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*.

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.

[FR Doc. E9–10012 Filed 4–30–09; 8:45 am]

BILLING CODE 4140–01–P

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*, 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.

[FR Doc. E9–9959 Filed 4–30–09; 8:45 am]

BILLING CODE 4120–01–P

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

long-term housing outcomes of youth after exiting the program. In addition to collecting information on housing outcomes, the study will also consider the living, employment, education, and family situation of the youth before and after their time in the TLP. This information will be used to better understand the most effective practices in improving long-term outcomes of youth in an effort to guide program improvements.

Respondents: (1) Youth ages 16–21 participating in Transitional Living Programs and (2) the Executive Director and Program Manager representing TLP grantees.

ANNUAL BURDEN ESTIMATES

Instrument	Number of respondents	Number of responses per respondent	Average burden hours per response	Total burden hours
Grantee Survey	70	1	1	70
Youth Baseline Survey	760	1	0.50	380
Youth Exit Survey	760	1	0.50	380
Youth 6-Month Follow Up	760	1	0.50	380
Youth 12-Month Follow Up	760	1	0.50	380
Service Log	760	1	0.25	190

Estimated Total Annual Burden Hours: 1,780.

Additional Information:

Copies of the proposed collection may be obtained 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. All requests should be identified by the title of the information collection. E-mail 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: 202–395–7245, Attn: Desk Officer for the Administration for Children and Families.

Dated: April 28, 2009.

Janean Chambers,

Reports Clearance Officer.

[FR Doc. E9–10020 Filed 4–30–09; 8:45 am]

BILLING CODE 4184–01–P

DEPARTMENT OF HEALTH AND HUMAN SERVICES

Food and Drug Administration

[Docket No. FDA–2009–D–0181]

Draft Guidance for Industry on Label Comprehension Studies for Nonprescription Drug Products; Availability

AGENCY: Food and Drug Administration, HHS.

ACTION: Notice.

SUMMARY: The Food and Drug Administration (FDA) is announcing the availability of a draft guidance for industry entitled ''Label Comprehension Studies for Nonprescription Drug Products.'' The draft guidance provides recommendations on the design of label comprehension studies, which can be used to assess the extent to which consumers understand the information conveyed by proposed nonprescription drug product labeling and then apply that information when making hypothetical drug product use decisions.

DATES: Although you can comment on any guidance at any time (see 21 CFR 10.115(g)(5)), to ensure that the agency considers your comment on this draft guidance before it begins work on the final version of the guidance, submit written or electronic comments on the draft guidance by July 30, 2009.

ADDRESSES: Submit written requests for single copies of the draft guidance to the Division of Drug Information, Center for Drug Evaluation and Research, Food and Drug Administration, 10903 New Hampshire Ave., Bldg. 51, rm. 2201, Silver Spring, MD 20993–0002. Send one self-addressed adhesive label to assist that office in processing your requests. Submit written comments on the draft guidance to the Division of Dockets Management (HFA–305), Food and Drug Administration, 5630 Fishers Lane, rm. 1061, Rockville, MD 20852. Submit electronic comments to *http://www.regulations.gov*. See the **SUPPLEMENTARY INFORMATION** section for electronic access to the draft guidance document.

FOR FURTHER INFORMATION CONTACT: Laura Shay, Center for Drug Evaluation and Research, Food and Drug Administration, 10903 New Hampshire Ave., Bldg. 22, rm. 5466, Silver Spring, MD 20993–0002, 301–796–0994.

SUPPLEMENTARY INFORMATION:

I. Background

FDA is announcing the availability of a draft guidance for industry entitled ''Label Comprehension Studies for Nonprescription Drug Products.'' This draft guidance is intended for individuals or organizations involved in the development of label comprehension studies for nonprescription drug products. This draft guidance discusses general concepts to be considered in the design and conduct of a label comprehension study. This draft guidance also incorporates advice obtained from the September 25, 2006, meeting of the Nonprescription Drug Advisory Committee that considered issues related to the analysis and interpretation of consumer behavior studies conducted to support marketing of nonprescription drug products.

This draft guidance is being issued consistent with FDA's good guidance practices regulation (21 CFR 10.115). The draft guidance, when finalized, will represent the agency's current thinking on label comprehension studies for nonprescription drug products. It does not create or confer any rights for or on any person and does not operate to bind