ADDRESSES: When commenting, please reference the document identifier or OMB control number. To be assured consideration, comments and recommendations must be submitted in any one of the following ways:

1. *Electronically.* You may send your comments electronically to *https://www.regulations.gov.* Follow the instructions for ''Comment or Submission'' or ''More Search Options'' to find the information collection document(s) that are accepting comments.

2. *By regular mail.* You may mail written comments to the following address: CMS, Office of Strategic Operations and Regulatory Affairs, Division of Regulations Development.

Attention: Document Identifier/OMB Control Number: ____, Room C4–26–05, 7500 Security Boulevard, Baltimore, Maryland 21244–1850.

To obtain copies of a supporting statement and any related forms for the proposed collection(s) summarized in this notice, please access the CMS PRA website by copying and pasting the following web address into your web browser: *https://www.cms.gov/Regulations-and-Guidance/Legislation/PaperworkReductionActof1995/PRA-Listing.*

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

SUPPLEMENTARY INFORMATION:

Contents

This notice sets out a summary of the use and burden associated with the following information collections. More detailed information can be found in each collection's supporting statement and associated materials (see **ADDRESSES**).

CMS–10110 Manufacturer Submission of Average Sales Price (ASP) Data for Medicare Part B Drugs and Biologicals

CMS–10242 Emergency Ambulance Transports and Beneficiary Signature

Under the 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 requires federal agencies to publish a 60-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.

Information Collection

1. *Type of Information Collection Request:* Revision of a currently approved collection; *Title of Information Collection:* Manufacturer Submission of Average Sales Price (ASP) Data for Medicare Part B Drugs and Biologicals; *Use:* Section 401 of Division CC of Title IV of the Consolidated Appropriations Act (CAA), 2021 amended section 1847A of the Social Security Act (the Act) to add new section 1847A(f)(2) of the Act, which requires manufacturers without a Medicaid drug rebate agreement to report average sales price (ASP) information to CMS for calendar quarters beginning on January 1, 2022, for drugs or biologicals payable under Medicare Part B and described in sections 1842(o)(1)(C), (E), or (G) or 1881(b)(14)(B) of the Act, including items, services, supplies, and products that are payable under Part B as a drug or biological. The reported ASP data are used to establish the Medicare payment amounts. *Form Number:* CMS–10110 (OMB control number: 0938–0921); *Frequency:* Quarterly; *Affected Public:* Private sector, Business or other for-profit; *Number of Respondents:* 500; *Total Annual Responses:* 2,000; *Total Annual Hours:* 26,000. (For policy questions regarding this collection contact Felicia Brown at 410–786–9287.)

2. *Type of Information Collection Request:* Extension of a currently approved collection; *Title of Information Collection:* CAHPS Hospice Survey; *Use:* CMS is required to collect and publicly report information on the quality of services provided by hospices under provisions in the Social Security Act. Specifically, sections 1814(i)(5)(A) through (C) of the Act, as added by section 3132(a) of the Patient Protection and Affordable Care Act (PPACA) (Pub. L. 111–148), required hospices to begin submitting quality data, based on measures specified by the Secretary of the Department of Health and Human Services (the Secretary) for FY 2014 and subsequent FYs.

The goal of the survey is to measure the experiences of patients and their caregivers with hospice care. The survey was developed to:

- Provide a source of information from which selected measures could be publicly reported to beneficiaries and their family members as a decision aid for selection of a hospice program;
- Aid hospices with their internal quality improvement efforts and external benchmarking with other facilities;
- Provide CMS with information for monitoring the care provided.

Form Number: CMS–10537 (OMB control number: 0938–1257); *Frequency:* Once; *Affected Public:* Individuals and Households; *Number of Respondents:* 1,140,695; *Total Annual Responses:* 1,140,695; *Total Annual Hours:* 198,481. (For policy questions regarding this collection contact Lauren Fuentes at 410–786 2290 or 443–618–2123.)

Dated: January 25, 2023.

William N. Parham, III,

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

[FR Doc. 2023–01822 Filed 1–27–23; 8:45 am]

BILLING CODE 4120–01–P

DEPARTMENT OF HEALTH AND HUMAN SERVICES

Administration for Children and Families

[0970–0490]

Submission for OMB Review; Generic Program-Specific Performance Progress Report

AGENCY: Office of Planning, Research, and Evaluation, Administration for Children and Families, HHS.

