minimize the information collection burden. See **DATES** and **ADDRESSES** for instructions for submitting comments.

While we will review all comments received, we may choose not to post off-topic or inappropriate comments. Otherwise, all comments will be posted without edit under the applicable docket number, including any personal information that the commenter provides. Our response to such comments will be posted at reginfo.gov under the applicable OMB control number.

Medicaid and CHIP Program (MACPro)

At this time, MACPro is made up of the main umbrella (see collection number 1 in the following list) and nine individual generic collections of information (see collection numbers 2 through 10 in the following list). Details such as the collection's requirements and burden estimates can be found in the collection's supporting statement and associated materials (see **ADDRESSES** for instructions for obtaining such documents).

Docket Information

1. *Title:* Medicaid and CHIP Program (MACPro).
Type of Request: Revision of a currently approved collection.
CMS ID Number: CMS–10434.
OMB Control Number: 0938–1188.
eRulemaking Docket ID Number: CMS–2023–0080.
Docket Web Address: https://www.regulations.gov/docket/CMS-2023-0080.
For Policy Related Questions, Contact: William N. Parham at 410–786–4669.

2. *Title:* Initial Application.
Type of Request: Extension of a currently approved collection.
CMS ID Number: CMS–10434 #1.
OMB Control Number: 0938–1188.
eRulemaking Docket ID Number: CMS–2023–0081.
Docket Web Address: https://www.regulations.gov/docket/CMS-2023-0081.
For Policy Related Questions, Contact: Stephanie Bell at 410–786–0617.

3. *Title:* CHIP State Plan Eligibility.
Type of Request: Extension of a currently approved collection.
CMS ID Number: CMS–10434 #2.
OMB Control Number: 0938–1188.
eRulemaking Docket ID Number: CMS–2023–0082.
Docket Web Address: https://www.regulations.gov/docket/CMS-2023-0082.
For Policy Related Questions, Contact: Stephanie Bell at 410–786–0617.

4. *Title:* Alternative Benefit Plans (ABPs).
Type of Request: Extension of a currently approved collection.
CMS ID Number: CMS–10434 #3.
OMB Control Number: 0938–1188.
eRulemaking Docket ID Number: CMS–2023–0083.
Docket Web Address: https://www.regulations.gov/docket/CMS-2023-0083.
For Policy Related Questions, Contact: Adrienne Delozier at 410–786–0278.

5. *Title:* Medicaid State Plan Eligibility.
Type of Request: Extension of a currently approved collection.
CMS ID Number: CMS–10434 #15.
OMB Control Number: 0938–1188.
eRulemaking Docket ID Number: CMS–2023–0090.
Docket Web Address: https://www.regulations.gov/docket/CMS-2023-0090.
For Policy Related Questions, Contact: Suzette Seng at 410–786–4703.

6. *Title:* Health Home State Plan Amendment (SPA).
Type of Request: Extension of a currently approved collection.
CMS ID Number: CMS–10434 #22.
OMB Control Number: 0938–1188.
eRulemaking Docket ID Number: CMS–2023–0084.
Docket Web Address: https://www.regulations.gov/docket/CMS-2023-0084.
For Policy Related Questions, Contact: Mary Pat Farkas at 410–786–5731.

7. *Title:* Medicaid Adult and Child Core Set Measures.
Type of Request: Extension of a currently approved collection.
CMS ID Number: CMS–10434 #26.
OMB Control Number: 0938–1188.
eRulemaking Docket ID Number: CMS–2023–0085.
Docket Web Address: https://www.regulations.gov/docket/CMS-2023-0085.
For Policy Related Questions, Contact: Virginia (Gigi) Raney at 410–786–6117.

8. *Title:* Maternal and Infant Health Quality.
Type of Request: Extension of a currently approved collection.
CMS ID Number: CMS–10434 #45.
OMB Control Number: 0938–1188.
eRulemaking Docket ID Number: CMS–2023–0086.
Docket Web Address: https://www.regulations.gov/docket/CMS-2023-0086.
For Policy Related Questions, Contact: Virginia (Gigi) Raney at 410–786–6117.

9. *Title:* Health Home Core Sets.
Type of Request: Extension of a currently approved collection.
CMS ID Number: CMS–10434 #47.
OMB Control Number: 0938–1188.
eRulemaking Docket ID Number: CMS–2023–0087.
Docket Web Address: https://www.regulations.gov/docket/CMS-2023-0087.
For Policy Related Questions, Contact: Mary Pat Farkas at 410–786–5731.

