running away that occurred after a child's adoption) as well as family functioning, perceptions of the adoption relationship, and services and support received after adoption. Due to the COVID–19 pandemic, initial activities to contact potential respondents were delayed. As a result, ACF is requesting an extension to collect data beyond the current OMB expiration date of September 30, 2021.

Respondents: Adopted youth, young adults, adults, and their associated adoptive parents who participated in NSCAW I or II.

ANNUAL BURDEN ESTIMATES

Instrument	No. of respondents (total over request period)	No. of responses per respondent (total over request period)	Avg. burden per response (in hours)	Total burden (in hours)	Annual burden (in hours)
Survey of NSCAW Adopted Youth, Young Adults, and Adults	588	1	.5	294	294
Survey of NSCAW Adoptive Parents	554	1	.5	277	277

Estimated Total Annual Burden Hours: 571.

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: Child Abuse Prevention and Treatment and Adoption Reform Act of 1978.

Mary B. Jones,

ACF/OPRE Certifying Officer.

[FR Doc. 2021–07420 Filed 4–9–21; 8:45 am]

BILLING CODE 4184–44–P

DEPARTMENT OF HEALTH AND HUMAN SERVICES

Administration for Children and Families

[OMB No. 0970–0323]

Proposed Information Collection Activity; Child Care Improper Payments Data Collection Instructions

AGENCY: Office of Child Care, Administration for Children and Families, HHS.

ACTION: Request for public comment.

SUMMARY: The Administration for Children and Families is proposing revisions to an approved information collection, Child Care Improper Payments Data Collection Instructions (OMB #0970–0323, expiration 10/31/2021). There are minor changes requested to the form.

DATES: *Comments due within 60 days of publication.* In compliance with the requirements of the Paperwork Reduction Act of 1995, ACF 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 (OPRE), 330 C Street SW, Washington, DC 20201, Attn: ACF Reports Clearance Officer. All requests, emailed or written, should be identified by the title of the information collection.

SUPPLEMENTARY INFORMATION:

Description: Section 2 of the Payment Integrity Information Act of 2019 (PIIA) provides for estimates and reports of improper payments by federal agencies. Subpart K of 45 CFR, Part 98 of the Child Care and Development Fund (CCDF) requires states to prepare and submit a report of errors occurring in the administration of CCDF grant funds once every 3 years.

The Office of Child Care (OCC) is completing the fifth 3-year cycle of case record reviews to meet the requirements for reporting under PIIA. The current data collection forms and instructions expire October 31, 2021. As part of the renewal process, OCC has revised the document with minor changes that do not change the methodology, but provide respondents with additional guidance, clarification, and support to facilitate completeness and accuracy of the required data submissions.

Clarifying language and a question have been added to the revised document to support Lead Agencies that administer all or part of the CCDF program through other governmental or non-governmental agencies to include the following:

- In Section 1 *Introduction* on page 2, a subsection ''Considerations for Administering CCDF Through Other Agencies'' was added to describe how Lead Agency responsibilities in administering the CCDF program through other entities apply to the error rate review process.
- In Section III *Creating the Sampling Decisions, Assurances, and Fieldwork Preparation Plan* on page 11, and the *Sampling Decisions, Assurances, and Fieldwork Preparation Plan Report template* (Attachment 1), a new item was added at Item 3g Case Review Logistics to request information about how a Lead Agency accesses documents stored by other entities if part of eligibility is determined by the other entity.

OCC is particularly interested in feedback about the clarity of these instructions and the ease and accuracy with which respondents can provide information on accessing documents stored by other entities.

Respondents: State grantees, the District of Columbia, and Puerto Rico.

ANNUAL BURDEN ESTIMATES

Instrument	Total number of respondents	Total number of responses per respondent	Average burden hours per response	Total burden hours	Annual burden hours
Sampling Decisions, Assurances, and Fieldwork Preparation Plan	52	1	106	5,512	1,837
Record Review Worksheet	52	276	6.33	90,848	30,283
State Improper Payments Report	52	1	639	33,228	11,076
Corrective Action Plan	5	2 [a]	156	1,560	520
Estimated Total Annual Burden Hours					43,716

[a] The total number of responses per respondent ranges from one to three, depending on how long it takes respondents to reduce the Improper Payment Rate to below the threshold. Respondents submit a *Corrective Action Plan* that covers a 1-year period; at the end of each year, if respondents have not reduced the Improper Payment Rate to below the threshold, they submit a new *Corrective Action Plan* for the following year. An average of two responses per respondent is used to calculate annual burden estimates.

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: 45 CFR part 98, subpart K.

Mary B. Jones,

ACF/OPRE Certifying Officer.

[FR Doc. 2021–07425 Filed 4–9–21; 8:45 am]

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DEPARTMENT OF HEALTH AND HUMAN SERVICES

Food and Drug Administration

[Docket No. FDA–2011–N–0655]

Animal Generic Drug User Fee Act; Public Meeting; Request for Comments

AGENCY: Food and Drug Administration, HHS.

ACTION: Notice of public meeting; request for comments.

SUMMARY: The Food and Drug Administration (FDA, the Agency, or we) is announcing the following virtual public meeting entitled ''Animal Generic Drug User Fee Act.'' The purpose of the public meeting is to invite public comment on the Animal Generic Drug User Fee Act (AGDUFA) program and suggestions regarding the features FDA should consider for the next reauthorization of the AGDUFA program. The meeting will be open to the public.

DATES: The public meeting will be hosted via a live virtual webcast on Thursday, May 20, 2021, from 11 a.m. to 1 p.m. Eastern Time. See the **SUPPLEMENTARY INFORMATION** section for registration dates and information. To permit the widest possible opportunity to obtain comments on all aspects of the public meeting, the docket will remain open for comment throughout the reauthorization process of AGDUFA, until December 1, 2022. In addition to being publicly viewable at *http://www.regulations.gov,* comments received by June 21, 2021, suggesting changes to the program, will also be published on *https://www.fda.gov/industry/animal-generic-drug-user-fee-act-agdufa/agdufa-meetings.*

ADDRESSES: You may submit comments as follows. Please note that late, untimely filed comments will not be considered. Electronic comments must be submitted on or before December 1, 2022. The *https://www.regulations.gov* electronic filing system will accept comments until 11:59 p.m. Eastern Time at the end of December 1, 2022. Comments received by mail/hand delivery/courier (for written/paper submissions) will be considered timely if they are postmarked or the delivery service acceptance receipt is on or before that date.

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 comments will be made public, you are solely responsible for ensuring that your comments do 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.
- For written/paper comments submitted to the Dockets Management Staff, FDA will post your comment, as well as any attachments, except for information submitted, marked and identified, as confidential, if submitted as detailed in ''Instructions.''

Instructions: All submissions received must include the Docket No. FDA–2011–N–0655 for ''Animal Generic Drug User Fee Act.'' Received comments, those filed in a timely manner, 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, 240–402–7500.

- Confidential Submissions—To submit a comment with confidential information that you do not wish to be made publicly available, submit your