## GENERAL SERVICES ADMINISTRATION

**[OMB Control No. 3090–0290; Docket No. 2019–0001; Sequence No. 15]**

### Submission for OMB Review; System for Award Management Registration Requirements for Financial Assistance Recipients

**AGENCY:** Office of Systems Management, General Services Administration (GSA).

**ACTION:** Notice.

**SUMMARY:** Under the provisions of the Paperwork Reduction Act of 1995, the Regulatory Secretariat Division has submitted to the Office of Management and Budget (OMB) a request to review and approve a renewal of a previously approved information collection requirement regarding the pre-award registration requirements for Prime Grant Recipients. The updated information collection title is based on the Office of Management and Budget's (OMB) proposed expansion of SAM registration requirements to include all entities that receive financial assistance.

**DATES:** Submit comments on or before September 24, 2020.

**ADDRESSES:** Written comments and recommendations for this 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 Review—Open for Public Comments'' or by using the search function.

**FOR FURTHER INFORMATION CONTACT:** Ms. Nancy Goode, Program Manager, IAE Outreach and Stakeholder Engagement Division, at telephone number 703–605–2175; or via email at *nancy.goode@gsa.gov.*

**SUPPLEMENTARY INFORMATION:**

#### A. Purpose

This information collection requires information necessary for prime applicants and recipients, excepting individuals, of Federal financial assistance to register in the System for Award Management (SAM) and maintain an active SAM registration with current information at all times during which they have an active Federal award or an application or plan under consideration by an agency pursuant to 2CFR Subtitle A, Chapter I, and Part 25 (75 FR 5672). This facilitates prime awardee reporting of sub-award and executive compensation data pursuant to the Federal Funding Accountability and Transparency Act (Pub. L. 109–282, as amended by section 6202(a) of Pub. L. 110–252). This information collection requires that all prime financial assistance awardees, subject to reporting under the Transparency Act register and maintain their registration in SAM.

This information collection is being amended to meet a statutory requirement of the National Defense Authorization Act (NDAA) of FY 2013. The NDAA of 2013 requires that the Federal Awardee Performance and Integrity Information System (FAPIIS)(currently located in SAM) include information on a non-Federal entity's parent, subsidiary, or successor entities. Applicants will need to provide information in SAM on their immediate and highest level owner as well as predecessors that have been awarded a Federal contract, grant, or cooperative agreement within the last three years. Additionally, the information collection is being amended to increase transparency regarding Federal spending and to support implementation of the Digital Accountability and Transparency Act of 2014 (DATA ACT).

OMB proposes to expand the requirement to register in SAM beyond grants, cooperative agreements, and contracts, to entities that receive financial assistance such as loans, insurance, and direct appropriations. This information collection requirement (published in the **Federal Register** at 85 FR 49506 on August 13, 2020) is included in OMB's proposed revision to guidance in 2 CFR Subtitle A, Chapter I, and Parts 25, 170, and 200.

#### B. Annual Reporting Burden

*Respondents:* 172,084.
*Responses per Respondent:* 1.
*Total Annual Responses:* 172,084.
*Hours per Response:* 2.5.
*Total Burden Hours:* 430,210.

#### C. Public Comments

A 60-day notice published in the **Federal Register** at 85 FR 3690 on January 22, 2020. No comments were received.

*Obtaining Copies of Proposals:* Requesters may obtain a copy of the information collection documents from the General Services Administration, Regulatory Secretariat Division (MVCB) at *GSARegSec@gsa.gov.* Please cite OMB Control No. 3090–0290, System for Award Management Registration Requirements for Financial Assistance Recipients, in all correspondence.

**Beth Anne Killoran,**

*Deputy Chief Information Officer.*

[FR Doc. 2020–18618 Filed 8–24–20; 8:45 am]

**BILLING CODE 6820–WY–P**

## DEPARTMENT OF HEALTH AND HUMAN SERVICES

### Administration for Children and Families

**[OMB #0970–0428]**

### Submission for OMB Review; Case Plan Requirement, Title IV–E of the Social Security Act

**AGENCY:** Administration on Children, Youth and Families, Administration for Children and Families, HHS.

**ACTION:** Request for public comment.

**SUMMARY:** The Administration on Children, Youth and Families (ACYF), Administration for Children and Families (ACF), U.S. Department of Health and Human Services (HHS), is requesting a 3-year extension of the information collection—Case Plan Requirement, Title IV–E of the Social Security Act, (OMB #0970–0428, expiration 3/31/2021). ACF is reporting a change to the information collection—the burden estimates in the previously-approved request were based on the children in foster care as the respondent instead of the agency completing the case plan. The burden estimates, therefore, are adjusted accordingly.

