Because FDA is developing two new training programs, Trainee Program and Reagan-Udall Fellowship, our estimated burden for the information collection reflects an overall increase of 2 hours. FDA has removed the Commissioner's Fellowship Program and Regulatory Science Internship Program from this information collection as these programs have been discontinued.

FDA published a 30-day notice for this information collection on February 3, 2020 (85 FR 5966). FDA is reopening the 30-day comment period in order to satisfy PRA requirements. No changes have been made to the information collection.

Dated: December 14, 2020.

Lauren K. Roth,

Acting Principal Associate Commissioner for Policy.

[FR Doc. 2020–27963 Filed 12–17–20; 8:45 am]

BILLING CODE 4164–01–P

DEPARTMENT OF HEALTH AND HUMAN SERVICES

Food and Drug Administration

[Docket No. FDA–2020–N–1359]

Sugars That Are Metabolized Differently Than Traditional Sugars; Extension of Comment Period

AGENCY: Food and Drug Administration, HHS.

ACTION: Notification; extension of comment period.

SUMMARY: The Food and Drug Administration (FDA or we) is extending the comment period for the notice entitled ''Sugars that Are Metabolized Differently than Traditional Sugars'' that appeared in the **Federal Register** of October 19, 2020. We are taking this action in response to a request from stakeholders to extend the comment period to allow additional time for interested persons to develop and submit data, information, and/or comments for this request for information and comments.

DATES: FDA is extending the comment period on the notice published October 19, 2020 (85 FR 66335). Submit either electronic or written comments on the notice by February 16, 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 February 16, 2021. The *https://www.regulations.gov* electronic filing system will accept comments until 11:59 p.m. Eastern Time at the end of February 16, 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–2020–N–1359 for ''Sugars that Are Metabolized Differently than Traditional Sugars.'' Received comments, those filed in a timely manner (see **ADDRESSES**), will be placed in the docket and, except for those submitted as ''Confidential Submissions,'' publicly viewable at *https://www.regulations.gov* or at the Dockets Management Staff between 9 a.m. and 4 p.m. Eastern Time, Monday through Friday, 240–402–7500.

- Confidential Submissions—To submit a comment with confidential information that you do not wish to be made publicly available, submit your comments only as a written/paper submission. You should submit two copies total. One copy will include the information you claim to be confidential with a heading or cover note that states ''THIS DOCUMENT CONTAINS CONFIDENTIAL INFORMATION.'' We will review this copy, including the claimed confidential information, in our consideration of comments. The second copy, which will have the claimed confidential information redacted/blacked out, will be available for public viewing and posted on *https://www.regulations.gov.* Submit both copies to the Dockets Management Staff. If you do not wish your name and contact information to be made publicly available, you can provide this information on the cover sheet and not in the body of your comments and you must identify this information as ''confidential.'' Any information marked as ''confidential'' will not be disclosed except in accordance with 21 CFR 10.20 and other applicable disclosure law. For more information about FDA's posting of comments to public dockets, see 80 FR 56469, September 18, 2015, or access the information at: *https://www.govinfo.gov/content/pkg/FR-2015-09-18/pdf/2015-23389.pdf*.

Docket: For access to the docket to read background documents or the electronic and written/paper comments received, go to *https://www.regulations.gov* and insert the docket number, found in brackets in the heading of this document, into the ''Search'' box and follow the prompts and/or go to the Dockets Management Staff, 5630 Fishers Lane, Rm. 1061, Rockville, MD 20852, 240–402–7500.

FOR FURTHER INFORMATION CONTACT: Blakeley Fitzpatrick, Office of Nutrition and Food Labeling, Center for Food Safety and Applied Nutrition, Food and Drug Administration, 5001 Campus Dr., College Park, MD 20740, 240–402–1450.

SUPPLEMENTARY INFORMATION:

I. Background

In the *Federal Register* of October 19, 2020 (85 FR 66335), we published a notice requesting information and comments entitled ''Sugars that Are Metabolized Differently Than Traditional Sugars.'' This action opened a docket with a 60-day comment period to receive information and comments related to the nutrition labeling of sugars that are metabolized differently than traditional sugars.

FDA has received a request for a 60-day extension for this comment period in order to allow additional time for

interested persons to develop and submit data, information, and/or comments for this notice. We have concluded that it is reasonable to extend the comment period for 60 days. FDA believes that this extension allows adequate time for any interested persons to submit data, information, and/or comments.

Dated: December 11, 2020.

Lauren K. Roth,

Acting Principal Associate Commissioner for Policy.

