statistical bases. The C series of worksheets computes the average cost per visit for HHA services. The D series of worksheets are Medicare specific and are used to determine reimbursement due to the provider or program. The F series of worksheets collect data from a provider's balance sheet and income statement. *Form Number:* CMS–1780–20 (OMB control number: 0938–0022); *Frequency:* Yearly; *Affected Public:* Private Sector—Business or other for-profits, Not-for-profit institutions; *Number of Respondents:* 10,944; *Total Annual Responses:* 10,944; *Total Annual Hours:* 2,134,080. (For policy questions regarding this collection contact LuAnn Piccione at (410) 786–5423.)

Dated: May 5, 2023.

William N. Parham, III,

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

[FR Doc. 2023–10026 Filed 5–10–23; 8:45 am]

BILLING CODE 4120–01–P

DEPARTMENT OF HEALTH AND HUMAN SERVICES

Administration for Children and Families

Proposed Information Collection Activity; Mother and Infant Home Visiting Program Evaluation: Long-Term Follow-Up, Third Grade Data Collection (Office of Management and Budget #0970–0402)

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

ACTION: Request for public comments.

SUMMARY: The Administration for Children and Families (ACF), in partnership with the Health Resources and Services Administration, both of the U.S. Department of Health and Human Services (HHS), is proposing to collect data as part of the elementary school phase of the Mother and Infant Home Visiting Program Evaluation (MIHOPE). MIHOPE is a longitudinal study of the effects of Maternal, Infant, and Early Childhood Home Visiting (MIECHV)-funded home visiting on child and family outcomes. The purpose of the MIHOPE Long-Term Follow-Up, Third grade (MIHOPE–3G) data collection, which will focus on children when they are in approximately third grade, is to assess the long-term effects of MIECHV-funded home visiting on families and children when participating children are in elementary school.

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: You can obtain copies of the proposed collection of information and submit comments by emailing *OPREinfocollection@acf.hhs.gov.* Identify all requests by the title of the information collection.

SUPPLEMENTARY INFORMATION:

Description: To date, MIHOPE has been collecting data through the time children in the study were in approximately kindergarten. The currently approved materials under this OMB number include those for the kindergarten follow-up. The kindergarten direct data collection with study participants is complete, but some administrative data are still being collected from states and local education agencies.

This request is to complete administrative data collection for the kindergarten follow-up and to conduct the following data collection activities for MIHOPE 3G: (1) child welfare records data collection from states and (2) school records data collection from states and local education agencies. Future information collection requests and related **Federal Register** notices will describe future data collection efforts for this project.

Data collected during the third-grade follow-up study is being used to estimate the effects of MIECHV-funded programs on the following seven domains: (1) maternal health, (2) child health, (3) child development and school performance, (4) child maltreatment, (5) parenting, (6) crime or domestic violence, and (7) family economic self-sufficiency.

Respondents: For MIHOPE–3G, we will seek to obtain child welfare data from 11 states and school records data from up to 48 states and local education agencies. This data will be collected for 4,105 families who are currently participating in MIHOPE.

ANNUAL BURDEN ESTIMATES

Instrument	Number of respondents (total over request period)	Number of responses per respondent (total over request period)	Average burden per response (in hours)	Total burden (in hours)	Annual burden (in hours)
Remaining Burden for Kindergarten Follow-up					
State and local education agency school records: data file submission ..	8	1	15.6	125	42
State child welfare records: data file submission	5	1	8.4	42	14
New Request for MIHOPE–3G					
State and local education agency school records: data file submission ..	48	2	33.75	3,240	1,080
State child welfare records: data file submission	11	[a] 2.1	21.43	495	165

Note: [a] The 2.1 responses is a weighted average that reflects that the study team expects to collect 2 data extracts from 10 states and 3 data extracts from 1 state.

Estimated Total Annual Burden Hours: 1,301.

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: Social Security Act Title V 511 [42 U.S.C. 711]. As amended by Section 6101 of the Consolidated Appropriations Act, 2023 (Pub. L. 117–328).

Mary B. Jones,

ACF/OPRE Certifying Officer.

[FR Doc. 2023–10095 Filed 5–10–23; 8:45 am]

BILLING CODE 4184–74–P

DEPARTMENT OF HEALTH AND HUMAN SERVICES

Food and Drug Administration

[Docket No. FDA–2023–N–1619]

Pediatric Oncology Subcommittee of the Oncologic Drugs Advisory Committee; Notice of Meeting; Establishment of a Public Docket; Request for Comments

AGENCY: Food and Drug Administration, HHS.

ACTION: Notice; establishment of a public docket; request for comments.

SUMMARY: The Food and Drug Administration (FDA) announces a forthcoming public advisory committee meeting of the Pediatric Oncology Subcommittee of the Oncologic Drugs Advisory Committee. The general function of the committee is to provide advice and recommendations to FDA on regulatory issues. The meeting will be open to the public. FDA is establishing a docket for public comment on this document.

DATES: The meeting will be held virtually on June 16, 2023, from 10 a.m. to 3:30 p.m. Eastern Time.

ADDRESSES: Please note that due to the impact of the COVID–19 pandemic, all meeting participants will be joining this advisory committee meeting via an online teleconferencing platform. Answers to commonly asked questions about FDA advisory committee meetings, including information regarding special accommodations due to a disability, may be accessed at: *https://www.fda.gov/AdvisoryCommittees/AboutAdvisoryCommittees/ucm408555.htm.*

FDA is establishing a docket for public comment on this meeting. The docket number is FDA–2023–N–1619. Please note that late, untimely filed comments will not be considered. The docket will close on June 15, 2023. The *https://www.regulations.gov* electronic filing system will accept comments until 11:59 p.m. Eastern Time at the end of June 15, 2023. Comments received by mail/hand delivery/courier (for written/paper submissions) will be considered timely if they are received on or before that date.

Comments received on or before June 2, 2023, will be provided to the subcommittee. Comments received after that date will be taken into consideration by FDA. In the event that the meeting is canceled, FDA will continue to evaluate any relevant applications or information, and consider any comments submitted to the docket, as appropriate.

You may submit comments 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.
- 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–2023–N–1619 for ''Pediatric Oncology Subcommittee of the Oncologic Drugs Advisory Committee; Notice of Meeting; Establishment of a Public Docket; Request for Comments.'' Received comments, those filed in a timely manner (see **ADDRESSES**), 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 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.'' FDA 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 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 the 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.govinfo.gov/content/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 heading of this document, into the ''Search'' box and follow the prompts