official rules posted on the challenge site, vetoviolence.challenge.gov. Prior to entering a submission to the challenge, contestants must follow the challenge rules before the end of the submission period.

Amount of the Prize: One prize winner for each category, including General Public View, Student View, and Violence Prevention Professional View, will receive an award in the amount of $500 after the notification of the winners. A total of $1,500 will be distributed as awards by the contractor.

Payment of the Prize: Prizes under this competition will be paid by electronic funds transfer by Westat Health Communications as part of their VetoViolence Facebook contract with the HHS/CDC Injury Center's Division of Violence Prevention.

Basis Upon Which Winners Will be Selected: Submissions to the challenge will be assessed by a panel of judges composed of HHS/CDC Injury Center teen dating violence subject matter experts and communications staff and external injury and violence professionals in compliance with the requirements of the America COMPETES Act. Judges will be named after the commencement of the challenge on July 15, 2013. The judging panel will make decisions based on the following criteria:

(1) Creativity: Each entry will be judged on creativity demonstrated in the delivery of teen dating violence prevention messages.

(2) Communication of teen dating violence prevention messages: Each entry will be judged on the expression of positive prevention of teen dating violence messages. The submissions should not contain real or simulated acts of violence, profane language, inappropriate content, or personal or professional attacks.

(3) Length of Video: Each entry should be 60 seconds or less.

(4) Video and Audio Quality: Each entry should be visually focused and have audible sound quality. Submissions should not be difficult to watch because of an unclear image or to hear because of a poor audio recording.

(5) Fulfilling contest purpose: Each entry will be judged on its overall success in meeting the contest goal: Development of video public service announcements (PSA) that increase the understanding (1) that teen dating violence is a public health problem and (2) that prevention efforts can stop it before it starts.

One prize winner for each category—General Public View, Student View, and Violence Prevention Professional View—will receive an award in the amount of $500 after the notification of the winners. A total of $1,500 will be distributed among the three winners.

Additional Information: Finalists and the contest winners must comply with all terms and conditions of the official rules posted on the challenge site, *vetoviolence.challenge.gov,* and winning is contingent upon fulfilling all requirements herein. The finalists will be notified by email, telephone, or mail after the date of the judging.

Contestant information provided during registration will be used to respond to contestants in matters regarding their submission, announcements of entrants, finalists, and winners of the contest. Information is not collected for commercial marketing. Winners are permitted to cite that they won this contest.

HHS/CDC reserves the right to cancel, suspend, and/or modify the contest, or any part of it, for any reason, at HHS/CDC's sole discretion.

More information on teen dating violence may be found at *http://www.cdc.gov/violenceprevention/intimatepartnerviolence/teen_dating_violence.html.* More information on VetoViolence may be found at *http://vetoviolence.cdc.gov/* and *http://www.facebook.com/vetoviolence.*

Dated: July 5, 2013.

Tanja Popovic,

Deputy Associate Director for Science, Centers for Disease Control and Prevention.

[FR Doc. 2013–16619 Filed 7–10–13; 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

Title: Evaluation of the Transitional Living Program (TLP)

OMB No.: 0970–0383

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	15	1	1	15
Youth Baseline Survey	1250	1	0.75	937.50
Youth 6-Month Follow Up	1250	1	0.33	412.50
Youth 12-Month Follow Up	1250	1	0.33	412.50
Youth 18-Month Follow Up	1250	1	0.75	937.50

Estimated Total Annual Burden Hours: 2,715.

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. 2013–16643 Filed 7–10–13; 8:45 am]

BILLING CODE 4184–01–P

DEPARTMENT OF HEALTH AND HUMAN SERVICES

Food and Drug Administration

[Docket No. FDA–2013–N–0115]

Agency Information Collection Activities; Submission for Office of Management and Budget Review; Comment Request; Manufactured Food Regulatory Program Standards

AGENCY: Food and Drug Administration, HHS.

ACTION: Notice.

SUMMARY: The Food and Drug Administration (FDA) is announcing that a proposed collection of information has been submitted to the Office of Management and Budget (OMB) for review and clearance under the Paperwork Reduction Act of 1995.

DATES: Fax written comments on the collection of information by August 12, 2013.

ADDRESSES: To ensure that comments on the information collection are received, OMB recommends that written comments be faxed to the Office of Information and Regulatory Affairs, OMB, Attn: FDA Desk Officer, FAX: 202–395–7285, or emailed to *oira_submission@omb.eop.gov.* All comments should be identified with the OMB control number 0910–0601. Also include the FDA docket number found in brackets in the heading of this document.

FOR FURTHER INFORMATION CONTACT: Ila S. Mizrachi, Office of Information Management, Food and Drug Administration, 1350 Piccard Dr., PI50–400B, Rockville, MD 20850, 301–796–7726, *Ila.Mizrachi@fda.hhs.gov*.

SUPPLEMENTARY INFORMATION: In compliance with 44 U.S.C. 3507, FDA has submitted the following proposed collection of information to OMB for review and clearance.

Manufactured Food Regulatory Program Standards—(OMB Control Number 0910–0601)—Extension

In the **Federal Register** of July 20, 2006 (71 FR 41221), FDA announced the availability of a draft document entitled ''Manufactured Food Regulatory Program Standards (MFRPS).'' These draft program standards are the framework that States should use to design and manage its manufactured food program. The implementation of the standards will be negotiated as an option for payment under the State food contract. States that are awarded this option will receive up to $25,000 over a period of 5 years to fully implement the program standards. Additionally, 26 States may receive up to $300,000 each year for a period of 5 years to be in compliance with the 10 standards.

In the first year of implementing the program standards, the State program conducts a baseline self-assessment to determine if they meet the elements of each standard. The State program should use the worksheets and forms contained herein; however, it can use alternate forms that are equivalent. The State program maintains the documents and verifying records required for each standard. The information contained in the documents must be current and fit-for-use. If the State program fails to meet all program elements and documentation requirements of a standard, it develops a strategic plan which includes the following: (1) The individual element of documentation requirement of the standard that was not met; (2) improvements needed to meet the program element or documentation requirement of the standard; and (3) projected completion dates for each task.

In the **Federal Register** of February 19, 2013 (78 FR 11651), FDA published a 60-day notice requesting public comment on the proposed collection of information. No comments were received.

FDA estimates the burden of this collection of information as follows:

TABLE 1—ESTIMATED ANNUAL REPORTING BURDEN [1]

Respondent	Number of respondents	Number of responses per respondent	Total annual responses	Average burden per response	Total hours
State Departments of Agriculture or Health	44	1	44	303	13,332

[1] There are no capital costs or operating and maintenance costs associated with this collection of information.

The burden has been calculated to 303 hours per respondent. This burden was determined by capturing the average amount of time for each respondent to assess the current state of the program and work toward implementation of each of the 10 standards contained in MFRPS. The hours per respondent will remain the same as implementation to account for continuing improvement and self-sufficiency in the program.

Dated: July 5, 2013.

Leslie Kux,

Assistant Commissioner for Policy.

[FR Doc. 2013–16620 Filed 7–10–13; 8:45 am]

BILLING CODE 4160–01–P