The proposed consent order would terminate 20 years after the date it is issued.

Donald S. Clark,

Secretary.

Concurring Statement of Commissioner Mary L. Azcuenaga in Montana Associated Physicians, Inc.

[File No. 911–0008]

I concur in the decision to issue the complaint and accept the order for public comment and write separately to emphasize two points. First, the complaint and order do not directly challenge the organization and conduct of the Billings Physician Hospital Alliance, Inc., as a physician hospital organization (PHO), and in my view, this order should cast no shadow on the activities of PHO's. Second, although I concur in the unusual and complicated fencing-in relief in the particular circumstances of this case, in my view, this negotiated order is not, and should not be viewed as, a guide for what a PHO can and cannot do.

[FR Doc. 96–28277 Filed 11–1–96; 8:45 am]

BILLING CODE 6750–01–M

GENERAL SERVICES ADMINISTRATION

Federal Acquisition Policy Division, FAR Secretariat Stocking Change of a Standard Form

AGENCY: General Services Administration.

ACTION: Notice.

SUMMARY: The General Services Administration/FAR Secretariat is changing the stocking of the following Standard form because of low user demand: SF 25B, Continuation Sheet for SF 24, 25, and 25A.

Since this form is now authorized for local reproduction, you can obtain the updated camera copy in two ways:

On the internet. Address: http://www.gsa.gov/forms, or;

From CARM, Attn.: Barbara Williams, (202) 501–0581.

FOR FURTHER INFORMATION CONTACT: FAR Secretariat, (202) 501–4755.

DATES: EFFECTIVE NOVEMBER 4, 1996.

Dated October 8, 1996.

Theodore D. Freed,

Standard and Optional Forms Management Officer.

[FR Doc. 96–28188 Filed 11–1–96; 8:45 am]

BILLING CODE 6820–34–M

Revision and Stocking Changes of Standard Forms

AGENCY: Public Building Service, General Services Administration.

ACTION: Notice.

SUMMARY: The General Services Administration is changing the stocking requirement of SF 118, Report of Excess Real Property, SF 118A, Buildings, Structures, Utilities, and Miscellaneous Facilities (Schedule A—Supplement to Report of Excess), SF 118B, Land (Schedule B—Supplement to Report of Excess Real Property) and SF 118C, Related Personal Property (Schedule C—Supplement to Report of Excess Real Property). These forms are revised to include metric measurements and authorized for local reproduction. Since these forms are authorized for local reproduction, you can obtain the updated camera copy in two ways.

On the Internet. Address: http://www.gsa.gov/forms, or;

From CARM, Attn.: Barbara Williams, (202) 501–0581.

FOR FURTHER INFORMATION CONTACT: Ronald Rice, (202) 501–0074. This contact is for information on completing the form only.

DATES: Effective November 4, 1996.

Dated: October 24, 1996.

Barbara M. Williams,

Deputy Standard and Optional Forms Management Officer.

[FR Doc. 96–28166 Filed 11–1–96; 8:45 am]

BILLING CODE 6820–34–M

DEPARTMENT OF HEALTH AND HUMAN SERVICES

Administration for Children and Families

Proposed Information Collection Activity; Comment Request

Proposed Projects

Title: Detailed Case Data Component (DCDC) of the National Child Abuse and Neglect Data System.

OMB No.: 0980–0256.

Description: The Detailed Case Data Component of the National Child Abuse and Neglect Data System compiles automated case-level data on child maltreatment investigated by State child protective services agencies. Data are collected on reports of abuse and neglect, characteristics of victims, risk factors associated with victims and their families, and the development of polices and programs relating the child abuse and neglect at the National, State and local levels.

Respondents: State, Local or Tribal Govt.

ANNUAL BURDEN ESTIMATES

Instrument	Number of respondents	Number of responses per respondent	Average burden hours per response	Total burden hours
DCDC	56	1	110	6,160
Estimated Total Annual Burden Hours:				6,160

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, D.C. 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 proposed collection of information; (c) 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: 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.