available to Medicare beneficiaries, as well as to the general public, to provide information to assist them in making decisions about their health care. *Form Number:* CMS–10530 (OMB control number: 0938–NEW); *Frequency:* Annually; *Affected Public:* Business or other for-profits and not-for-profit institutions; *Number of Respondents:* 5,250; *Total Annual Responses:* 744,816; *Total Annual Hours:* 444,790. (For policy questions regarding this collection contact Anita Bhatia at 410–786–7236.)

2. *Type of Information Collection Request:* Extension without change of a currently approved collection. *Title of Information Collection:* Certification as a Supplier of Portable X-Ray and Portable X-Ray Survey Report Form and Supporting Regulations. *Use:* CMS–1880 is initially completed by suppliers of portable X-ray services, expressing an interest in and requesting participation in the Medicare program. This form initiates the process of obtaining a decision as to whether the conditions of coverage are met as a portable X-ray supplier. It also promotes data reduction or introduction to, and retrieval from, the Certification and Survey Provider Enhanced Reporting (CASPER) by the CMS Regional Offices (ROs).

The CMS–1882 is used by the State survey agency to provide data collected during an on-site survey of a supplier of portable X-ray services to determine compliance with the applicable conditions of participation and to report this information to the Federal Government. The form is primarily a coding worksheet designed to facilitate data reduction and retrieval into the ASPEN system at the CMS Regional Offices. The form includes basic information on compliance (i.e., met, not met, explanatory statements) and does not require any descriptive information regarding the survey activity itself. We have the responsibility and authority for certification decisions which are based on supplier compliance with the applicable conditions of participation. The information needed to make these decisions is available to us only through the use of information abstracted from the survey report form. *Form Numbers:* CMS–1880 and CMS–1882 (OMB control number: 0938–0027); *Frequency:* Occasionally; *Affected Public:* State, Local, or Tribal Governments; *Number of Respondents:* 579; *Total Annual Responses:* 86; *Total Annual Hours:* 151. (For policy questions regarding this collection contact James Cowher at 410–786–1948.)

Dated: November 12, 2014.

Martique Jones,

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

[FR Doc. 2014–27137 Filed 11–14–14; 8:45 am]

BILLING CODE 4120–01–P

DEPARTMENT OF HEALTH AND HUMAN SERVICES

Centers for Medicare & Medicaid Services

[Document Identifier CMS–10422]

Agency Information Collection Activities: Submission for OMB Review; Comment Request

ACTION: Notice.

SUMMARY: The Centers for Medicare & Medicaid Services (CMS) is announcing an opportunity for the public to comment on CMS' intention to collect information from the public. Under the Paperwork Reduction Act of 1995 (PRA), federal agencies are required to publish notice in the **Federal Register** concerning each proposed collection of information, including each proposed extension or reinstatement of an existing collection of information, and to allow a second opportunity for public comment on the notice. Interested persons are invited to send comments regarding the 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.

DATES: Comments on the collection(s) of information must be received by the OMB desk officer by December 17, 2014.

ADDRESSES: When commenting on the proposed information collections, please reference the document identifier or OMB control number. To be assured consideration, comments and recommendations must be received by the OMB desk officer via one of the following transmissions: OMB, Office of Information and Regulatory Affairs, Attention: CMS Desk Officer, Fax Number: (202) 395–5806 or Email: *OIRA_submission@omb.eop.gov.*

To obtain copies of a supporting statement and any related forms for the proposed collection(s) summarized in this notice, you may make your request using one of following:

1. Access CMS' Web site address at *http://www.cms.hhs.gov/PaperworkReductionActof1995.*

2. Email your request, including your address, phone number, OMB number, and CMS document identifier, to *Paperwork@cms.hhs.gov.*

3. Call the Reports Clearance Office at (410) 786–1326.

FOR FURTHER INFORMATION CONTACT: Reports Clearance Office at (410) 786–1326.

SUPPLEMENTARY INFORMATION: Under the Paperwork Reduction Act of 1995 (PRA) (44 U.S.C. 3501–3520), federal agencies must obtain approval from the Office of Management and Budget (OMB) for each collection of information they conduct or sponsor. The term ''collection of information'' is defined in 44 U.S.C. 3502(3) and 5 CFR 1320.3(c) and includes agency requests or requirements that members of the public submit reports, keep records, or provide information to a third party. Section 3506(c)(2)(A) of the PRA (44 U.S.C. 3506(c)(2)(A)) requires federal agencies to publish a 30-day notice in the **Federal Register** concerning each proposed collection of information, including each proposed extension or reinstatement of an existing collection of information, before submitting the collection to OMB for approval. To comply with this requirement, CMS is publishing this notice that summarizes the following proposed collection(s) of information for public comment:

1. *Type of Information Collection Request:* Revision of a currently approved collection; *Title of Information Collection:* Payments for Services Furnished by Certain Primary Care Providers and Supporting Regulations in 42 CFR 438.804, 447.400, and 447.410; *Use:* The information will be used to document expenditures for the specified primary care services in the baseline period for the purpose of then calculating the expenditure eligible for 100 federal matching funds in calendar years 2015 and 2016, should Congress extend the availability of such funding and make no additional changes in statutory language necessitating programmatic alterations. *Form Number:* CMS–10422 (OMB control number: 0938–1170); *Frequency:* Yearly, once, and occasionally; *Affected Public:* State, Local, or Tribal Governments; *Number of Respondents:* 51; *Total Annual Responses:* 126,021; *Total Annual Hours:* 63,240. (For policy questions regarding this collection

contact Linda Tavener at 410–786–3838).

