comments and suggestions submitted within 60 days of this publication.

Dated: October 28, 1996.

Bob Sargis,

Acting Reports Clearance Officer.

[FR Doc. 96–28139 Filed 11–1–96; 8:45 am]

BILLING CODE 4184–01–M

Submission for OMB Review; Comment Request

Title: Federal Parent Locator Service.
OMB No.: 0970–0142.

Description: The Office of Child support Enforcement (OCSE) operates the Federal Parent Locator Services (FPLS), a computerized national location network which provides address and social security number information to State and local child support enforcement agencies upon request to locate parents in order to establish or enforce a child support order and to assist authorized persons in resolving parental kidnapping and child custody cases.

State and local agency requests to the FPLS can be made by tape, cartridge, electronic file transfer or by dialing-up using a personal computer. The FPLS serves as a conduit between child support enforcement offices and Federal and State agencies by conducting weekly, biweekly, or monthly matches of the collected information with various agencies and distributing the information back to the requesting State or local child support office.

Respondents: State, Local, Tribal or Federal Govt. Governments.

ANNUAL BURDEN ESTIMATE

Instrument	Number of respondents	Number of responses per respondent	Average burden hours per response	Total burden hours
Standard Forms	200	24	1	4,800
Estimated Total Annual Burden Hours:				4,800

Explanation

*The specific number of annual burden hours per respondent will vary depending on individual circumstance including a States' frequency in submitting requests and their mode of submission.

*Burden hour for initial collection of information included in the submission are not considered as part of their day-to-day operation of the child support enforcement program.

Additional Information: Copies of the proposed collection may be obtained 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, D.C. 20447, Attn: ACF Reports Clearance Officer.

OMB Comment: OMB is required to make a decision concerning the collection of information between 30 and 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: Ms. Wendy Taylor.

Dated: October 28, 1996.

Douglas J. Godesky,

Reports Clearance Officer.

[FR Doc. 96–28140 Filed 11–1–96; 8:45 am]

BILLING CODE 4184–01–M

Food and Drug Administration

[Docket No. 96N–0298]

Agency Information Collection Activities: Proposed Collection; Comment Request; Extension

AGENCY: Food and Drug Administration, HHS.

ACTION: Notice.

SUMMARY: The Food and Drug Administration (FDA) is announcing an opportunity for public comment on the proposed collection of certain information by the agency. Under the Paperwork Reduction Act of 1995, Federal agencies are required to publish notice in the Federal Register concerning each proposed collection of information, including each proposed extension of an existing collection of information, and to allow 60 days for public comment in response to the notice. This notice solicits comments on the voluntary collection of information for the Medical Devices Standards Activities Report, a comprehensive listing of current national and international standards for medical devices.

DATES: Submit written comments on the collection of information by January 3, 1997.

ADDRESSES: Submit written comments on the collection of information to the Dockets Management Branch (HFA–305), Food and Drug Administration, 12420 Parklawn Dr., rm. 1–23, Rockville, MD 20857. All comments should be identified with the docket number found in brackets in the heading of this document.

FOR FURTHER INFORMATION CONTACT: Charity B. Smith, Office of Information Resources Management (HFA–250), Food and Drug Administration, 5600 Fishers Lane, rm. 16B–19, Rockville, MD 20857, 301–827–1686.

SUPPLEMENTARY INFORMATION: Section 3506(c)(2)(A) of the Paperwork Reduction Act of 1995 (44 U.S.C. 3506(c)(2)(A)) requires Federal agencies to provide a 60-day notice in the Federal Register concerning each proposed collection of information, including each proposed extension of an existing collection of information. ''Collection of information'' is defined in 44 U.S.C. 3502(3) and 5 CFR 1320.3(c). To comply with this requirement, FDA is publishing notice of the proposed collection of information listed below.

With respect to the following collection of information, FDA invites comments on: (1) Whether the proposed collection of information is necessary for the proper performance of FDA's functions, including whether the information will have practical utility; (2) the accuracy of FDA's estimate of the burden of the proposed collection of information, including the validity of the methodology and assumptions used; (3) ways to enhance the quality, utility, and clarity of the information to be collected; and (4) ways to minimize the burden of the collection of information on respondents, including through the use of automated collection techniques, when appropriate, and other forms of information technology.