In compliance with the requirements of Section 3506(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. This information collection is a supplement to the Maternal, Infant and Early Childhood Home Visiting Evaluation collection described in a 60 day **Federal Register** Notice, published on December 12, 2011 (Volume 76, No. 238, Page 77236). Per OMB guidance, ACF requests comments on this supplemental information collection within 30 days of this publication. Comments on and requests for copies of the proposed information collection may be forwarded by writing to the Administration for Children and Families, Office of Planning, Research and Evaluation, 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 30 days of this publication.

Steven M. Hanmer,

Reports Clearance Officer.

[FR Doc. 2012–15796 Filed 6–27–12; 8:45 am]

BILLING CODE 4184–22–M

DEPARTMENT OF HEALTH AND HUMAN SERVICES

Administration for Children and Families

Proposed Information Collection Activity; Comment Request

Proposed Projects

Title: Plan for Foster Care and Adoption Assistance—Title IV–E.

OMB No.: 0980–0141.

Description: A title IV–E plan is required by section 471, part IV–E of the Social Security Act (the Act) for each public child welfare agency requesting Federal funding for foster care, adoption assistance and guardianship assistance under the Act. Section 479B of the Act provides for an Indian tribe, tribal organization or tribal consortium (Tribe) to operate a title IV–E program in the same manner as a State with minimal exceptions. The Tribe must have an approved title IV–E Plan. The title IV–E plan provides assurances the programs will be administered in conformity with the specific requirements stipulated in title IV–E. The plan must include all applicable State or Tribal statutory, regulatory, or policy references and citations for each requirement as well as supporting documentation. A title IV–E agency may use the pre-print format prepared by the Children's Bureau of the Administration for Children and Families or a different format, on the condition that the format used includes all of the title IV–E plan requirements of the law.

Respondents: Title IV–E agencies administering or supervising the administration of the title IV–E programs.

ANNUAL BURDEN ESTIMATES

Instrument	Number of respondents	Number of responses per respondent	Average burden hours per response	Total burden hours
Title IV–E Plan	17	1	16	272

Estimated Total Annual Burden Hours: 272.

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 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. 2012–15770 Filed 6–27–12; 8:45 am]

BILLING CODE 4184–01–P

DEPARTMENT OF HEALTH AND HUMAN SERVICES

Administration for Children and Families

Announcement of the Award of Single-Source Cooperative Agreement to Rubicon Programs, Inc., in Richmond, CA

AGENCY: Office of Family Assistance, ACF, HHS.

ACTION: Announcement of the award of a single-source cooperative agreement to Rubicon Programs, Inc, in Richmond, CA, to support Community-Centered Responsible Fatherhood Ex-Prisoner Reentry activities to promote responsible fatherhood, family reunification, and economic stability designed to move individuals and families to self-sufficiency.

CFDA Number: 93.086.

Statutory Authority: The award is made under the authority of Claims

Resettlement Act of 2010 (Pub. L. 111–291).

SUMMARY: The Administration for Children and Families (ACF), Office of Family Assistance (OFA), Division of State and Territory TANF Management (DSTTM) announces the award of a single-source cooperative agreement of $1,500,000 to Rubicon Programs, Inc., in Richmond, CA.

The cooperative agreement will support a demonstration pilot project for responsible fatherhood activities authorized by the Claims Resolution Act of 2010 (Pub. L. 111–291). The Community-Centered Responsible Fatherhood Ex-Prisoner Reentry Pilot Project supports programs that are designed to offer community-centered, pre- and post-release responsible fatherhood and supportive services to formerly incarcerated fathers. The primary purpose of the program is to eliminate barriers to social and economic self-sufficiency for individuals preparing to reenter their communities, or those who have recently returned to their communities following incarceration. The project will implement three legislatively specified activities: Healthy marriage, responsible parenting, and economic stability.

