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 Control and Prevention.*
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**DEPARTMENT OF HEALTH AND
 HUMAN SERVICES**

**Administration for Children and
 Families**

**Submission for OMB Review;
 Comment Request**

Title: Tribal TANF Data Report, TANF
 Annual Report, and Reasonable Cause/

Corrective Action Documentation
 Process—Final.
OMB No.: 0970-0215.
Description: 42 U.S.C. 612 (Section
 412 of the Social Security Act as
 amended by Pub. L. 104-193, the
 Personal Responsibility and Work
 Opportunity Reconciliation Act of 1996
 (PRWORA)), mandates that federally
 recognized Indian Tribes with an
 approved Tribal TANF program collect
 and submit to the Secretary of the
 Department of Health and Human
 Services data on the recipients served
 by the Tribes' programs. This
 information includes both aggregated
 and disaggregated data on case
 characteristics and individual
 characteristics. In addition, Tribes that

are subject to a penalty are allowed to
 provide reasonable cause justifications
 as to why a penalty should not be
 imposed or may develop and implement
 corrective compliance procedures to
 eliminate the source of the penalty.
 Finally, there is an annual report, which
 requires the Tribes to describe program
 characteristics. All of the above
 requirements are currently approved by
 OMB and the Administration for
 Children and Families is simply
 proposing to extend them without any
 changes.

Respondents: Indian Tribes.

ANNUAL BURDEN ESTIMATES

Instrument	Number of respondents	Number of responses per respondent	Average burden hours per response	Total burden hours
Final Tribal TANF Data Report	66	4	451	119,064
Tribal TANF Annual Report	66	1	40	2,640
Tribal TANF Reasonable Cause/Corrective	66	1	60	3,960

*Estimated Total Annual Burden
 Hours:* 125,664.

Additional Information: Copies of the
 proposed collection may be obtained by
 writing to the Administration for
 Children and Families, Office of
 Planning, Research and Evaluation, 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. Email 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: 202-395-7285,
 Email:
OIRA_SUBMISSION@OMB.EOP.GOV.
 Attn: Desk Officer for the

Administration for Children and
 Families.

Robert Sargis,
Reports Clearance Officer.
 [FR Doc. 2012-13630 Filed 6-5-12; 8:45 am]
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**DEPARTMENT OF HEALTH AND
 HUMAN SERVICES**

**Administration for Children and
 Families**

**Submission for OMB Review;
 Comment Request**

Proposed Projects:
Title: Performance Measures for
 Community-Centered Healthy Marriage,
 Pathways to Responsible Fatherhood
 and Community-Centered Responsible
 Fatherhood Ex-Prisoner Reentry Grant
 Programs.
OMB No.: 0970-0365.
Description: The Office of Family
 Assistance (OFA), Administration for
 Children and Families (ACF), U.S.
 Department of Health and Human
 Services (HHS), intends to request
 approval from the Office of Management
 and Budget (OMB) to renew OMB Form
 0970-0365 for the collection of
 performance measures from grantees for
 the Community-Centered Healthy

Marriage, Pathways to Responsible
 Fatherhood and Community-Centered
 Responsible Fatherhood Ex-Prisoner
 Reentry discretionary grant programs.
 The performance measure data obtained
 from the grantees will be used by OFA
 to report on the overall performance of
 these grant programs. Data will be
 collected from all 61 Community-
 Centered Healthy Marriage, 53 Pathways
 to Responsible Fatherhood and 4
 Community-Centered Responsible
 Fatherhood Ex-Prisoner Reentry
 grantees in the OFA programs. Grantees
 will report on program and participant
 outcomes in such areas as participants'
 improvement in knowledge skills,
 attitudes, and behaviors related to
 healthy marriage and responsible
 fatherhood. Grantees will be asked to
 input data for selected outcomes for
 activities funded under the grants.
 Grantees will extract data from program
 records and will report the data twice
 yearly through an on-line data
 collection tool. Training and assistance
 will be provided to grantees to support
 this data collection process.

Respondents: Office of Family
 Assistance Funded Community-
 Centered Healthy Marriage, Pathways to
 Responsible Fatherhood and
 Community-Centered Responsible
 Fatherhood Ex-Prisoner Reentry
 Grantees.

ANNUAL BURDEN ESTIMATES

Instrument	Number of respondents	Number of responses per respondent	Average burden hours per response	Total annual burden hours
Performance measure reporting form (for private sector affected public)	103	2	0.8	165
Performance measure reporting form (for State, local, and tribal government affected public)	15	2	0.8	24
Estimated Total Annual Burden Hours				189

Additional Information: Copies of the proposed collection may be obtained by writing to the Administration for Children and Families, Office of Planning, Research and Evaluation, 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. Email 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: 202-395-7285, Email: OIRA_SUBMISSION@OMB.EOP.GOV, Attn: Desk Officer for the Administration for Children and Families.

Robert Sargis,

Reports Clearance Officer.

[FR Doc. 2012-13602 Filed 6-5-12; 8:45 am]

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DEPARTMENT OF HEALTH AND HUMAN SERVICES

Food and Drug Administration

[Docket No. FDA-2012-N-0536]

Agency Information Collection Activities; Proposed Collection; Comment Request; Medical Device User Fee Cover Sheet, Form FDA 3601

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 Form FDA 3601, entitled "Medical Device User Fee Cover Sheet," which must be submitted along with certain medical device product applications, supplements, and fee payment of those applications.

DATES: Submit either electronic or written comments on the collection of information by August 6, 2012.

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:

Daniel Gittleston, Office of Information Management, Food and Drug Administration, 1350 Piccard Dr., PI50-400B, Rockville, MD 20850, 301-796-5156, Daniel.Gittleston@fda.hhs.gov.

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.

Medical Device User Fee Cover Sheet—Form FDA 3601 (OMB Control Number 0910-0511)—Extension

The Federal Food, Drug, and Cosmetic Act, as amended by the Medical Device User Fee and Modernization Act of 2002 (Pub. L. 107-250), and the Medical Device User Fee Amendments of 2007 (Title II of the Food and Drug Administration Amendments Act of 2007), authorizes FDA to collect user fees for certain medical device applications. Under this authority, companies pay a fee for certain new medical device applications or supplements submitted to the Agency for review. Because the submission of user fees concurrently with applications and supplements is required, the review of an application cannot begin until the fee is submitted. Form FDA 3601, the "Medical Device User Fee Cover Sheet," is designed to provide the minimum necessary information to determine whether a fee is required for review of an application, to determine the amount of the fee required, and to account for and track user fees.

The form provides a cross-reference between the fees submitted for an application with the actual submitted application by using a unique number