[FR Doc. 2020–27749 Filed 12–17–20; 8:45 am]

BILLING CODE 4164–01–P

DEPARTMENT OF HEALTH AND HUMAN SERVICES

Health Resources and Services Administration

Agency Information Collection Activities: Proposed Collection: Public Comment Request; Information Collection Request Title: The Maternal, Infant, and Early Childhood Home Visiting Program Performance Measurement Information System, OMB No. 0906–0017, Revision

AGENCY: Health Resources and Services Administration (HRSA), Department of Health and Human Services.

ACTION: Notice.

SUMMARY: In compliance with the requirement for opportunity for public comment on proposed data collection projects of the Paperwork Reduction Act of 1995, HRSA announces plans to submit an Information Collection Request (ICR), described below, to the Office of Management and Budget (OMB). Prior to submitting the ICR to OMB, HRSA seeks comments from the public regarding the burden estimate, below, or any other aspect of the ICR.

DATES: Comments on this ICR should be received no later than February 16, 2021.

ADDRESSES: Submit your comments to *paperwork@hrsa.gov* or mail the HRSA Information Collection Clearance Officer, 14N136B, 5600 Fishers Lane, Rockville, MD 20857.

FOR FURTHER INFORMATION CONTACT: To request more information on the proposed project or to obtain a copy of the data collection plans and draft instruments, email *paperwork@hrsa.gov* or call Lisa Wright-Solomon, the HRSA Information Collection Clearance Officer at (301) 443–1984.

SUPPLEMENTARY INFORMATION:

Information Collection Request Title: The Maternal, Infant, and Early Childhood Home Visiting Program Performance Measurement Information System, OMB No. 0906–0017, Revision.

Abstract: This clearance request is for continued approval of the Maternal, Infant, and Early Childhood Home Visiting (MIECHV) Program Performance Measurement Information System.

The MIECHV Program, administered by HRSA in partnership with the Administration for Children and Families, supports voluntary, evidence-based home visiting services to pregnant women and to parents with young children up to kindergarten entry. States, certain non-profit organizations, and Tribal entities are eligible to receive funding from the MIECHV program and have the flexibility to tailor the program to serve the specific needs of their communities. HRSA is revising the data collection forms for the MIECHV program by making the following changes:

- *Form 1, Table 1:* Update table to include reporting for gender non-conforming participants and unknown/did not report participant gender.
- *Form 1, Tables 3, 4, 6, 7, 8, 9, 10, 11, and 18:* Update tables to include reporting for gender non-conforming participants and unknown/did not report adult participant gender.
- *Form 1, Tables 3, 5, 6, 7, 18, 19, and 20:* Update tables to remove index child gender reporting.
- *Form 1, Table 15:* Change table title to ''Home Visits''.
- *Form 1, Table 15:* Update table to collect the number of home visits completed virtually.
- *Form 1, Tables 4, 9, 10, and 18:* Update tables to include reporting for new and continuing adult participants.
- *Form 1, Tables 5, 19, and 20:* Update tables to include reporting for new and continuing index children.
- *Form 1, Table 16:* Add new table to include reporting on father and additional caregiver engagement.
- *Form 1, Tables 16, 17, 18, 19, 20, and 21:* Update table numbers to reflect the addition of Table 16.
- *Form 2, Measure 13:* Change measure name to ''Behavioral Concern Inquiries''
- *Form 2, Measure 16:* Update measure to reflect caregiver health insurance coverage status.
- *Form 2:* Add two measures to collect information on substance use screening and referrals.

Need and Proposed Use of the Information: HRSA uses performance information to demonstrate program accountability and continuously monitor and provide oversight to MIECHV Program awardees. The information is also used to provide quality improvement guidance and technical assistance to awardees and help inform the development of early childhood systems at the national, state, and local level. HRSA is seeking to revise demographic, service utilization, and select clinical indicators for participants enrolled in home visiting services. In addition, HRSA will collect a set of standardized performance and outcome indicators that correspond with the statutorily identified benchmark areas.

This information will be used to demonstrate awardees' compliance with legislative and programmatic requirements. It will also be used to monitor and provide continued oversight for awardee performance and to target technical assistance resources to awardees. In the future, HRSA anticipates that MIECHV funding decisions may be allocated, in part, based on awardee performance, including on benchmark performance areas. This notice is subject to the appropriation of funds, and is a contingency action taken to ensure that, should funds become available for this purpose, information can be collected in a timely manner.

Likely Respondents: MIECHV Program awardees that are states, territories, and, where applicable, nonprofit organizations providing home visiting services within states.

Burden Statement: Burden in this context means the time expended by persons to generate, maintain, retain, disclose, or provide the information requested. This includes the time needed to review instructions; to develop, acquire, install, and utilize technology and systems for the purpose of collecting, validating, and verifying information, processing and maintaining information, and disclosing and providing information; to train personnel and to be able to respond to a collection of information; to search data sources; to complete and review the collection of information; and to transmit or otherwise disclose the information. The total annual burden hours estimated for this ICR are summarized in the table below.