or OMB control number. To be assured consideration, comments and recommendations must be received by the OMB desk officer via one of the following transmissions: OMB, Office of Information and Regulatory Affairs, Attention: CMS Desk Officer, Fax Number: (202) 395–5806 *OR* Email: *OIRA_submission@omb.eop.gov.*

To obtain copies of a supporting statement and any related forms for the proposed collection(s) summarized in this notice, you may make your request using one of following:

1. Access CMS' Web site address at Web site address at *https://www.cms.gov/Regulations-and-Guidance/Legislation/PaperworkReductionActof1995/PRA-Listing.html.*

2. Email your request, including your address, phone number, OMB number, and CMS document identifier, to *Paperwork@cms.hhs.gov.*

3. Call the Reports Clearance Office at (410) 786–1326.

FOR FURTHER INFORMATION CONTACT: William Parham at (410) 786–4669.

SUPPLEMENTARY INFORMATION: Under the Paperwork Reduction Act of 1995 (PRA) (44 U.S.C. 3501–3520), federal agencies must obtain approval from the Office of Management and Budget (OMB) for each collection of information they conduct or sponsor. The term ''collection of information'' is defined in 44 U.S.C. 3502(3) and 5 CFR 1320.3(c) and includes agency requests or requirements that members of the public submit reports, keep records, or provide information to a third party. Section 3506(c)(2)(A) of the PRA (44 U.S.C. 3506(c)(2)(A)) requires federal agencies to publish a 30-day notice in the **Federal Register** concerning each proposed collection of information, including each proposed extension or reinstatement of an existing collection of information, before submitting the collection to OMB for approval. To comply with this requirement, CMS is publishing this notice that summarizes the following proposed collection(s) of information for public comment:

1. *Type of Information Collection Request:* Extension without change of a currently approved collection; *Title of Information Collection:* Quarterly Medicaid and CHIP Budget and Expenditure Reporting for the Medical Assistance Program, Administration and CHIP; *Use:* The Medicaid Budget and Expenditure System (MBES) and the Child Health Budget and Expenditure System (CBES) is a financial reporting system that produces Budget and expenditures for Medical Assistance and Children's Health Insurance Program. All forms are to be filed on a quarterly basis and need to be certified by the States to the CMS. The forms consist of CMS–21 and –21B, CMS–37, and CMS–64.

Forms CMS–21 and –21B provide CMS with the information necessary to issue quarterly grant awards, monitor current year expenditure levels, determine the allowability of state claims for reimbursement, develop Children's Health Insurance Program (CHIP) financial management information, provide for state reporting of waiver expenditures, and ensure that the federally established allotment is not exceeded. They are also necessary in the redistribution and reallocation of unspent funds over the federally mandated timeframes.

Form CMS–37 due dates are November 15, February 15, May 15 and August 15 of each fiscal year. While all submissions represent equally important components of the grant award cycle, the May and November submissions are particularly significant for budget formulation. The November submission introduces a new fiscal year to the budget cycle and serves as the basis for the formulation of the Medicaid portion of the President's Budget, which is presented to Congress in January. The February and August submissions are used primarily for budget execution in providing interim updates to our Office of Financial Management, the Department of Health and Human Services, the Office of Management and Budget, and Congress depending on the scheduling of the national budget review process in a given fiscal year. The submissions provide us with base information necessary to track current year obligations and expenditures in relation to the current year appropriation and to notify senior managers of any impending surpluses or deficits.

Form CMS–64 is used to issue quarterly grant awards, monitor current year expenditure levels, determine allowed state claims for reimbursement, develop Medicaid financial management information provide for state reporting of waiver expenditures, ensure that the federally-established limit is not exceeded for HCBS waivers, and to allow for the implementation of the Assignment of Rights and Part A and Part B Premium (*i.e.,* accounting for overdue Part A and Part B Premiums under state buy-in agreements)—Billing Offsets. *Form Number:* CMS–10529 (OMB control number: 0938–1265); *Frequency:* Quarterly; *Affected Public:* State, Local, or Tribal Governments; *Number of Respondents:* 56; *Total Annual Responses:* 672; *Total Annual Hours:* 17,920. (For policy questions regarding this collection contact Chris Kessler at 410–786–7168.)

Dated: December 6, 2017.

William N. Parham, III,

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

[FR Doc. 2017–26575 Filed 12–8–17; 8:45 am]

BILLING CODE 4120–01–P

DEPARTMENT OF HEALTH AND HUMAN SERVICES

Administration for Children and Families

Submission for OMB Review; Comment Request

Title: Case Plan Requirement, Title IV–E of the Social Security Act.

