receiving access to the Medicare eligibility information for the purpose of conducting real-time 270/271 inquiry/response transactions. *Form Number:* CMS–10157 (OMB control number: 0938–0960); *Frequency:* Yearly; *Affected Public:* Private Sector; Business or other for profits, Not-for-Profits Institutions; *Number of Respondents:* 1000; *Total Annual Responses:* 1000; *Total Annual Hours:* 250. (For policy questions regarding this collection contact Rupinder Singh at 410 786–7484.)

Dated: May 21, 2019.

William N. Parham, III,

Director, Paperwork Reduction Staff, Office of Strategic Operations and Regulatory Affairs.

[FR Doc. 2019–10964 Filed 5–23–19; 8:45 am]

BILLING CODE 4120–01–P

DEPARTMENT OF HEALTH AND HUMAN SERVICES

Administration for Children and Families

[CFDA Number: 93.647]

Announcement of an Unsolicited Single-Source Grant Award to the Woodson Center in Washington, DC

AGENCY: Office of Planning, Research and Evaluation, Administration for Children and Families, Department of Health and Human Services.

ACTION: Notice of issuance.

SUMMARY: The Administration for Children and Families (ACF), Office of Planning, Research and Evaluation announces the award of a grant in the amount of $150,000 to the Woodson Center of Washington, DC, to support an environmental scan of models of service for privately managed foster care.

DATES: The project period will be May 15, 2019, to November 14, 2019.

FOR FURTHER INFORMATION CONTACT: Wendy DeCourcey, Senior Social Science Analyst; Office of Planning, Research and Evaluation; 330 C Street SW, Washington, DC 20201. Telephone: 202–260–2039.

SUPPLEMENTARY INFORMATION: The Woodson Center plans to complete an environmental scan of privately managed foster care systems in multiple states. This environmental scan is expected to inform future networking efforts among similar organizations and identification and dissemination of best practices.

Statutory Authority: Social Security Act, Title XI, Section 1110, 42 U.S.C. 1310.

Elizabeth Leo,

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

[FR Doc. 2019–10897 Filed 5–23–19; 8:45 am]

BILLING CODE 4184–07–P

DEPARTMENT OF HEALTH AND HUMAN SERVICES

Administration for Children and Families

Proposed Information Collection Activity; Extension of Assets for Independence (AFI) Performance Progress Report (PPR) (OMB #0970–0483)

AGENCY: Office of Community Services; Administration for Children and Families; HHS.

ACTION: Request for public comment.

SUMMARY: The Office of Community Services (OCS), Administration for Children and Families (ACF) is requesting approval of a three-year extension of the Assets for Independence (AFI) Performance Progress Report (PPR) Long Form and AFI PPR Short Form (OMB #0970–0483, expiration 8/31/2019). There are no changes requested to the forms.

DATES: *Comments due within 60 days of publication.* 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.

ADDRESSES: Copies of the proposed collection of information can be obtained and comments may be forwarded by emailing *infocollection@acf.hhs.gov.* Alternatively, copies can also be obtained by writing to the Administration for Children and Families, Office of Planning, Research, and Evaluation, 330 C Street SW, Washington, DC 20201, Attn: OPRE Reports Clearance Officer. All requests, emailed or written, should be identified by the title of the information collection.

SUPPLEMENTARY INFORMATION:

Description: The Assets for Independence (AFI) Act (Title IV of the Community Opportunities, Accountability, and Training and Educational Services Act of 1998, Pub. L. 105–285, [42 U.S.C. 604 note]) requires that organizations operating AFI projects submit annual progress reports.

This request is for approval and extension of the current AFI PPR that expires August 31, 2019. OCS will continue to use the data collected in the AFI PPR to prepare the annual AFI Report to Congress, to evaluate and monitor the performance of the AFI program overall and of individual projects, and to inform and support technical assistance efforts. The AFI PPR will continue to fulfill AFI Act reporting requirements and program purposes.

AFI program grantees are required to submit Standard Form Performance Progress Reports (SF–PPR) semiannually: One time per year using an abbreviated short form and one time using a long form. Both data collection instruments are available for review online at: *https://www.acf.hhs.gov/ocs/resource/afi-ppr-long-form*, *https://www.acf.hhs.gov/ocs/resource/afi-ppr-short-form.*

Note: This request does not affect financial reporting requirements for AFI grantees. The SF–425 will still be required semiannually throughout the grant project period with a final report due 90 days after the grant project period ends.

Respondents: Assets for Independence (AFI) program grantees.

ANNUAL BURDEN ESTIMATES

Instrument	Total number of respondents	Annual number of responses per respondent	Average burden hours per response	Annual burden hours
AFI PPR Short Form	145	1	0.5	72.5
AFI PPR Long Form	145	1	3.8	551

Estimated Total Annual Burden Hours: 623.5.

Comments: 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.

Authority: Pub. L. 105–285, [42 U.S.C. 604 note].

Mary B. Jones,

ACF/OPRE Certifying Officer.

[FR Doc. 2019–10863 Filed 5–23–19; 8:45 am]

BILLING CODE 4184–24–P

DEPARTMENT OF HEALTH AND HUMAN SERVICES

Food and Drug Administration

[Docket No. FDA–2019–N–1281]

General and Plastic Surgery Devices Panel of the Medical Devices Advisory Committee; Amendment of Notice

AGENCY: Food and Drug Administration, HHS.

ACTION: Notice.

SUMMARY: The Food and Drug Administration (FDA) is announcing an amendment to the notice of meeting of the General and Plastic Surgery Devices Panel of the Medical Devices Advisory Committee. This meeting was announced in the **Federal Register** of April 24, 2019. The amendment is being made to reflect a change in the **DATES** portion of the document. There are no other changes.

FOR FURTHER INFORMATION CONTACT: Patricio Garcia, Center for Devices and Radiological Health, Food and Drug Administration, 10903 New Hampshire Ave., Bldg. 66, Rm. G610, Silver Spring, MD 20993, 301–796–6875, *Patricio.garcia@fda.hhs.gov;* or FDA Advisory Committee Information Line, 1–800–741–8138 (301–443–0572 in the Washington, DC area). Please call the Information Line for up-to-date information on this meeting.

SUPPLEMENTARY INFORMATION: In the **Federal Register** of April 24, 2019 (84 FR 17173), FDA announced that a meeting of the General and Plastic Surgery Devices Panel of the Medical Devices Advisory Committee would be held on May 30, 2019, from 10 a.m. to 4 p.m. On page 17173, in the third column, in the **DATES** section, the sentence ''The meeting will be held on May 30, 2019, from 10 a.m. to 4 p.m. and on May 31, 2019, from 8 a.m. to 4 p.m.'' is changed to read as follows:

The meeting will be held on May 30, 2019, from 9 a.m. to 4 p.m. and on May 31, 2019, from 8 a.m. to 4 p.m.

This notice is issued under the Federal Advisory Committee Act (5 U.S.C. app. 2) and 21 CFR part 14, relating to the advisory committees.

Dated: May 21, 2019.

Lowell J. Schiller,

Principal Associate Commissioner for Policy.

[FR Doc. 2019–10900 Filed 5–23–19; 8:45 am]

BILLING CODE 4164–01–P

DEPARTMENT OF HEALTH AND HUMAN SERVICES

Food and Drug Administration

[Docket No. FDA–2019–N–2037]

Electronic Nicotine Delivery System Device and E-Liquid Manufacturer Site Tours Program

AGENCY: Food and Drug Administration, HHS.

ACTION: Notice.

SUMMARY: The Food and Drug Administration (FDA), Center for Tobacco Products (CTP), is announcing an invitation for participation in its voluntary Electronic Nicotine Delivery System (ENDS) Device and E-Liquid Manufacturer Site Tours Program. This program is intended to give CTP staff an opportunity to visit facilities that develop, manufacture, or test ENDS devices or e-liquids (including pods or cartridges) to gain a better understanding of the processes involved in the development, manufacturing, and testing of ENDS devices and e-liquids. The site tours in this program are not intended as regulatory inspections. The purpose of this document is to invite ENDS device or e-liquid manufacturers that can demonstrate assembly process and present supply chain information, and laboratories that conduct ENDS aerosol and e-liquid testing, that are interested in participating in the ENDS Device and E-Liquid Manufacturer Site Tours Program to submit requests to CTP.

DATES: Submit either an electronic or written request for participation in this program by July 23, 2019. See section IV of this document for information on requests for participation.

ADDRESSES: If your facility is interested in participating in a facility visit, please submit a request either electronically to *https://www.regulations.gov* or in writing to the Dockets Management Staff (HFA–305), Food and Drug Administration, 5630 Fishers Lane, Rm. 1061, Rockville, MD 20852.

FOR FURTHER INFORMATION CONTACT: Karla Price, Office of Science, Center for Tobacco Products, Food and Drug Administration, Document Control Center, 10903 New Hampshire Ave., Bldg. 71, Rm. G335, Silver Spring, MD 20993–0002, 1–877–287–1373, email: *AskCTP@fda.hhs.gov.*

SUPPLEMENTARY INFORMATION:

I. Background

On June 22, 2009, the Family Smoking Prevention and Tobacco Control Act (Pub. L. 111–31) (Tobacco Control Act) was signed into law, amending the Federal Food, Drug, and Cosmetic Act (FD&C Act) by, among other things, adding a new chapter (chapter IX) granting FDA the authority to regulate tobacco product manufacturing, distribution, and marketing. The Tobacco Control Act provides FDA authority to regulate cigarettes, cigarette tobacco, roll-your-own tobacco, smokeless tobacco, and any other tobacco products that the Agency by regulation deems to be subject to the law.

On May 10, 2016, FDA published a final rule entitled ''Deeming Tobacco Products To Be Subject to the Federal Food, Drug, and Cosmetic Act, as Amended by the Family Smoking Prevention and Tobacco Control Act; Restrictions on the Sale and Distribution of Tobacco Products and Required Warning Statements for Tobacco Products'' (81 FR 28974), which became effective on August 8, 2016. Under this rule, all products, such as ENDS, that meet the statutory definition of ''tobacco product'' set forth in section 201(rr) of the FD&C Act (21 U.S.C. 321(rr)), including components and parts, but excluding accessories of newly deemed products, are now subject to chapter IX of the FD&C Act.

CTP's Office of Science is conducting the ENDS Device and E-Liquid Manufacturer Site Tours Program to provide its staff an opportunity to visit facilities that develop, manufacture, or test ENDS devices or e-liquids (including pods or cartridges). The ENDS device and e-liquid facilities are regulated by FDA if they, among other things, manufacture products that meet the statutory definition of a ''tobacco product'' set forth in section 201(rr) of the FD&C Act. The site tours will aid the Agency in gaining a better understanding of the processes involved in developing, manufacturing, and