food and Drug Administration, 2094 Gaither Rd., Rockville, MD 20850, 301–594–4618.

SUPPLEMENTARY INFORMATION:

I. Background

Section 211 of FDAMA (Pub. L. 105–115) amended the tracking provisions of section 519(e) of the act (21 U.S.C. 360i(e)) to authorize FDA, at its discretion, to issue orders that require a manufacturer to track a class II or class III device if the failure of the device would be reasonably likely to have serious adverse health consequences, or the device is intended to be implanted in the human body for more than 1 year, or is life sustaining or life supporting and used outside a device user facility. The FDAMA tracking provisions became effective on February 19, 1998.

On January 15, 1998, FDA conducted a public meeting to discuss FDAMA changes in section 519(e) of the act. Comments were received concerning factors FDA should consider in determining what devices are subject to FDAMA tracking requirements. On February 11, 1998, FDA issued tracking orders, under the revised FDAMA tracking provisions which became effective on February 19, 1998, to manufacturers of devices that were subject to tracking previously under the Safe Medical Devices Act of 1990 (the SMDA) provisions (21 CFR 821.20(b)(1), (b)(2), and (c)). Additionally, tracking orders were issued to manufacturers of intraocular lenses and arterial stents that had not been subject to tracking under the SMDA provisions (63 FR