OMB No.: 0970–0428.

Description: Under section 471(a)(16) of title IV–E of the Social Security Act (the Act), to be eligible for payments, states and tribes must have an approved title IV–E plan that provides for the development of a case plan for each child for whom the State or Tribe receives foster care maintenance payments and that provides a case review system that meets the requirements in section 475(5) and 475(6) of the Act.

The case review system assures that each child has a case plan designed to achieve placement in a safe setting that is the least restrictive (most family-like) setting available and in close proximity to the child's parental home, consistent with the best interest and special needs of the child. Through these requirements, States and Tribes also comply, in part, with title IV–B section 422(b) of the Act, which assures certain protections for children in foster care.

The case plan is a written document that provides a narrative description of the child-specific program of care. Federal regulations at 45 CFR 1356.21(g) and section 475(1) of the Act delineate the specific information that should be addressed in the case plan. The Administration for Children and Families (ACF) does not specify a recordkeeping format for the case plan nor does ACF require submission of the document to the Federal government. Case plan information is recorded in a format developed and maintained by the State or Tribal child welfare agency.

Respondents: State and Tribe title IV–B and title IV–E agencies.

ANNUAL BURDEN ESTIMATES

Instrument	Number of respondents	Number of responses per respondent	Average burden hours per response	Total burden hours
Case Plan	544,098	1	4.80	2,626,436

Estimated Total Annual Burden Hours: 2,626,436.

Additional Information: Copies of the proposed collection may be obtained by writing to the Administration for Children and Families, Office of Planning, Research and Evaluation, 330 C Street SW., Washington, DC 20201. Attention Reports Clearance Officer. All requests should be identified by the title of the information collection. Email address: *infocollection@acf.hhs.gov.*

OMB Comment: OMB is required to make a decision concerning 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. Written comments and recommendations for the proposed information collection should be sent directly to the following: Office of Management and Budget, Paperwork Reduction Project, Email: *OIRA_SUBMISSION@OMB.EOP.GOV.* Attn: Desk Officer for the Administration for Children and Families.

Robert Sargis,

Reports Clearance Officer.

[FR Doc. 2017–26553 Filed 12–8–17; 8:45 am]

BILLING CODE 4184–25–P

DEPARTMENT OF HEALTH AND HUMAN SERVICES

Food and Drug Administration

[Docket No. FDA–2014–N–1072]

Agency Information Collection Activities; Submission for Office of Management and Budget Review; Comment Request; Application for Participation in Food and Drug Administration Fellowship Programs

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.

DATES: Fax written comments on the collection of information by January 10, 2018.

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 0910–0780. Also include the FDA docket number found in brackets in the heading of this document.

FOR FURTHER INFORMATION CONTACT: Amber Sanford, Office of Operations, Food and Drug Administration, Three White Flint North, 10A–12M, 11601 Landsdown St., North Bethesda, MD 20852, 301–796–8867, *PRAStaff@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.

Application for Participation in FDA Fellowship Programs (Formerly Application for Participation in the FDA Commissioner's Fellowship Program)—OMB Control Number 0910–0780—Extension

Sections 1104, 1302, 3301, 3304, 3320, 3361, 3393, and 3394 of Title 5 of the United States Code authorize Federal agencies to rate applicants for Federal jobs. The proposed information collection involves brief online applications completed by applicants applying to FDA's fellowship programs. These voluntary online applications will allow the Agency to easily and efficiently elicit and review information from students and healthcare professionals who are interested in becoming involved in FDA-wide activities. The process will reduce the time and cost of submitting written documentation to the Agency and lessen the likelihood of applications being misrouted within the Agency mail system. It will assist the Agency in promoting and protecting the public health by encouraging outside persons to share their expertise with FDA.

In the **Federal Register** of June 20, 2017 (82 FR 28075), FDA published a 60-day notice requesting public comment on the proposed collection of information. Although one comment was received, it wasn't responsive to the four collection of information topics solicited and therefore will not be discussed in this document.

FDA estimates the burden of this collection of information as follows:

TABLE 1—ESTIMATED ANNUAL RECORDKEEPING BURDEN [1]

Activity	Number of respondents	Number of responses per respondent	Total annual responses	Average burden per response	Total hours
Commissioner's Fellowship Program	600	1	600	1.33	798
Regulatory Science Internship Program	250	1	250	1	250
Medical Device Fellowship Program	250	1	250	1	250
Total					1,298

[1] There are no capital costs or operating and maintenance costs associated with this collection of information.