Parents are also asked to collect cheek cells from themselves and their infant for DNA. The collection of cheek cells by the mother, father, and infant takes about 10 minutes per person. Each person rubs 1 brush inside the left cheek and 1 brush inside the right cheek for a total of 2 brushes per person. Collection of the cheek cells takes approximately 1–2 minutes, but the estimate of burden is 10 minutes to account for reading and understanding the consent form and specimen collection instructions and mailing back the completed kits. The anticipated maximum burden for collection of the cheek cells is 200 hours per center per year.

Information gathered from both the interviews and the DNA specimens have been and will continue to be used to study independent genetic and environmental factors as well as gene-environment interactions for a broad range of carefully classified birth defects.

This request is submitted to obtain OMB clearance for three additional years.

There are no costs to the respondents other than their time.

ESTIMATED ANNUALIZED BURDEN HOURS PER CENTER

Respondents	Number of respondents	Number of responses per respondent	Average burden per response	Total burden hours
NBDPS case/control interview	400	1	1	400
Biologic Specimen Collection	1,200	1	10/60	200
Total				600

Kimberly S. Lane,

Acting Reports Clearance Officer.

[FR Doc. 2012–4170 Filed 2–22–12; 8:45 am]

BILLING CODE 4163–18–P

DEPARTMENT OF HEALTH AND HUMAN SERVICES

Administration for Children and Families

Proposed Information Collection Activity; Comment Request

Proposed Projects: Data Collection for some of the Children's Bureau Funded Discretionary Programs.

Title: Performance Measurement On-Line Tool (PMOTOOL).

OMB No.: New Collection.

Description: The Performance Measurement On-Line Tool (PMOTOOL) was designed by the Children's Bureau to collect data, in an automated format, from specified discretionary grants funded by the Children's Bureau. The data collected by this instrument will be submitted by individual discretionary grantees funded under the following programs: Abandoned Infants Assistance Program, Infant Adoption Awareness Program, Adoption Opportunities Program, Child Abuse and Neglect Program and the Child Welfare Training Program. Grantees will submit this information on a semi-annual basis in conjunction with their semi-annual program progress report.

The purpose of this data collection is to assist the Children's Bureau in using the aggregated data to examine the social impact or public benefit under each funded federal program. These measurable outcomes will serve as evidence that the federally funded programs are making progress toward achieving broad, legislated program goals.

Respondents: Selected clusters of competitive grant programs funded by the Children's Bureau.

ANNUAL BURDEN ESTIMATES

Instrument	Number of respondents	Number of responses per respondent	Average burden hours per response	Total burden hours
Performance Measurement On-Line Tool.	Abandoned Infants Assistance Program Range 20–30.	2 per fiscal year	One hour per response field ..	Range 40–60
Performance Measurement On-Line Tool.	Infant Adoption Awareness Program Range 6.	2 per fiscal year	One hour per response field ..	Range 12
Performance Measurement On-Line Tool.	Adoption Opportunities Program Range 45–55.	2 per fiscal year	One hour per response field ..	Range 90–110
Performance Measurement On-Line Tool.	Child Abuse and Neglect Program Range 30–40.	2 per fiscal year	One hour per response field ..	Range 60–80
Performance Measurement On-Line Tool.	Child Welfare Training Program Range 40–50.	2 per fiscal year	One hour per response field ..	Range 80–100

Estimated Total Annual Burden Hours: 282–350.

In compliance with the requirements of Section 506(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 Planning, Research and Evaluation, 370 L'Enfant Promenade SW., Washington, DC 20447, Attn: ACF Reports Clearance Officer. Email 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.

Robert Sargis,

Reports Clearance Officer.

[FR Doc. 2012–4143 Filed 2–22–12; 8:45 am]

BILLING CODE 4184–01–P

DEPARTMENT OF HEALTH AND HUMAN SERVICES

Administration for Children and Families

Proposed Information Collection Activity; Comment Request

Title: Assets for Independence (AFI) Program Evaluation.

OMB No.: New Collection.

Description: The U.S. Department of Health and Human Services, Administration for Children and Families (ACF) is proposing a data collection activity as part of an experimental evaluation of the Assets for Independence (AFI) Program. The purpose of this study is to assess the impact of participation in AFI-funded individual development account (IDA) projects on the savings, asset purchases, and economic well-being of low-income individuals and families. The two primary research questions are:

- What is the impact of AFI project participation on short-term outcomes such as savings, asset purchases, and material hardship?
- How do specific API project design features affect short-term participant outcomes?

While some evaluations suggest that IDAs help low-income families save, rigorous experimental research is limited. Few studies have focused on API-funded IDAs, and few have tested alternative design features.

This evaluation—the first experimental evaluation of IDA projects operating under the Assets for Independence Act—will contribute importantly to understanding the effects of IDA project participation on project participants, particularly effects that occur within the first 12 months of participation, and how these short-term effects differ under alternative project designs. The evaluation will be conducted in two sites, with the random assignment of API-eligible cases to program and control groups. The evaluation consists of both an impact study and an implementation study. Data collection activities will span a three-year period.

Respondents: Respondent groups will include: (1) API-eligible participants and (2) API project administrators and staff members of the participating API grantees and their partnering organizations.

ANNUAL RESPONSE BURDEN ESTIMATES

Instrument	Number of respondents	Number of responses per respondent	Average burden hours per response	Estimated burden hours
Baseline survey: AFI-eligible participants	567	1	.50	284
Follow-up survey: AFI-eligible participants	482	1	.50	241
Implementation interview: Administrators and staff	10	1	1.00	10

Estimated Annual Response Burden Hours: 535.

In compliance with the requirement of section 3506(c)(2)(A) of the Paperwork Reduction Act of 1995, the Administration for Children and Families (ACF), Department of Health and Human Services, 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 in writing to the Administration for Children and Families, Office of Planning, Research and Evaluation, 370 L'Enfant Promenade SW., Washington, DC 20447, Attn: OPRE Reports Clearance Officer. Email address: *OPREinfocollection@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: February 14, 2012.

Steven M. Hanmer,

Office of Planning Research and Evaluation; ACF, Reports Clearance Officer.

[FR Doc. 2012–3946 Filed 2–22–12; 8:45 am]

BILLING CODE 4184–24–M

DEPARTMENT OF HEALTH AND HUMAN SERVICES

Food and Drug Administration

[Docket No. FDA–2012–N–0129]

Agency Information Collection Activities; Proposed Collection; Comment Request; General Licensing Provisions; Section 351(k) Biosimilar Applications; Correction

AGENCY: Food and Drug Administration, HHS.

ACTION: Notice; correction.

SUMMARY: The Food and Drug Administration (FDA) is correcting a notice that appeared in the **Federal Register** of February 15, 2012 (77 FR 8880). The document announced an opportunity for public comment on the proposed collection of certain information by the Agency. The document was published with an incorrect docket number. This document corrects that error.

FOR FURTHER INFORMATION CONTACT: Joyce Strong, Office of Policy, Food and Drug Administration, 10903 New Hampshire Ave., Bldg. 32, rm. 3208,