dog from Egypt, including a dog from Egypt that is being imported from a third-party country. Such approvals will be granted on a limited and case-by-case basis and at CDC’s discretion.

Individuals seeking to import a dog from Egypt must submit the Application for a Permit to Import a Dog Inadequately Immunized Against Rabies, which is currently approved under OMB Control Number 0920–0134 Foreign Quarantine Regulations (exp. 03/31/2022).

To request the advance written approval of the CDC, you must send an approval to the Executive Secretary, Centers for Disease Control and Prevention.

Dated: May 6, 2019.

Sandra Cashman, Executive Secretary, Centers for Disease Control and Prevention.

[FR Doc. 2019–09654 Filed 5–9–19; 8:45 am]

DEPARTMENT OF HEALTH AND HUMAN SERVICES

Administration for Children and Families

Proposed Information Collection Activity: State Temporary Assistance for Needy Families Case Studies (New Collection)

AGENCY: Office of Planning, Research, and Evaluation; Administration for Children and Families; HHS.

ACTION: Request for public comment.

SUMMARY: The Office of Planning, Research, and Evaluation (OPRE) is proposing a data collection activity as part of the State Temporary Assistance for Needy Families (TANF) Case Studies project. This study seeks to document innovative employment and training programs for low-income individuals including TANF recipients and examine the ways the programs provide or link families to wraparound services. Over a three-year period, the study will conduct up to 12 comprehensive qualitative case studies and up to 20 profiles of innovative programs to showcase promising approaches.

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 OPREinfo@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 20291. Attn: OPRE Reports Clearance Officer. All requests, emailed or written, should be identified by the title of the information collection.

SUPPLEMENTARY INFORMATION:

Description: The State TANF Case Studies project will involve several phases including: (1) Identifying innovative programs through a scan of the field and engagement with stakeholders; (2) visiting up to 12 selected programs to collect detailed information and produce comprehensive case studies of these programs to enhance policymakers’ and other stakeholders’ understanding of promising programs helping low-income individuals to succeed in the labor force; and (3) gathering information through telephone interviews to produce up to 20 shorter case studies. The proposed information collection activities are: (1) Semi-structured interviews with program and partner administrators and frontline staff; (2) in-depth interviews with participants to better inform and enhance understanding of client experiences and perspectives; (3) a guided case review with frontline staff to capture information about client characteristics as well as intensity, frequency, duration, and sequencing of services; and (4) an observation of program services, such as case management sessions, intake and referrals, services delivered in a classroom setting, and work sites. The study will take place over a three year period.

Respondents: Respondents include program administrators, frontline program staff, and program participants. Program administrators include staff who administer and supervise the case study program under review; TANF and employment and training programs; child care and other wraparound supports; and other workforce programs and partners such as community colleges, adult basic education providers, and employers; and state decision makers, as appropriate. Frontline program staff include intake workers, case managers, job developers, and other direct service providers who work at TANF agencies and American Job Centers, employment and training providers such as community colleges, and providers of wraparound supports, such as child care subsidy frontline staff. TANF and other low-income program participants will also be respondents. All participants will be able to opt out of participating in the data collection activities.
### ANNUAL BURDEN ESTIMATES

<table>
<thead>
<tr>
<th>Instrument</th>
<th>Total number of respondents</th>
<th>Annual number of respondents</th>
<th>Number of responses per respondent</th>
<th>Average burden hours per response</th>
<th>Annual burden hours</th>
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<tbody>
<tr>
<td>Semi-structured program staff interview guide</td>
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<td>67</td>
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<td>67</td>
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<tr>
<td>In-depth participant interview guide</td>
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<td>1</td>
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<tr>
<td>Case review guide</td>
<td>24</td>
<td>8</td>
<td>2</td>
<td>.75</td>
<td>12</td>
</tr>
</tbody>
</table>

**Estimated Total Annual Burden Hours:** 91.

**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. 413, Pub. L. 115–31.

Mary B. Jones,

ACF/OPRE Certifying Officer.