Dated: November 12, 2014.

Martique Jones,

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

[FR Doc. 2014–27135 Filed 11–14–14; 8:45 am]

BILLING CODE 4120–01–P

DEPARTMENT OF HEALTH AND HUMAN SERVICES

Administration for Children and Families

Announcing the Award of a Single-Source Program Expansion Supplement Grant to the Futures Without Violence in San Francisco, CA

AGENCY: Family and Youth Services Bureau, ACYF, ACF, HHS.

ACTION: Notice of the award of a single-source program expansion supplement grant under the Family Violence Prevention and Services Act (FVPSA) Technical Assistance (TA) Project to the Futures Without Violence to support training and technical assistance activities.

CFDA Number: 93.592.

SUMMARY: The Administration for Children and Families (ACF), Administration on Children, Youth and Families (ACYF), Family and Youth Services Bureau (FYSB), Division of Family Violence Prevention and Services (DFVPS) announces the award of $270,000 as a single-source program expansion supplement to Futures Without Violence in San Francisco, CA. The grantee, funded under the Family Violence Protection and Services Act (FVPSA) program, is a technical assistance (TA) provider that serves as the FVPSA-funded National Health Resource Center on Domestic Violence.

DATES: The period of support is September 30, 2014 through September 29, 2015.

FOR FURTHER INFORMATION CONTACT: Shawndell Dawson, Senior Program Specialist, Family Violence Prevention and Services Program, 1250 Maryland Avenue SW., Suite 8219, Washington, DC 20024. Telephone: 202–205–1476; Email: *Shawndell.Dawson@acf.hhs.gov.*

SUPPLEMENTARY INFORMATION: Supplemental award funds will support the grantee in providing training and technical assistance to domestic violence service and health care providers. A portion of the supplemental award is contributed by the Health Resources and Services Administration (HRSA) and the Office on Women's Health (OWH) at the Department of Health and Human Services (HHS).

This award will expand the scope of Futures Without Violence's technical assistance activities to include additional activities on the following issues: Assessing and responding to domestic violence in health clinics; addressing dating violence and sexual assault on college campuses; and supporting children/youth experiencing domestic violence. This additional technical assistance and training may involve such activities as:

- Planning, coordinating, and evaluating a pre-conference institute on Sexual Assault and Dating Violence on College Campuses, as part of the 2015 National Conference on Health and Domestic Violence;
- providing technical assistance for three health centers to create health system changes that support providers and create sustainable responses to victims of intimate partner violence;
- providing training on comprehensive, culturally competent responses to domestic violence within a Patient Centered Medical Home model.
- creating new technical assistance resources that promote protective factors and resilience when working with children, youth, and teens impacted by domestic violence which includes fostering stronger relationships with their non-abusive parents or caregivers;
- providing training to domestic violence programs that improves consistent implementation of evidence-informed, trauma-informed, and culturally relevant programming for children, youth, and abused parents; and,
- developing new resources for the Web site, *www.PromisingFutures WithoutViolence.org.*

Statutory Authority: The statutory authority for the FVSPA Program is under section 310 of the FVPSA, as amended by Section 201 of the CAPTA Reauthorization Act of 2010, Pub. L. 111–320. The Office on Women's Health authority for its additional funds is through Sections 1701(a)(3)(A), 1701(a)(5), and 1701(a)(8) of the Public Health Service Act; and the Economy Act (31 U.S.C. 1535/FAR 17.5). HRSA's authority for its funds is through Section 330 of the Public Health Service Act (42 U.S.C. § 254b).

Christopher Beach,

Senior Grants Policy Specialist, Office of Administration, Office of Financial Services, Division of Grants Policy.

[FR Doc. 2014–27131 Filed 11–14–14; 8:45 am]

BILLING CODE 4184–32–P

DEPARTMENT OF HEALTH AND HUMAN SERVICES

Food and Drug Administration

[Docket No. FDA–2014–D–1461]

Rare Pediatric Disease Priority Review Vouchers, Draft Guidance for Industry; 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 ''Rare Pediatric Disease Priority Review Vouchers.'' Under the Federal Food, Drug, and Cosmetic Act (the FD&C Act), FDA will award priority review vouchers to sponsors of certain rare pediatric disease product applications that meet the criteria specified in that section. These vouchers can be used when submitting future human drug marketing applications that would not otherwise qualify for priority review. These vouchers can be sold or transferred for use to another sponsor any number of times before the voucher is used, as long as the sponsor making the transfer has not yet submitted the application. Because there exists a need for products for rare pediatric diseases, this program is intended to encourage development of new drug and biological products for prevention and treatment of certain rare pediatric diseases.

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 either electronic or written comments on the draft guidance by January 16, 2015. Submit either electronic or written comments concerning the collection of information proposed in the draft guidance by January 16, 2015.

ADDRESSES: Submit written requests for single copies of the draft guidance to the Office of Communications, 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, or Office of Communication, Outreach, and Development, Center for Biologics Evaluation and Research, Food and Drug Administration, Bldg. 71, Rm. 3128, 10903 New Hampshire Ave., Silver Spring, MD 20993–0002; or Office of Orphan Products Development, Office of Special Medical Programs,