

variables contained in the Uniform Hospital Discharge Data Set (UHDDS) in addition to two data items (admission type and source) which are identical to those needed for billing of inpatient services for Medicare patients. In the 2003 NHDS, 426 hospitals participated.

Data for approximately forty-four percent of the responding hospitals (186) are abstracted from medical records. The remaining hospitals supply data through in-house tapes or printouts (80 hospitals) or are hospitals that belong to commercial abstract service

organizations or state data systems (160 hospitals) from which electronic data files are purchased. There is no actual cost to respondents since hospital staff who actively participate in the data collection effort are compensated by the government for their time.

Medical record abstracts	Number of respondents (hospitals)	Number of responses/respondent	Average burden/response (in hrs.)	Total burden hours
Primary Procedure Hospitals	62	250	5/60	1,292
Alternate Procedure Hospitals	124	250	1/60	517
In-House Tape or Printout Hospitals	80	12	12/60	192
Induction Forms	15	1	2	30
Non-response Study	50	1	2	100
Total	2,131

Dated: November 12, 2004.

B. Kathy Skipper,

Acting Director, Management Analysis and Services Office, Centers for Disease Control and Prevention.

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DEPARTMENT OF HEALTH AND HUMAN SERVICES

Centers for Disease Control and Prevention

[60Day-05AH]

Proposed Data Collections Submitted for Public Comment and Recommendations

In compliance with the requirement of Section 3506(c)(2)(A) of the Paperwork Reduction Act of 1995 for opportunity for public comment on proposed data collection projects, the Centers for Disease Control and Prevention (CDC) will publish periodic summaries of proposed projects. To request more information on the proposed projects or to obtain a copy of the data collection plans and instruments, call 404-498-1210 or send comments to Sandi Gambescia, CDC Assistant Reports Clearance Officer, 1600 Clifton Road, MS-E11, Atlanta, GA 30333 or send an e-mail to omb@cdc.gov.

Comments are invited 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) 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

A Comprehensive Evaluation of an Approach to Self-Management: "Diabetes: Living My Best Life"—New—National Center for Chronic Disease Prevention and Health Promotion (NCCDPHP), Centers for Disease Control and Prevention (CDC).

Background and Brief Description: African-American women are more than twice as likely as white women to be diagnosed with diabetes, and two and one-half times as likely to die from diabetic complications. The onset of type 2 diabetes in African-American adults is attributable not only to a genetic link, but also to an unhealthy lifestyle. The vast number of African-American women with type 2 diabetes report having a sedentary lifestyle and eating a diet high in fat. In addition to taking medications, lifestyle modifications, such as changes in diet, weight loss and participating in a low-impact exercise program, can significantly reduce the complications experienced by women with type 2 diabetes. Unfortunately, there is a scarcity of training and educational materials on type 2 diabetes targeting the African-American woman. The limited availability of targeted educational materials has undoubtedly contributed to an inability to manage and control this disease in this population and has resulted in a higher prevalence of disease-related comorbidities. There is a need for innovative interventions that can be used in a variety of settings and which

feature culturally appropriate information that will engage African-American women with type 2 diabetes in a proactive role in the treatment and management of their disease.

The proposed project is the evaluation of a CD-ROM educational program: "Diabetes: Living My Best Life." This product has been developed to teach African American women with type 2 diabetes self-management skills. Social Learning Theory (SLT) was used in the development of the product and the selection of the media elements. Selection of the information and tools was guided by input from an advisory board composed of professionals in the field and African American women with type 2 diabetes.

To evaluate this program there will be two questionnaires: a pre-test and a post-test. The two questionnaires will include questions on:

- Respondent demographic information (pre-test only).
- Respondent use of computers (pre-test only).
- Knowledge of diabetes.
- Self-efficacy in addressing diabetes self-management issues.
- Diabetes self-care activities.
- Feeling of empowerment around diabetes self-management.
- Social learning theory elements (post-test only).

Pre and post intervention data will be collected by computer. Burden estimates are based on observation of African-American women with type 2 diabetes who completed a formal pilot test of the pre and post-test forms. There are no costs to respondents except their time to participate in the survey.

ANNUALIZED BURDEN TABLE

Respondent	Number of respondents	Number of responses per respondent	Average burden per response (in hours)	Total burden (in hours)
African American women with Type 2 diabetes—Pre-test	66	1	20/60	22
African American women with Type 2 diabetes—Posttest	66	1	20/60	22
Total				44

Dated: November 18, 2004.
Alvin Hall,
 Director, Management Analysis and Services
 Office, Centers for Disease 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: Refugee Resettlement Program
 Estimates: CMA, ORR-1.
OMB No.: 0970-0030.
Description: The Office of Refugee Resettlement (ORR) reimburses, to the extent of available appropriations,

certain non-Federal costs for the provision of cash and medical assistance to refugees, along with allowable expenses in the administration of the Refugee Resettlement Program. ORR needs sound State estimates of likely expenditures for refugee cash, medical, and administrative (CMA) expenditures so that it can anticipate Federal costs in upcoming quarters. If Federal costs are anticipated to exceed budget allocations, ORR must take steps to reduce Federal expenses, such as limiting the number of months of eligibility for Refugee Cash Assistance and Refugee Medical Assistance.

To meet the need for reliable State estimates of anticipated expenses, ORR has developed a single-page form in which States estimate the average number of recipients for each category of assistance, the average unit cost over

the next 12 months, and the expense for the overall administration of the program. This form, the ORR-1, must be submitted prior to the beginning of each Federal fiscal year. Without this information, ORR would be out of compliance with the intent of its legislation and otherwise unable to estimate program costs adequately.

In addition, the ORR-1 serves as the State's application for reimbursement of its CMA expenses. Submission of this form is thus required by section 412(a)(4) of the Immigration and Nationality Act, which provides that "no grant or contract may be awarded under this section unless an appropriate proposal and application * * * are submitted to, and approved by, the appropriate administering official."

Respondents: State governments.

ANNUAL BURDEN ESTIMATES

Instrument	Number of respondents	Number of responses per respondent	Average burden hours per response	Total burden hours
ORR-1	48	1	.5	24

Estimated Total Annual Burden Hours: 24.

Additional Information: Copies of the proposed collection may be obtained by writing to the Administration for Children and Families, 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: grjohnson@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, 725 17th Street, NW., Washington, DC 20503, Attn: Desk Officer for ACF, Katherine_T._Astrich@omb.eop.gov.

Dated: November 17, 2004.
Robert Sargis,
 Reports Clearance Officer.
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DEPARTMENT OF HEALTH AND HUMAN SERVICES

Administration for Children and Families

Proposed Information Collection Activity; Comment Request

Proposed Projects:

Title: Developmental Disabilities Protection and Advocacy Statement of Goals and Priorities.

OMB No.: 0980-0270.

Description: As required by Federal statute and regulation, each State Protection and Advocacy System must prepare and submit to public comment a Statement of Goals and Priorities (SGP). The final version of this SGP for the coming fiscal year is submitted to the Administration on Developmental Disabilities (ADD). The information in the SGP will be aggregated into a national prospective profile of where Protection and Advocacy Systems are going. It will provide ADD with a tool for monitoring the public input requirement. Further, it will provide an overview of program direction, and permit ADD to track accomplishments against goals/targets, permitting the formulation of technical assistance and