

*Estimated Total Annual Burden Hours: 5,417.40.*

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 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: April 28, 2009.  
**Janean Chambers,**  
*Reports Clearance Officer.*  
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**DEPARTMENT OF HEALTH AND HUMAN SERVICES**

**National Institutes of Health**

**Proposed Collection; Comment Request; Collection of Customer Service, Demographic, and Smoking/Tobacco Use Information From NCI Cancer Information Service (CIS) Clients (NCI)**

**SUMMARY:** 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 National Cancer Institute (NCI), the National Institutes of Health (NIH) will publish periodic summaries of proposed projects to be submitted to the Office of Management and Budget (OMB) for review and approval.

Proposed Collection: *Title:* Collection of Customer Service, Demographic, and Smoking/Tobacco Use Information from NCI Cancer Information Service (CIS) Clients. *Type of Information Collection Request:* Revision of currently approved collection 0925-0208 (expiration 09/30/2009). *Need and Use of Information Collection:* The National Cancer Institute's Cancer Information Service (CIS) provides the latest information on

cancer, clinical trials, and tobacco cessation in English and Spanish. Clients are served by calling 1-800-4-CANCER for cancer information; 1-877-44U-QUIT for smoking cessation services; and using the NCI's LiveHelp, a Web-based chat service. CIS currently conducts a brief survey of a sample of telephone and LiveHelp clients at the end of usual service—a survey that includes three customer service and twelve demographic questions (age, sex, race, ethnicity, education, household income, number in household, and five questions about health care/coverage). Characterizing clients and how they found out about the CIS is essential to customer service, program planning, and promotion. The NCI also conducts a survey of individuals using the CIS's smoking cessation services—a survey that includes 20 smoking/tobacco use "intake" questions that serve as a needs assessment that addresses smoking history, previous quit attempts, and motivations to quit smoking. An additional question is used with callers who want to receive proactive call-back services. Responses to these questions enable Information Specialists to provide effective individualized counseling. *Frequency of Response:* Once. *Affected Public:* Individuals or households. *Type of Respondents:* People with cancer; their relatives and friends; and general public, including smokers/tobacco users. Annualized estimates for numbers of respondents and respondent burden are presented in Table 1.

TABLE 1—ESTIMATE OF ANNUAL BURDEN HOURS

Type of respondents	Survey instrument	Number of respondents	Frequency of responses	Average time per response (minutes/hour)	Annual burden hours
Telephone Clients: <sup>1</sup>	Customer Service .....	62,000	1	1/60	1,033.33
	Demographic Questions .....	22,000	1	2/60	733.33
Smoking Cessation "Quitline" Clients: <sup>1,2</sup>					
Reactive Service Clients .....	Smoking Cessation "Intake" Questions.	4,641	1	5/60	386.75
Proactive Callback Service Clients <sup>3</sup>	Demographic Questions .....	1,300	1	2/60	43.33
	Follow-Up .....	928	4	1/60	61.87
LiveHelp Clients: <sup>4</sup>	Demographic questions .....	7,014	1	2/60	233.80
Total .....		97,883			2524.00

<sup>1</sup> Approximately 36% of telephone and quitline clients will be sampled for the demographic questions, and 100% of telephone clients will be sampled for the customer service questions. Estimates based on 77.5% response rate.

<sup>2</sup> 100% of smoking cessation clients will be asked the smoking intake questions. Estimates for quitline callers answering demographic questions are based on 77.8% response rate.

<sup>3</sup> 100% of smoking cessation clients participating in the proactive callback service (about 20% of all smoking callers) will be asked the smoking follow-up question (at up to 4 callbacks).

<sup>4</sup> Approximately 50% of LiveHelp clients will be sampled for the demographic questions.

The annualized cost to the respondents is estimated at \$48,752.

There are no Capital Costs, Operating

Costs, and/or Maintenance Costs to report.

Request For Comments: Written comments and/or suggestions from the public and affected agencies should address one or more of the following points: (1) Evaluate whether the proposed collection of information is necessary for the proper performance of the function of the agency, including whether the information will have practical utility; (2) Evaluate the accuracy of the agency's estimate of the burden of the proposed collection of information, including the validity of the methodology and assumptions used; (3) Enhance the quality, utility, and clarity of the information to be collected; and (4) Minimize the burden of the collection of information on those who are to respond, including the use of appropriate automated, electronic, mechanical, or other technological collection techniques or other forms of information technology.

**FOR FURTHER INFORMATION CONTACT:** To request more information on the proposed project or to obtain a copy of the data collection plans and instruments, contact Mary Anne Bright, Associate Director, Office of Public Information and Resource Management, Office of Communications and Education, National Cancer Institute, 6116 Executive Blvd., Room 3049, MSC 8322, Bethesda, MD 20892-8322 or call 301-594-9048 or e-mail your request, including your address, to: [brightma@mail.nih.gov](mailto:brightma@mail.nih.gov).