ACTION: Request for public comments.

SUMMARY: This notice describes the proposal to extend data collection under the Administration for Children and Families (ACF) Generic Program-Specific Performance Progress Report (PPR) (0970–0490). This overarching generic currently allows ACF program offices to collect performance and progress data from recipients and sub-recipients who receive funding from ACF under a discretionary grant or cooperative agreement. This generic mechanism provides the opportunity for ACF program offices to tailor requests for performance and progress data to specific funding recipients. ACF is proposing to include performance and progress data reporting for mandatory funding recipients in addition to discretionary funding recipients. ACF is also requesting an increase in burden.

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: ACF is primarily a grant-making agency that promotes the economic and social well-being of families, children, individuals and communities with partnerships, funding, guidance, training and technical assistance.

Prior to the use of this generic program-specific PPR, a standard ACF PPR (#0970–0406) was used for all ACF discretionary grant and cooperative agreement awards for post award reporting. Historically, on the standard ACF PPR form, ACF required grantees to only respond to a common set of broad questions, which often solicited qualitative or incomplete information. This one-size-fits-all approach did not adequately collect the specific data needed for particular grant programs or allow program offices to assess continuous quality improvement. Different grant programs vary in purpose, target population, and activities. Therefore, a need for program offices to customize performance measurements was identified and the generic program-specific PPR was developed. Non-discretionary funding recipients have historically provided performance and progress data through program-specific information collection requests. When subject to the Paperwork Reduction Act, these collections have been approved through full information collection requests.

ACF program offices have provided feedback that the ability to efficiently customize performance measurements would also be helpful for these funding recipients and therefore, ACF would like to expand this generic to cover these non-discretionary funding recipients as well.

ACF program offices have benefited from the ability to create and use a program-specific PPR that is more effective and includes specific data elements that reflects a specific program's indicators, demographics, priorities and objectives.

A generic program-specific PPR that can be tailored for program-specific needs allows program offices to collect useful data in a uniform and systematic manner. The reporting format allows program offices to gather uniform program performance data from each grantee, allowing aggregation at the program level to calculate outputs and outcomes, providing a snapshot and allowing for longitudinal analysis.

Data from a tailored program-specific PPR that demonstrates a program's successes and challenges have been useful for accountability purposes, such as required reports to Congress. Moreover, it has been useful for program management and oversight, such as identifying grantees' technical assistance needs and ensuring compliance with federal and programmatic regulations and policies. To review currently approved PPRs under this generic, see: *https://www.reginfo.gov/public/do/PRAICList?ref_nbr=202206-0970-004.*

Respondents: ACF funding recipients.

Annual Burden Estimates: ACF is requesting an increase in burden account for the potential use by non-discretionary programs and to reflect use over the past 3 years and anticipated use in the next 3 years.

Instrument	Total number of respondents	Total number of responses per respondent	Average burden hours per response	Total burden hours
Program Specific PPRs	900	3	6	16,200

John M. Sweet Jr.,

ACF/OPRE Certifying Officer.

[FR Doc. 2023–01762 Filed 1–27–23; 8:45 am]

BILLING CODE 4184–79–P

DEPARTMENT OF HEALTH AND HUMAN SERVICES

Food and Drug Administration

[Docket No. FDA–2023–N–0187]

Agency Information Collection Activities; Proposed Collection; Comment Request; Premarket Approval of Medical Devices

AGENCY: Food and Drug Administration, HHS.

ACTION: Notice.

SUMMARY: The Food and Drug Administration (FDA or Agency) is announcing an opportunity for public comment on the proposed collection of certain information by the Agency. Under the Paperwork Reduction Act of 1995 (PRA), Federal Agencies are required to publish notice in the **Federal Register** concerning each proposed collection of information, including each proposed extension of an existing collection of information, and to allow 60 days for public comment in response to the notice. This notice solicits comments on information collection associated with premarket approval of medical devices.

DATES: Submit either electronic or written comments on the collection of information by March 31, 2023.

ADDRESSES: You may submit comments as follows. Please note that late, untimely filed comments will not be considered. Electronic comments must be submitted on or before March 31, 2023. The *https://www.regulations.gov* electronic filing system will accept comments until 11:59 p.m. Eastern Time at the end of March 31, 2023. Comments received by mail/hand delivery/courier (for written/paper submissions) will be considered timely if they are postmarked or the delivery service acceptance receipt is on or before that date.

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