The project will implement a program that includes comprehensive case management to strengthen father, couple, and family relationships and that connect formerly incarcerated fathers to employment, housing (when necessary), and other needed support services to help reduce the likelihood of recidivism. It is expected that the full project period will be 24 months so that, based on performance; the recipient may receive an additional noncompetitive award in Fiscal Year 2013.

DATES: September 30, 2012–September 29, 2013.

FOR FURTHER INFORMATION CONTACT: Robin Y. McDonald, Division Director, Office of Family Assistance, 370 L'Enfant Promenade SW., 5th Floor East, Washington, DC 20047. Telephone: (202) 401–5587 Email: *robin.mcdonald@acf.hhs.gov.*

Earl S. Johnson,

Director, Office of Family Assistance, Administration for Children and Families.

[FR Doc. 2012–15783 Filed 6–27–12; 8:45 am]

BILLING CODE 4184–35–P

DEPARTMENT OF HEALTH AND HUMAN SERVICES

Food and Drug Administration

[Docket No. FDA–2012–N–0021]

Agency Information Collection Activities; Submission for Office of Management and Budget Review; Comment Request; Substances Generally Recognized as Safe; Notification Procedure

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 July 30, 2012.

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 0910–0342. Also include the FDA docket number found in brackets in the heading of this document.

FOR FURTHER INFORMATION CONTACT: Domini Bean, Office of Information Management, Food and Drug Administration, 1350 Piccard Dr., PI50–400T, Rockville, MD 20850, 301–796–5733, *domini.bean@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.

Substances Generally Recognized as Safe: Notification Procedure—21 CFR 170.36 and 570.36 (OMB Control Number 0910–0342)—Revision

I. Background

Section 409 of the Federal Food, Drug, and Cosmetic Act (the FD&C Act) (21 U.S.C. 348) establishes a premarket approval requirement for ''food additives;'' section 201(s) of the FD&C Act (21 U.S.C. 321) provides an exemption from the definition of ''food additive'' and thus from the premarket approval requirement, for uses of substances that are GRAS by qualified experts. In the **Federal Register** of April 17, 1997 (62 FR 18938) (the 1997 proposed rule), FDA published a proposed rule that would establish a voluntary procedure whereby manufacturers would notify FDA about a view that a particular use (or uses) of a substance is not subject to the statutory premarket approval requirements based on a determination that such use is GRAS. The proposed regulations (proposed 21 CFR 170.36 and 21 CFR 570.36) provide a standard format for the voluntary submission of a notice. The notice would include a detailed summary of the data and information that support the GRAS determination, and the notifier would maintain a record of such data and information. FDA would make the information describing the subject of the GRAS notice, and the Agency's response to the notice, available in a publicly accessible file; the entire GRAS notice would be publicly available consistent with the Freedom of Information Act and other Federal disclosure statutes. In the **Federal Register** of December 28, 2010 (75 FR 81536) (the GRAS reopener), FDA announced the reopening of the comment period for the 1997 proposed rule. The Agency requested that comments be submitted by March 28, 2011.

FDA's Center for Food Safety and Applied Nutrition (CFSAN) has recently developed a form that prompts a notifier to include certain elements of a GRAS notice in a standard format. New Form FDA 3667 is entitled ''Generally Recognized as Safe (GRAS) Notice.'' The form, and elements that would be prepared as attachments to the form, may be submitted in electronic format via the Electronic Submissions Gateway (ESG), or may be submitted in paper format, or as electronic files on physical media with paper signature page. CFSAN expects that most if not all businesses filing GRAS notices in the next 3 years will choose to take advantage of the option of electronically submitting their GRAS notice. Thus, the burden estimate in Table 1, line 1 is based on the expectation of 100 percent participation in the electronic submission process.

FDA's Center for Veterinary Medicine (CVM) continues to comply with the GRAS Pilot Program procedures announced on June 4, 2010 (75 FR 31800).

II. GRAS Information on Form FDA 3667

The GRAS notice submitted to CFSAN includes the following information on Form FDA 3667 and in attachments to the form: