The Chief Operating Officer, Centers for Disease Control and Prevention, has been delegated the authority to sign Federal Register notices pertaining to announcements of meetings and other committee management activities, for both the Centers for Disease Control and Prevention and the Agency for Toxic Substances and Disease Registry.

Sherri A. Berger,
Chief Operating Officer, Centers for Disease Control and Prevention.

[FR Doc. 2019–15589 Filed 7–22–19; 8:45 am]
BILLING CODE 4163–18–P

DEPARTMENT OF HEALTH AND HUMAN SERVICES

Administration for Children and Families

Proposed Information Collection Activity: State Court Improvement Program (OMB # 0970–0307)

AGENCY: Children’s Bureau Administration for Children and Families; HHS.

ACTION: Request for public comment.

SUMMARY: The Administration for Children and Families (ACF) is requesting a three-year extension of the Court Improvement Program (CIP) Program Instruction, Strategic Plan Template, and Annual CIP Self-Assessment (OMB #0970–0307, expiration 8/31/2019). There are minimal updates to the form to reflect new legislation. The collections are necessary to continue operating the program in compliance with congressional reauthorization.

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.

ADDRESS: 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 proposed collection is a continuation of the current collection and comprised of two components: An application including a strategic plan that is due once every five years, and an annual self-assessment. The next annual self-assessment will be due June 30, 2020. The next five-year application will be due in 2021.

Respondents: We anticipate the highest state court of every state, Puerto Rico and the U.S. Virgin Islands to respond. All 52 jurisdictions currently participate in the program.

ANNUAL BURDEN ESTIMATES

<table>
<thead>
<tr>
<th>Instrument</th>
<th>Total number of respondents</th>
<th>Total number of responses per respondent</th>
<th>Average burden hours per response</th>
<th>Total burden hours</th>
<th>Annual burden hours</th>
</tr>
</thead>
<tbody>
<tr>
<td>5-year Application</td>
<td></td>
<td></td>
<td>52</td>
<td>1</td>
<td>92</td>
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<tr>
<td>Annual self-assessment</td>
<td></td>
<td></td>
<td>52</td>
<td>3</td>
<td>77</td>
</tr>
</tbody>
</table>

Estimated Total Annual Burden Hours: 4004 hours in 2020 and 2022; 8788 hours in 2021 (when both the self-assessment and the 5-year application are due within the year)

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: Sec. 50761, Pub. L. 115–123

Mary B. Jones,
ACF/OPRE Certifying Officer.

[FR Doc. 2019–15639 Filed 7–22–19; 8:45 am]
BILLING CODE 4184–01–P

DEPARTMENT OF HEALTH AND HUMAN SERVICES

Administration for Children and Families

Proposed Information Collection Activity: Plan for Foster Care and Adoption Assistance—Title IV-E (OMB #0970–0433)

AGENCY: Children’s Bureau, Administration for Children and Families, HHS.

ACTION: Request for public comment.

SUMMARY: The Administration for Children and Families (ACF) is requesting to revise the existing information collection Plan for Foster Care and Adoption Assistance (OMB #0970–0433) to include two new information collections specific to two new programs.

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.

ADDRESS: 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,
SUPPLEMENTARY INFORMATION: Title IV–E of the Social Security Act (the Act) was amended by Public Law 115–123, which included the Family First Prevention Services Act (FFPSA). The FFPSA authorized new optional title IV–E funding for time-limited (one year) prevention services for mental health/ substance abuse and in-home parent skill-based programs for: (1) A child who is a candidate for foster care (as defined in section 475(13) of the Act), (2) pregnant/parenting foster youth, and (3) the parents/kin caregivers of those children and youth (sections 471(e), 474(a)(6), and 475(13) of the Act). Title IV–E prevention services must be rated as promising, supported, or well-supported in accordance with HHS criteria and be approved by HHS (section 471(e)(4)(C) of the Act) as part of the Title IV–E Prevention Services Clearinghouse (section 476(d)(2) of the Act). A state or tribal title IV–E agency electing to participate in the program must submit a five-year title IV–E prevention program plan that meets the statutory requirements. (See Program Instructions ACYF–CB–PI–18–09 and ACYF–CB–PI–18–10 for more information.)

The FFPSA also amended Section 474(a)(7) of the Act to reimburse state and tribal IV–E agencies for a portion of the costs of operating kinship navigator programs that meet certain criteria. To qualify for funding under the title IV–E Kinship Navigator program, the program must meet the requirements of a kinship navigator program described in section 427(a)(1) of the Act. The kinship navigator program must also meet practice criteria of promising, supported, or well-supported in accordance with HHS criteria and be approved by HHS (section 471(e)(4)(C) of the Act). To begin participation in the title IV–E Kinship Navigator Program, a title IV–E agency must submit an attachment to its title IV–E plan that specifies the Kinship Navigator model it has chosen to implement, the date on which the provision of program services began or will begin, and that provides an assurance that the model meets the requirements of section 427(a)(1) of the Act as well as a brief narrative describing how the program will be operated. (Please see Program Instruction ACYF–CB–PI–18–11 for additional information.)

Respondents: State and tribal title IV–E agencies.

### ANNUAL BURDEN ESTIMATES

<table>
<thead>
<tr>
<th>Instrument</th>
<th>Total number of respondents</th>
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<th>Average burden hours per response</th>
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</tr>
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<td>5</td>
<td>150</td>
</tr>
<tr>
<td>Attachment to Title IV–E plan for Kinship navigator program</td>
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<td>1</td>
<td>1</td>
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</tr>
</tbody>
</table>

**Estimated Total Annual Burden Hours:** 195.

**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:** Title IV–E of the Social Security Act as amended by Public Law (Pub. L.) 115–123 enacted February 9, 2018.

**Mary B. Jones,**

ACF/OPRE Certifying Officer.

[FR Doc. 2019–15603 Filed 7–22–19; 8:45 am]

BILLING CODE 4184–25–P

### DEPARTMENT OF HEALTH AND HUMAN SERVICES

#### Food and Drug Administration

[Docket No. FDA–2019–N–0430]

**Agency Information Collection Activities; Submission for Office of Management and Budget Review; Comment Request; Generic Clearance for Quick Turnaround Testing of Communication Effectiveness**

**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 (PRA).

**DATES:** Fax written comments on the collection of information by August 22, 2019.

**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-New and title “Generic Clearance for Quick Turnaround Testing of Communication Effectiveness.” Also include the FDA docket number found in brackets in the heading of this document.

**FOR FURTHER INFORMATION CONTACT:** Ila S. Mizrachi, Office of Operations, Food and Drug Administration, Three White Flint North, 10A–12M, 11601 Landsdown St., North Bethesda, MD 20852, 301–796–7226, PHAStaff@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.

**Generic Clearance for Quick Turnaround Testing of Communication Effectiveness**

**OMB Control Number 0910–NEW**

This notice announces the FDA information collection request to OMB for approval of a generic clearance that will allow FDA to use quick turnaround surveys, focus groups, and in-depth interviews collected from consumers and other stakeholders to communicate FDA issues of immediate and important public health significance. For example,