**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 case plan information collection is authorized in sections 422(b)(8)(A)(ii) and 471(a)(16), and defined in sections 475 and 475A of the Social Security Act (the Act). Statutory requirements in the Act mandate that states, territories, and tribes with an approved title IV–E plan develop a case plan and case review system for each child in the foster care system for whom the state, territory, or tribe receives title IV–E reimbursement of foster care maintenance payments.

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. States, territories, and tribes meeting these requirements also partly comply 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 sections 475 and 475A of the Act delineate the specific information that must be addressed in the case plan. ACF does not specify a 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, territorial, or tribal title IV–E agency.

*Respondents:* State, territorial, and tribal title IV–E 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 |
|---|---|---|---|---|---|
| Case Plan | 64 | 26,427 | 4.8 | 8,118,374 | 2,706,125 |

*Estimated Total Annual Burden Hours:* 2,706,125.

**Authority:** 42 U.S.C. 622; 42 U.S.C. 671; 42 U.S.C. 675.

**John M. Sweet Jr.,**

*ACF/OPRE Certifying Officer.*

[FR Doc. 2020–18652 Filed 8–24–20; 8:45 am]

**BILLING CODE 4184–01–P**

## DEPARTMENT OF HEALTH AND HUMAN SERVICES

### Food and Drug Administration

**[Docket No. FDA–2020–N–0908]**

### Agency Information Collection Activities; Submission for Office of Management and Budget Review; Comment Request; Submission of Petitions—Food Additive, Color Additive (Including Labeling), Submission of Information to a Master File in Support of Petitions; and Electronic Submission Using Food and Drug Administration Form 3503

**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:** Submit written comments (including recommendations) on the collection of information by September 24, 2020.

**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 0910–0016. Also include the FDA docket number found in brackets in the heading of this document.

**FOR FURTHER INFORMATION CONTACT:** Domini Bean, Office of Operations, Food and Drug Administration, Three White Flint North, 10A–12M, 11601 Landsdown St., North Bethesda, MD 20852, 301–796–5733, *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.

### Submission of Petitions: Food Additive, Color Additive (Including Labeling); Submission of Information to a Master File in Support of Petitions; Electronic Submission Using Form FDA 3503—21 CFR 70.25, 71.1, 171.1, 172, 173, 179, and 180

*OMB Control Number 0910–0016—Revision*

Section 409(a) of the Federal Food, Drug, and Cosmetic Act (FD&C Act) (21 U.S.C. 348(a)) provides that a food additive shall be deemed to be unsafe, unless: (1) The additive and its use, or intended use, are in conformity with a regulation issued under section 409 that describes the condition(s) under which the additive may be safely used; (2) the additive and its use, or intended use, conform to the terms of an exemption for investigational use; or (3) a food contact notification submitted under section 409(h) of the FD&C Act is effective. Food additive petitions (FAPs) are submitted by individuals or companies to obtain approval of a new food additive or to amend the conditions of use permitted under an existing food additive regulation. Section 171.1 of FDA's regulations (21 CFR 171.1) specifies the information that a petitioner must submit to establish that the proposed use of a food additive is safe and to secure the publication of a food additive regulation describing the conditions under which the additive may be safely used. Parts 172, 173, 179, and 180 (21 CFR parts 172, 173, 179, and 180) contain labeling requirements for certain food additives to ensure their safe use.

Section 721(a) of the FD&C Act (21 U.S.C. 379e(a)) provides that a color additive shall be deemed to be unsafe unless the additive and its use are in conformity with a regulation that describes the condition(s) under which the additive may safely be used, or the additive and its use conform to the terms of an exemption for investigational use issued under section 721(f) of the FD&C Act. Color additive petitions (CAPs) are submitted by individuals or companies to obtain approval of a new color additive or a change in the conditions of use permitted for a color additive that is already approved. Section 71.1 of the Agency's regulations (21 CFR 71.1) specifies the information that a petitioner must submit to establish the safety of a color additive and to secure the issuance of a regulation permitting its use. FDA's color additive labeling requirements in § 70.25 (21 CFR 70.25) require that color additives that are to be used in food, drugs, devices, or cosmetics be labeled with sufficient information to ensure their safe use.

FDA scientific personnel reviews FAPs to ensure the safety of the intended use of the additive in or on food, or that may be present in food as a result of its use in articles that contact food. Likewise, FDA personnel review CAPs to ensure the safety of the color