

partners to make sure that the evidence is understood and used. Seven current members' terms will expire in November 2021.

DATES: Nominations should be received on or before 60 days after date of publication.

ADDRESSES: Nominations should be sent to Jaime Zimmerman via email at NationalAdvisoryCouncil@ahrq.hhs.gov.

FOR FURTHER INFORMATION CONTACT: Jaime Zimmerman, AHRQ, at (301) 427-1456.

SUPPLEMENTARY INFORMATION: 42 U.S.C. 299c provides that the Secretary shall appoint to the Council twenty one appropriately qualified individuals. At least seventeen members shall be representatives of the public and at least one member shall be a specialist in the rural aspects of one or more of the professions or fields listed below. In addition, the Secretary designates, as ex officio members, representatives from other Federal agencies, principally agencies that conduct or support health care research, as well as Federal officials the Secretary may consider appropriate. 42 U.S.C. 299c(c)(3).

Seven current members' terms will expire in November 2021. To fill these positions, we are seeking individuals who: (1) Are distinguished in the conduct of research, demonstration projects, and evaluations with respect to health care; (2) are distinguished in the fields of health care quality research or health care improvement; (3) are distinguished in the practice of medicine; (4) are distinguished in other health professions; (5) represent the private health care sector (including health plans, providers, and purchasers) or are distinguished as administrators of health care delivery systems; (6) are distinguished in the fields of health care economics, information systems, law, ethics, business, or public policy; and (7) represent the interests of patients and consumers of health care. 42 U.S.C. 299c(c)(2). Individuals are particularly sought with experience and success in these activities. AHRQ will accept nominations to serve on the Council in a representative capacity.

The Council meets in the Washington, DC, metropolitan area, generally in Rockville, Maryland, approximately three times a year to provide broad guidance to the Secretary and AHRQ's Director on the direction of and programs undertaken by AHRQ.

Seven individuals will be selected by the Secretary to serve on the Council beginning with the meeting in the spring of 2022. Members generally serve 3-year terms. Appointments are

staggered to permit an orderly rotation of membership.

Interested persons may nominate one or more qualified persons for membership on the Council. Self-nominations are accepted. Nominations shall include: (1) A copy of the nominee's resume or curriculum vitae; and (2) a statement that the nominee is willing to serve as a member of the Council. Selected candidates will be asked to provide detailed information concerning their financial interests, consultant positions and research grants and contracts, to permit evaluation of possible sources of conflict of interest. Please note that once a candidate is nominated, AHRQ may consider that nomination for future positions on the Council.

The Department seeks a broad geographic representation. In addition, AHRQ conducts and supports research concerning priority populations, which include: Inner city; rural; low income; minority; women; children; elderly; and those with special health care needs, including those who have disabilities, need chronic care, or need end-of-life health care. See 42 U.S.C. 299(c). AHRQ also includes in its definition of priority populations those groups identified in Section 2(a) of Executive Order 13985 as members of underserved communities: Black, Latino, and Indigenous and Native American persons, Asian Americans and Pacific Islanders and other persons of color; members of religious minorities; lesbian, gay, bisexual, transgender, and queer (LGBTQ+) persons; persons with disabilities; persons who live in rural areas; and persons otherwise adversely affected by persistent poverty or inequality. Nominations of persons with expertise in health care for these priority populations are encouraged.

Dated: June 24, 2021.

Marquita Cullom,

Acting Director.

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DEPARTMENT OF HEALTH AND HUMAN SERVICES

Administration for Children and Families

Submission for OMB Review; Head Start Program Performance Standards (OMB #0970-0148)

AGENCY: Office of Head Start, Administration for Children and Families, HHS.

ACTION: Request for public comment.

SUMMARY: The Office of Head Start (OHS), 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 requirements under the Head Start Program Performance Standards (OMB #0970-0148). There are no changes to the information collection.

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: Section 641A of the Head Start Act, 42 U.S.C. 9836A, directs HHS to develop "scientifically based and developmentally appropriate education performance standards related to school readiness" and "ensure that any such revisions in the standards do not result in the elimination of or any reduction in quality, scope, or types of health, educational, parental involvement, nutritional, social, or other services." The Office of Head Start (OHS) announced in the **Federal Register** in 2016 the first comprehensive revision of the Head Start Program Performance Standards (HSPPS) since their original release in 1975. This information collection was approved alongside the final rule for the HSPPS.

This information collection is entirely recordkeeping and does not contain any standardized instruments to provide flexibility for local programs. These records are intended to act as a tool for grantees and delegate agencies to be used in their day-to-day operations. For example, this includes the requirement that programs maintain a waiting list of eligible families. There are no changes to the record keeping requirements.

