

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) ways to enhance 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. Written comments should be received within 60 days of this notice.

Proposed Project

Understanding the Community Context of the Diabetes Education in Tribal Schools Project—New—National Center for Chronic Disease Prevention and Health Promotion/Division of Diabetes Translation (NCCDPHP/DDT),

Centers for Disease Control and Prevention (CDC).

Background and Brief Description: This study is part of a larger evaluation of the multi-year Diabetes Education in Tribal Schools (DETS) project to develop and pilot test a science based diabetes prevention curriculum for Native American school children. As part of the overall evaluation (before the curriculum is pilot tested), it will be important to understand the community context and identify implementation issues. Through a series of qualitative interviews with key informants, the study will obtain information about: (1) The community's experience with diabetes; (2) community readiness to adopt the DETS curriculum; (3) the connection between the DETS project and the community; and (4) the best fit between the curriculum and community schools.

The participants for this study will include key informants in five categories: Community leaders, DETS Advisory Board members, DETS Curriculum Subcommittee members, community teachers, and community parents. Potential participants will be identified by DETS Subcommittee members and invited to participate in this research activity. These individuals will be invited to participate because they are already involved in the project and are familiar with the curriculum.

A maximum of 18 individuals from each category will be interviewed for a total of 90 participants. All participants will be adults, both male and female, over the age of 18. It is expected that approximately 75% of participants will be Native American and 25% will be non-Native American. There is no cost to respondents other than their time.

Estimate of Annualized Burden Table:

Respondent	Number of respondents	Number of responses per respondent	Avg. burden per response (in hours)	Total burden hours
Community Leaders/Elders	18	1	45/60	13.5
Parents	18	1	45/60	13.5
Teachers	18	1	45/60	13.5
DETS Curriculum Subcommittee Members	18	1	45/60	13.5
DETS Advisory Board Members	18	1	45/60	13.5
Totals:	90			67.5

Dated: March 21, 2005.

Betsey Dunaway,

Acting Reports Clearance Officer, Centers for Disease Control and Prevention.

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DEPARTMENT OF HEALTH AND HUMAN SERVICES

Administration for Children and Families

Privacy Act of 1974, as Amended; Computer Matching Program

AGENCY: Office of Child Support Enforcement (OCSE), ACF, DHHS.

ACTION: Notice of a computer matching program.

SUMMARY: In compliance with the Privacy Act of 1974, as amended by Pub. L. 100-503, the Computer Matching and Privacy Protection Act of 1988, we are publishing a notice of a computer matching program that OCSE will conduct on behalf of itself and the District of Columbia Department of Human Services, Income Maintenance Administration (IMA) for verification of

continued eligibility for Public Assistance. The match will utilize National Directory of New Hire (NDNH) records and IMA records. The purpose of the computer matching program is to exchange personal data for purposes of identifying individuals who are employed and also are receiving payments pursuant to the Temporary Assistance for Needy Families (TANF) benefit program administered by IMA.

DATES: OCSE will file a report of the subject OCSE matching program with the Committee on Homeland Security and Governmental Affairs of the Senate and the Committee on Government Reform of the House of Representatives, and the Office of Information and Regulatory Affairs, Office of Management and Budget (OMB). The matching program will be effective as indicated below.

ADDRESSES: Interested parties may comment on this notice by writing to the Director, Office of Federal Systems, Office of Child Support Enforcement, Aerospace Building, 370 L'Enfant Promenade, SW., Washington, DC 20047. All comments received will be

available for public inspection at this address.

FOR FURTHER INFORMATION CONTACT:

Director, Office of Federal Systems, Office of Child Support Enforcement, Aerospace Building, 370 L'Enfant Promenade, SW., Washington, DC 20047. Telephone Number (202) 401-9271.