Comments Due Date: Comments regarding this information collection are best assured of having their full effect if received within 60 days of the date of this publication.

Dated: April 23, 2009.

**Vivian Horovitch-Kelley,**

*NCI Project Clearance Liaison, National Institutes of Health.*

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## DEPARTMENT OF HEALTH AND HUMAN SERVICES

### Centers for Medicare & Medicaid Services

[Document Identifier: CMS-10284 and CMS-2567]

### Agency Information Collection Activities: Proposed Collection; Comment Request

**AGENCY:** Centers for Medicare & Medicaid Services, Department of Health and Human Services.

In compliance with the requirement of section 3506(c)(2)(A) of the Paperwork Reduction Act of 1995, the

Centers for Medicare & Medicaid Services (CMS) is publishing the following summary of proposed collections for public comment. Interested persons are invited to send comments regarding this burden estimate or any other aspect of this collection of information, including any of the following subjects: (1) The necessity and utility of the proposed information collection for the proper performance of the agency's functions; (2) the accuracy of the estimated burden; (3) ways to enhance the quality, utility, and clarity of the information to be collected; and (4) the use of automated collection techniques or other forms of information technology to minimize the information collection burden.

1. *Type of Information Collection Request:* New Collection; *Title of Information Collection:* Children's Health Insurance Program Reauthorization Act (CHIPRA) of 2009, State Option Pre-print to Include Pregnant Women in Title XXI; *Use:* Section 111 of CHIPRA adds a new section 2112 to the Social Security Act which gives States the option of providing necessary prenatal, delivery and postpartum care to low-income uninsured pregnant women through an amendment to its State Child Health Plan (CHIP plan). The purpose of this draft State plan amendment template is to provide States with the format needed to enable a State to amend their CHIP plan to reflect the coverage of pregnant women. *Form Number:* CMS-10284 (OMB#: 0938-NEW); *Frequency:* Reporting—One-time and Occasionally; *Affected Public:* State, Local or Tribal Government; *Number of Respondents:* 40; *Total Annual Responses:* 40; *Total Annual Hours:* 3,200. (For policy questions regarding this collection contact Meredith Robertson at 410-786-6543. For all other issues call 410-786-1326.)

2. *Type of Information Collection Request:* Reinstatement without change of a previously approved collection; *Title of Information Collection:* Statement of Deficiencies and Plan of Correction; *Use:* The information from the CMS-2567 is used by the States and CMS regional offices to document and certify compliance. *Form Number:* CMS-2567 (OMB#: 0938-0391); *Frequency:* Reporting—Annually; *Affected Public:* State, Local or Tribal Government, Federal Government, Business or other for-profits and Not-for-profit Institutions; *Number of Respondents:* 60,000; *Total Annual Responses:* 60,000; *Total Annual Hours:* 120,000. (For policy questions regarding this collection contact Joanne Perry at

410-786-3336. For all other issues call 410-786-1326.)

To obtain copies of the supporting statement and any related forms for the proposed paperwork collections referenced above, access CMS' Web Site at <http://www.cms.hhs.gov/PaperworkReductionActof1995>, or E-mail your request, including your address, phone number, OMB number, and CMS document identifier, to [Paperwork@cms.hhs.gov](mailto:Paperwork@cms.hhs.gov), or call the Reports Clearance Office on (410) 786-1326.

In commenting on the proposed information collections please reference the document identifier or OMB control number. To be assured consideration, comments and recommendations must be submitted in one of the following ways by June 30, 2009:

1. *Electronically.* You may submit your comments electronically to <http://www.regulations.gov>. Follow the instructions for "Comment or Submission" or "More Search Options" to find the information collection document(s) accepting comments.

2. *By regular mail.* You may mail written comments to the following address:

CMS, Office of Strategic Operations and Regulatory Affairs, Division of Regulations Development, Attention: Document Identifier/OMB Control Number (CMS-10283), Room C4-26-05, 7500 Security Boulevard, Baltimore, Maryland 21244-1850.

Dated: April 23, 2009.

**Michelle Shortt,**

*Director, Regulations Development Group, Office of Strategic Operations and Regulatory Affairs.*

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## DEPARTMENT OF HEALTH AND HUMAN SERVICES

### Administration for Children and Families

### Submission for OMB Review; Comment Request

*Title:* Evaluation of the Transitional Living Program (TLP).

*OMB No.:* New Collection.

*Description:* The Runaway and Homeless Youth Act (RHYA), as amended by Public Law 106-71 (42 U.S.C. 5701 *et seq.*), provides for the Transitional Living Program (TLP), a residential program lasting up to 18 months designed to prepare older homeless youth ages 16-21 for a healthy and self-sufficient adulthood. Section 119 of RHYA requires a study on the