Respondents: Head Start Grantees. Depending on the standard, the calculated burden hours is based on the individual enrollee (1,054,720), family (956,120), program (3,020), or staff (265,030). In a few cases, only a proportion of one of these may apply.

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 |
|---------------------------------|-----------------------------|------------------------------------------|-----------------------------------|--------------------|---------------------|
| 1301.6(a) | 3,020 | 1 | 0.70 | 2,114 | 2,114 |
| 1302.12(k) | 1,054,720 | 1 | .166 | 175,084 | 175,084 |
| 1302.14(c) | 3,020 | 1 | 2.00 | 6,040 | 6,040 |
| 1302.16(b) | 3,020 | 1 | 5.00 | 15,100 | 15,100 |
| 1302.33(a)–(b) | 1,054,720 | 1 | 1.00 | 1,054,720 | 1,054,720 |
| 1302.33(c)(2) | 294,632 | 1 | 2.00 | 589,264 | 589,264 |
| 1302.42(a)–(b) | 1,054,720 | 1 | 0.66 | 696,115 | 696,115 |
| 1302.42(e) | 3,020 | 1 | 0.50 | 1,510 | 1,510 |
| 1302.47(b)(7)(iv) | 3,020 | 1 | 0.50 | 1,510 | 1,510 |
| 1302.53(b)–(d) | 3,020 | 1 | 0.166 | 501 | 501 |
| 1302.90(a) | 3,020 | 1 | 0.50 | 1,510 | 1,510 |
| 1302.90(b)(1)(i)–(iv), (b)(4) | 79,509 | 1 | 0.33 | 26,238 | 26,238 |
| 1302.93(a) | 26,503 | 1 | 0.25 | 6,626 | 6,626 |
| 1302.94(a) | 3,020 | 1 | 0.166 | 501 | 501 |
| 1302.101(a)(4), 1302.102(b)–(c) | 3,020 | 1 | 79.00 | 238,580 | 238,580 |
| 1302.102(d)(3) | 110 | 1 | 10.00 | 1,100 | 1,100 |
| 1303.12 | 3,020 | 1 | 0.166 | 501 | 501 |
| 1303.22–24 | 956,120 | 1 | 0.33 | 315,520 | 315,520 |
| 1303.42–53 | 260 | 1 | 40.00 | 10,400 | 10,400 |
| 1303.70(c) | 200 | 1 | 1.00 | 200 | 200 |
| 1303.72(a)(3) | 3,020 | 1 | 2.00 | 6,040 | 6,040 |
| 1304.13 | 75 | 1 | 60.00 | 4,500 | 4,500 |
| 1304.15(a) | 400 | 1 | 0.25 | 100 | 100 |

Estimated Total Annual Burden Hours: 3,153,774.

Authority: 42 U.S.C. 9836A.

Mary B. Jones,

ACF/OPRE Certifying Officer.

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DEPARTMENT OF HEALTH AND HUMAN SERVICES

Food and Drug Administration

[Docket No. FDA–2014–N–1960]

Agency Information Collection Activities; Proposed Collection; Comment Request; MedWatch: The Food and Drug Administration Medical Products Reporting Program

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 FDA’s MedWatch adverse experience reporting (AER) program.

DATES: Submit either electronic or written comments on the collection of information by August 30, 2021.

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 August 30, 2021. The <https://www.regulations.gov> electronic filing system will accept comments until 11:59 p.m. Eastern Time at the end of August 30, 2021.

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 that if you include your name, contact information, or other information that identifies you in the body of your comments, that information will be posted on <https://www.regulations.gov>.

- If you want to submit a comment with confidential information that you do not wish to be made available to the public, submit the comment as a written/paper submission and in the manner detailed (see “Written/Paper Submissions” and “Instructions”).

Written/Paper Submissions

Submit written/paper submissions as follows:

- **Mail/Hand delivery/Courier (for written/paper submissions):** Dockets Management Staff (HFA–305), Food and Drug Administration, 5630 Fishers Lane, Rm. 1061, Rockville, MD 20852.

- For written/paper comments submitted to the Dockets Management Staff, FDA will post your comment, as well as any attachments, except for information submitted, marked, and identified, as confidential, if submitted as detailed in “Instructions.”

Instructions: All submissions received must include the Docket No. FDA–2014–N–1960 for “MedWatch: The FDA Medical Products Reporting Program.” Received comments, those filed in a timely manner (see **ADDRESSES**), will be placed in the docket and, except for those submitted as “Confidential