10. *Title:* Medicaid Extended Postpartum Coverage and Continuous Eligibility for Children.
Type of Request: Extension of a currently approved collection.
CMS ID Number: CMS–10434 #77.
OMB Control Number: 0938–1188.
eRulemaking Docket ID Number: CMS–2023–0088.
Docket Web Address: https://www.regulations.gov/docket/CMS-2023-0088.
For Policy Related Questions, Contact: Alexa Turner at 410–786–8823.

William N. Parham, III,

Director, Division of Information Collections and Regulatory Impacts, Office of Strategic Operations and Regulatory Affairs.

[FR Doc. 2024–02243 Filed 2–2–24; 8:45 am]

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DEPARTMENT OF HEALTH AND HUMAN SERVICES

Administration for Children and Families

Submission for Office of Management and Budget (OMB) Review; Advance Planning Document (APD) Process (OMB #0970–0417)

AGENCY: Office of Child Support Services; Administration for Children and Families, Department of Health and Human Services.

ACTION: Request for public comments.

SUMMARY: The Administration for Children and Families' (ACF) Office of Child Support Services (OCSS) requests a 3-year extension for the Advance Planning Document (APD) process (OMB #0970–0417). No changes are proposed.

DATES: *Comments due within 30 days of publication.* OMB must make a decision about the collection of information between 30 and 60 days after publication of this document in the **Federal Register**. Therefore, a comment is best assured of having its full effect if OMB receives it within 30 days of publication.

ADDRESSES: Written comments and recommendations for the proposed information collection should be sent within 30 days of publication of this notice to *www.reginfo.gov/public/do/PRAMain.* Find this particular information collection by selecting ''Currently under 30-day Review—Open

for Public Comments'' or by using the search function. You can also obtain copies of the proposed collection of information by emailing *infocollection@acf.hhs.gov.* Identify all emailed requests by the title of the information collection.

SUPPLEMENTARY INFORMATION:

Description: State child support agencies are required to establish and operate a federally approved statewide Automated Data Processing (ADP) and information retrieval system to assist in child support services. The APD process, established at 45 CFR part 95, subpart F, is the procedure by which states request and obtain approval for Federal Financial Participation (FFP) in their cost of acquiring ADP equipment and services. The ACF OCSS Division of State and Tribal Systems (DSTS) oversees this process.

States are required to submit an initial APD, containing information to assist the Secretary of the Department of Health and Human Services (HHS) in determining if the state computerized support enforcement project planning and implementation meets federal certification requirements for approving FFP. States are then required to submit annual APD updates to HHS to report project status and request ongoing FFP for systems development, enhancements, operations, and maintenance. As-needed APDs are also submitted to acquire FFP when major milestones are missed or significant changes to project schedules occur. Based on an assessment of the information provided in the APD, states that do not meet the federal requirements necessary for approval are required to conduct periodic independent verification and validation services for high-risk project oversight.

In addition to the APDs providing HHS/ACF/OCSS with the information necessary to determine the allowable level of federal funding for state systems projects, states also submit associated procurement and data security documents, such as requests for proposals (RFPs), contracts, contract amendments, and the biennial security review reports.

Respondents: State child support agencies.

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
RFP and Contract	50	4.5	4	900	300
Emergency Funding Request	21	1	2	42	14
Biennial Reports	54	1.5	1.5	121.5	40.5
Advance Planning Document	44	3.6	120	19,008	6,336
Operational Advance Planning Document	10	3	30	900	300
Independent Verification and Validation (ongoing)	3	12	10	360	120
Independent Verification and Validation (semiannually)	4	6	16	384	128
Independent Verification and Validation (quarterly)	10	12	30	3,600	1,200
System Certification	3	3	240	2,160	720

Estimated Total Annual Burden Hours: 9,158.50.

Authority: 45 CFR part 95, subpart F.

Mary B. Jones,

ACF/OPRE Certifying Officer.

[FR Doc. 2024–02166 Filed 2–2–24; 8:45 am]

BILLING CODE 4184–41–P

DEPARTMENT OF HEALTH AND HUMAN SERVICES

Food and Drug Administration

[Docket No. FDA–2024–N–0018]

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 Oncologic Drugs Advisory Committee (the 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 on March 15, 2024, from 8:30 a.m. to 5:30 p.m. eastern time.

ADDRESSES: All meeting participants will be heard, viewed, captioned, and recorded for this advisory committee meeting via an online teleconferencing and/or video conferencing platform.

Answers to commonly asked questions about FDA advisory committee meetings 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–2024–N–0018. The docket will close on March 14, 2024. Please note that late, untimely filed comments will not be considered. The *https://www.regulations.gov* electronic filing system will accept comments until 11:59 p.m. eastern time at the end of March 14, 2024. 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 March 1, 2024, will be provided to the Committee. Comments received after that date will be taken into consideration by FDA. In the event that the meeting is cancelled, 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