12. UAC Case Status: This instrument is used by care providers to monitor the status of high-level milestones in a UAC's case.

Respondents: ORR grantee and contractor staff; UAC; sponsors; and child advocates.

ANNUAL BURDEN ESTIMATES

Instrument	Annual total number of respondents	Annual total number of responses per respondent	Average burden minutes per response	Annual total burden hours
Sponsor Assessment (Form S–5)	206	327	60	67,362
Home Study Report (Form S–6)	11	364	45	3,003
New Sponsor (Form S–7)	206	327	20	22,454
Initial Intakes Assessment (Form S–8)	206	363	15	18,695
Assessment for Risk (Form S–9)	206	794	30	81,782
UAC Assessment (Form S–11)	206	369	45	57,011
UAC Case Review (Form S–12)	206	764	30	78,692
Individual Service Plan (Form S–13)	206	985	15	50,728
UAC Long Term Foster Care Travel Request (Form S–14)	30	9	15	68
Child Advocate Referral and Appointment (Form S–15)	206	5	15	258
Summary Notes Thirty Day Restrictive Placement Case Review (Form S–16)	15	68	30	510
UAC Case Status	206	354	3	3,646
Estimated Annual Burden Total:				384,207

Authority: 6 U.S.C. 279; 8 U.S.C. 1232; Flores v. Reno Settlement Agreement, No. CV85–4544–RJK (C.D. Cal. 1996)

Mary B. Jones,

ACF/OPRE Certifying Officer.

[FR Doc. 2020–27898 Filed 12–17–20; 8:45 am]

BILLING CODE 4184–45–P

DEPARTMENT OF HEALTH AND HUMAN SERVICES

Administration for Children and Families

Submission for OMB Review; Advance Planning Document (APD) Process (OMB #0970–0417)

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

ACTION: Request for public comment.

SUMMARY: The Administration for Children and Families' (ACF) Office of Child Support Enforcement (OCSE) is requesting an update to the Advance Planning Document (APD) process (OMB #0970–0417). OCSE proposes revisions to the annual burden estimates to reflect an increase in the number of states seeking approval to implement modernization solutions in efforts to replace antiquated legacy child support enforcement systems and to address an excess demand for emergency funding requests due to the impacts of the COVID–19 pandemic.

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.

SUPPLEMENTARY INFORMATION:

Description: 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 Automated Data Processing (ADP) equipment and services.

State child support agencies are required to establish and operate a federally approved statewide ADP and information retrieval system to assist in child support enforcement. States are required to submit an initial APD containing information to assist the Secretary of the U.S. Department of Health and Human Services (HHS) in determining if the state computerized support enforcement project planning and implementation meets federal certification requirements needed for the approval of 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 milestone 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 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 the request 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. 2020–27916 Filed 12–17–20; 8:45 am]

BILLING CODE 4184–41–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 and Traineeship Programs

AGENCY: Food and Drug Administration, HHS.

ACTION: Notice.

SUMMARY: The Food and Drug Administration (FDA or Agency) 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: Submit written comments (including recommendations) on the collection of information by January 19, 2021.

ADDRESSES: To ensure that comments on the information collection are received, OMB recommends that written comments be submitted to *https://www.reginfo.gov/public/do/PRAMain.* Find this particular information collection by selecting ''Currently under Review—Open for Public Comments'' or by using the search function. The OMB control number for this information collection is 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 and Traineeship Programs

OMB Control Number 0910–0780—Revision

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 and Traineeship 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 October 19, 2018 (83 FR 53065), FDA published a 60-day notice requesting public comment on the proposed collection of information. Although one comment was received, it was not 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
Medical Device Fellowship Program	250	1	250	1	250
FDA Traineeship Program	1,000	1	1,000	1	1,000
Reagan-Udall Fellowship at FDA	50	1	50	1	50
Total					1,300

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