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Dated: April 5, 2023.

**William N. Parham, III,**

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

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**DEPARTMENT OF HEALTH AND HUMAN SERVICES**

**Administration for Children and Families**

**Submission for OMB Review; Child Care and Development Fund (CCDF) Consumer Education Website and Reports of Serious Injuries and Death**

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

**ACTION:** Request for Public Comments.

**SUMMARY:** The Office of Child Care (OCC), Administration for Children and Families (ACF), U.S. Department of Health and Human Services (HHS), is

requesting a 3-year extension of the CCDF Consumer Education website and Reports of Serious Incidents and Death (Office of Management and Budget (OMB) #: 0970-0473, expiration date: April 30, 2023). There are no changes requested to the reporting requirements.

**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](http://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](mailto:infocollection@acf.hhs.gov). Identify all emailed requests by the title of the information collection.

**SUPPLEMENTARY INFORMATION:**

*Description:* The existing Consumer Education Website reporting requirement will not be modified and requires states and territories to include information about state or territory policies (related to licensing, monitoring, and background checks) and provider-specific information, including results of monitoring and inspection reports and, if available, information about quality. The existing Reporting of Serious Injuries and Death reporting requirement will not be modified. CCDF Lead Agencies must establish procedures that require child care providers that care for children receiving CCDF subsidies to report to a designated state, territorial, or tribal entity any serious injuries or deaths of children occurring in child care. There are no standard federal forms associated with these reporting requirements.

*Respondents:* The Consumer Education website information collection requirement applies to the 50 states, the District of Columbia, and 5 territories that receive CCDF grants. Reporting of Serious Injuries and Death is a requirement for child care providers.

**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
Consumer Education Website .....	56	1	300	50,400	16,800
Reporting of Serious Injuries and Death .....	10,000	1	1	30,000	10,000

*Estimated Total Annual Burden Hours:* 26,800.

*Authority:* Pub. L. 113-186; 42 U.S.C. 9858 *et seq.*

**Mary B. Jones,**

ACF/OPRE Certifying Officer.

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**DEPARTMENT OF HEALTH AND HUMAN SERVICES**

**Food and Drug Administration**

[Docket No. FDA-2023-N-1006]

**Agency Information Collection Activities; Proposed Collection; Comment Request; Medical Devices; Reports of Corrections and Removals**

**AGENCY:** Food and Drug Administration, HHS.

**ACTION:** Notice.

**SUMMARY:** The Food and Drug Administration (FDA, Agency, or we) 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 reports of removals and corrections for medical and radiation emitting products regulated by FDA’s Center for Devices and Radiological Health.

**DATES:** Either electronic or written comments on the collection of information must be submitted by June 12, 2023.

**ADDRESSES:** You may submit comments as follows. 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 June 12, 2023. Comments received by mail/hand delivery/courier (for written/paper submissions) will be considered timely if they are received 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