SUPPLEMENTARY INFORMATION: Pub. L. 100-503, the Computer Matching and Privacy Protection Act of 1988, amended the Privacy Act (5 U.S.C. 552a) by adding certain protections for individuals applying for and receiving Federal benefits. The law regulates the use of computer matching by Federal agencies when records in a system of records are matched with other Federal, state and local government records.

The amendments require Federal agencies involved in computer matching programs to:

1. Negotiate written agreements with source agencies;
2. Provide notification to applicants and beneficiaries that their records are subject to matching;

3. Verify match findings before reducing, suspending, or terminating an individual's benefits or payments;

4. Furnish detailed reports to Congress and OMB; and

5. Establish a Data Integrity Board that must approve matching agreements.

This Computer Match meets the requirements of Pub. L. 100-503.

Dated: March 22, 2005.

David H. Siegel,

Acting Commissioner, Office of Child Support Enforcement.

Notice of Computer Matching Program

A. Participating Agencies

OCSE and IMA.

B. Purpose of the Match

To exchange personal data for purposes of identifying individuals who are employed and also are receiving payments pursuant to TANF benefit programs being administered by the IMA and to verify continuing eligibility for TANF benefits.

OCSE will match public assistance records, obtained from IMA, to the NDNH. After matching has been conducted, OCSE will provide matched data to IMA which will use this information to verify the continued eligibility of individuals to receive public assistance benefits and, if ineligible, to take such action, as may be authorized by law and regulation. Under the matching program, IMA will obtain data provided by OCSE.

C. Authority for Conducting the Match

The authority for conducting the matching program is contained in section 453(j)(3) of the Social Security Act (42 U.S.C. 653(j)(3)).

D. Records To Be Matched

The system of records maintained by the ACF under the Privacy Act of 1974, as amended, 5 U.S.C. 552a, from which records will be disclosed for the purpose of this computer match, is the Location and Collection System of Records, DHHS/OCSE No. 09-90-0074, last published in the **Federal Register** at 69 FR 31392 on June 3, 2004. The match is a routine use under this system of records.

OCSE, as the source agency, will collect from IMA electronic files containing the names and other personal identifying data of eligible public assistance beneficiaries. Upon receipt of the electronic files of IMA beneficiaries, OCSE will perform a computer match against the NDNH. The NDNH database consists of Quarterly Wage, New Hire, and Unemployment

Insurance information. The matches will be furnished by OCSE to IMA.

1. The electronic files provided by IMA will contain data elements of the client's name and SSN.

2. OCSE will match the SSN on the IMA file by computer against the NDNH database. Matching records, based on SSNs, will produce data elements of the individual's name; SSN; employer, and current work or home address, etc.

E. Inclusive Dates of the Matching Program

The effective date of the matching agreement and date when matching may actually begin shall be at the expiration of the 40-day review period for OMB and Congress, or 30 days after publication of the matching notice in the **Federal Register**, whichever date is later. By agreement between DHHS and IMA, the matching program will be in effect for 18 months from the effective date, with an option to renew for 12 additional months, unless one of the parties to the agreement advises the other by written request to terminate or modify the agreement.

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DEPARTMENT OF HEALTH AND HUMAN SERVICES

Food and Drug Administration

[Docket No. 2005N-0100]

Agency Information Collection Activities; Proposed Collection; Comment Request; Current Good Manufacturing Practice Regulations for Finished Pharmaceuticals

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 the information collection provisions of FDA's current good manufacturing practice (CGMP) regulations for finished pharmaceuticals.

DATES: Submit written or electronic comments on the collection of information by May 27, 2005.

ADDRESSES: Submit electronic comments on the collection of information to: <http://www.fda.gov/dockets/ecomments>. 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: Karen Nelson, Office of Management Programs (HFA-250), Food and Drug Administration, 5600 Fishers Lane, Rockville, MD 20857, 301-827-1482.

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.

CGMP Regulations for Finished Pharmaceuticals—21 CFR Parts 210 and 211 (OMB Control Number 0910-0139)—Extension