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 Administration, Office of Information Services, 370 L'Enfant Promenade, SW., Washington, DC 20447, Attn: ACF Reports Clearance Officer. E-mail address: *infocollection@acf.hhs.gov*. 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: July 9, 2008.

Janean Chambers,

Reports Clearance Officer.

[FR Doc. E8–15897 Filed 7–17–08; 8:45 am]

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DEPARTMENT OF HEALTH AND HUMAN SERVICES

Administration for Children and Families

Submission for OMB Review; Comment Request

Title: Mentoring Children of Prisoners Service Delivery Demonstration Project Data Collection (MCP) Program.

OMB No.: New Collection.

Description: The Promoting Safe and Stable Families Amendments, as reauthorized (2006), amended Title IV–B of the Social Security Act (42 U.S.C. 629–629e) providing funding for a service delivery demonstration project for the Mentoring Children of Prisoners (MCP) program. The grantee shall identify children of prisoners not being served by the grant program, provide families of identified children with a voucher for mentoring services and a list of quality mentoring programs, and monitor the delivery of mentoring services provided. The Family and Youth Services Bureau (FYSB) of the Administration for Children and Families, United States Department of Health and Human Services, administers the Mentoring Children of Prisoners (MCP) program. The MCP program provides children of prisoners with caring adult mentors, supporting one-to-one mentoring relationships. Research in other populations has shown that such relationships can lead to reductions in risk behaviors and improvements in academic, behavioral and psychological outcomes in children and youth. Although the MCP program was developed based on research documenting the efficacy of mentoring as a general intervention strategy, it is not yet known whether or not this particular intervention yields positive outcomes for the children of prisoners population. Little is known about how mentoring relationships work for these youth, and how effective mentoring relationships for children of prisoners differ from effective mentoring relationships for other youth. In addition, little is known about children of prisoners in general and thus a survey of MCP program youth has the potential to provide important data about this relatively unstudied population.

The evaluation and data collection proposed in this notice are to fulfill the statutory requirement under Section 8, subsection h(1) of the Child and Family Services Improvement Act of 2006, as amended, that the Secretary of the Department of Health and Human Services evaluate outcomes of the MCP service delivery demonstration project and report to Congress on the findings. The information collected will also be used for accountability monitoring, management improvement, and research. Data collection will ensure that the grantee knows that mentoring relationships are meeting the established milestones and that mentoring activities are faithful to characteristics established by research as essential to success. Data collected will allow the Administration for Children and Families to compare the MCP service delivery demonstration project with the MCP grant program. Data collected will also support the grantee as it carries out ongoing responsibilities and manages information for internal uses.

Respondents: Public, faith-based and community organizations applying to and implementing the MCP service delivery demonstration project.

ANNUAL BURDEN ESTIMATES

Instrument	Number of respondents	Number of responses per respondent	Average burden hours per response	Total burden hours
Program Application	325	1	2	650
MentorPRO Basic Mentoring Practices and Relationship Data	250	120	0.50	15,000
Child Application	4,200	1	0.50	2,100
Baseline Youth Survey	3,000	1	0.50	1,500
Follow-Up Youth Survey	2,000	1	0.50	1,000
Relationship Quality Survey	2,250	1	0.50	1,125
Program Survey	250	1	0.50	125
Mentor Survey	2,000	1	0.50	1,000
Payment Information	1	52	2	104

Estimated Total Annual Burden Hours: 22,604.

Additional Information: Copies of the proposed collection may be obtained by writing to the Administration for Children and Families, Office of Administration, Office of Information Services, 370 L'Enfant Promenade, SW., Washington, DC 20447, Attn: ACF Reports Clearance Officer. All requests should be identified by the title of the information collection. E-mail address: *infocollection@acf.hhs.gov*.

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, Fax: 202–395–6974, Attn: Desk Officer for the Administration for Children and Families.

Dated: July 9, 2008.

Janean Chambers,

Reports Clearance Officer.

[FR Doc. E8–15898 Filed 7–17–08; 8:45 am]

BILLING CODE 4184–01–P

DEPARTMENT OF HEALTH AND HUMAN SERVICES

Food and Drug Administration

[Docket No. FDA–2008–N–0397]

Agency Information Collection Activities; Proposed Collection; Comment Request; State Enforcement Notifications

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 (the PRA), 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 reporting requirements contained in existing FDA regulations governing State enforcement notifications.

DATES: Submit written or electronic comments on the collection of information by September 16, 2008.

ADDRESSES: Submit electronic comments on the collection of information to *http://www.regulations.gov*. Submit written comments on the collection of information to the Division of Dockets Management (HFA–305), Food and Drug Administration, 5630 Fishers Lane, rm. 1061, Rockville, MD 20852. All comments should be identified with the docket number found in brackets in the heading of this document.

FOR FURTHER INFORMATION CONTACT: Jonna Capezzuto, Office of the Chief Information Officer (HFA–250), Food and Drug Administration, 5600 Fishers Lane, Rockville, MD 20857, 301–796–3794.

SUPPLEMENTARY INFORMATION: Under the PRA (44 U.S.C. 3501–3520), Federal agencies must obtain approval from the Office of Management and Budget (OMB) for each collection of information they conduct or sponsor. ''Collection of information'' is defined in 44 U.S.C. 3502(3) and 5 CFR 1320.3(c) and includes agency requests or requirements that members of the public submit reports, keep records, or provide information to a third party. Section 3506(c)(2)(A) of the PRA (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, before submitting the collection to OMB for approval. To comply with this requirement, FDA is publishing notice of the proposed collection of information set forth in this document.

With respect to the following collection of information, FDA invites comments on these topics: (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.

State Enforcement Notifications—21 CFR 100.2(d) (OMB Control Number 0910–0275)—Extension

Section 310(b) of the Federal Food, Drug, and Cosmetic Act (the act) (21 U.S.C. 337(b)) authorizes States to enforce certain sections of the act in their own names, but provides that States must notify FDA before doing so. Section 100.2(d) (21 CFR 100.2 (d)) sets forth the information that a State must provide to FDA in a letter of notification when it intends to take enforcement action under the act against a particular food located in the State. The information required under § 100.2(d) will enable FDA to identify the food against which the State intends to take action and advise the State whether Federal action has been taken against it. With certain narrow exceptions, Federal enforcement action precludes State action under the act.

FDA estimates the burden of this collection of information as follows:

TABLE 1.—ESTIMATED ANNUAL REPORTING BURDEN[1]

21 CFR Section	No. of Respondents	Annual Frequency per Response	Total Annual Responses	Hours per Response	Total Hours
100.2(d)	1	1	1	10	10

[1]There are no capital costs or operating and maintenance costs associated with this collection of information.

The estimated reporting burden for § 100.2(d) is minimal because enforcement notifications are seldom used by States. During the last 3 years, FDA has not received any new enforcement notifications; therefore, the agency estimates that one or fewer notifications will be submitted annually. Although FDA has not received any new enforcement notifications in the last 3 years, it believes these information collection provisions should be extended to provide for the potential future need of a State government to submit enforcement notifications informing FDA when it intends to take enforcement action under the act against a particular food located in the State.

Please note that on January 15, 2008, the FDA Division of Dockets Management Web site transitioned to the Federal Dockets Management System (FDMS). FDMS is a Government-wide, electronic docket management system. Electronic comments or submissions will be accepted by FDA only through FDMS at *http://www.regulations.gov*.