[FR Doc. 2019–09658 Filed 5–9–19; 8:45 am]

**BILLING CODE 4184–09–P**

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### DEPARTMENT OF HEALTH AND HUMAN SERVICES

**Food and Drug Administration**

**[Docket No. FDA–2019–D–1798]**

**Maximal Usage Trials for Topically Applied Active Ingredients Being Considered for Inclusion in an Over-the-Counter Monograph: Study Elements and Considerations; Guidance for Industry; Availability**

**AGENCY:** Food and Drug Administration, HHS.

**ACTION:** Notice of availability.

**SUMMARY:** The Food and Drug Administration (FDA or Agency) is announcing the availability of a final guidance for industry entitled “Maximal Usage Trials for Topically Applied Active Ingredients Being Considered for Inclusion in an Over-the-Counter Monograph: Study Elements and Considerations.” This guidance represents FDA’s current thinking on the conduct of in vivo absorption trials for topically applied active ingredients that are under consideration for inclusion in an over-the-counter (OTC) monograph.

**DATES:** The announcement of the guidance is published in the Federal Register on May 10, 2019.

**ADDRESSES:** You may submit either electronic or written comments on Agency guidances at any time as follows:

**Electronic Submissions**

Submit electronic comments in the following way:

- **Federal eRulemaking Portal:** https://www.regulations.gov. Follow the instructions for submitting comments. Comments submitted electronically, including attachments, to https://www.regulations.gov will be posted to the docket unchanged. Because your comment will be made public, you are solely responsible for ensuring that your comment does not include any confidential information that you or a third party may not wish to be posted, such as medical information, your or anyone else’s Social Security number, or confidential business information, such as a manufacturing process. Please note that if you include your name, contact information, or other information that identifies you in the body of your comments, that information will be posted on https://www.regulations.gov.

- If you want to submit a comment with confidential information that you do not wish to be made available to the public, submit the comment as a written/paper submission and in the manner detailed (see “Written/Paper Submissions” and “Instructions”).

**Written/Paper Submissions**

Submit written/paper submissions as follows:

- **Mail/Hand Delivery/Courier (for written/paper submissions):** Dockets Management Staff (HFA–305), Food and Drug Administration, 5630 Fishers Lane, Rm. 1061, Rockville, MD 20852.

**Instructions:** All submissions received must include the Docket No. FDA–2019–D–1798 for “Maximal Usage Trials for Topically Applied Active Ingredients Being Considered for Inclusion in an Over-the-Counter Monograph: Study Elements and Considerations; Guidance for Industry.” Received comments will be placed in the docket and, except for those submitted as “Confidential Submissions,” publicly viewable at https://www.regulations.gov or at the Dockets Management Staff between 9 a.m. and 4 p.m., Monday through Friday.

- **Confidential Submissions**—To submit a comment with confidential information that you do not wish to be made publicly available, submit your comments only as a written/paper submission. You should submit two copies total. One copy will include the information you claim to be confidential with a heading or cover note that states “THIS DOCUMENT CONTAINS CONFIDENTIAL INFORMATION.” The Agency will review this copy, including the claimed confidential information, in its consideration of comments. The second copy, which will have the claimed confidential information redacted/blacked out, will be available for public viewing and posted on https://www.regulations.gov. Submit both copies to the Dockets Management Staff. If you do not wish your name and contact information to be made publicly available, you can provide this information on the cover sheet and not in the body of your comments and you must identify this information as “confidential.” Any information marked as “confidential” will not be disclosed except in accordance with 21 CFR 10.20 and other applicable disclosure law. For more information about FDA’s posting of comments to public dockets, see 80 FR 56469, September 18, 2015, or access the information at: https://www.gpo.gov/fdsys/pkg/FR-2015-09-18/pdf/2015-23389.pdf.

**Docket:** For access to the docket to read background documents or the electronic and written/paper comments received, go to https://www.regulations.gov and insert the docket number, found in brackets in the