To obtain copies of a supporting statement and any related forms for the proposed collection(s) summarized in this notice, you may make your request using one of following:

1. Access CMS' website address at website address at *https://www.cms.gov/Regulations-and-Guidance/Legislation/PaperworkReductionActof1995/PRA-Listing.html.*

2. Call the Reports Clearance Office at (410) 786–1326.

**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–10390 Hospice Quality Reporting Program**

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 without change; *Title of Information Collection:* Hospice Quality Reporting Program; *Use:* The Hospice Item Set (HIS) is a standardized, patient-level data collection tool developed specifically for use by hospices. It is currently used for the collection of quality measure data pertaining to the Hospice Quality Reporting Program (HQRP). Since April 1, 2017, hospices have been using the HIS V2.00.0 which specifies the collection of data items that support eight National Quality Forum (NQF) endorsed Quality Measures (QMs) and an additional measure pair for hospice. All Medicare-certified hospice providers are required to submit HIS admission and discharge records to CMS for each patient admission and discharge. The HIS contains data elements that are used by the CMS to calculate these measures and also allows CMS to collect quality data from hospices in compliance with Section 3004 of the Affordable Care Act. The information collection request was revised to remove Section O of the HIS discharge assessment now that we proposed to replace it with the claims-based Hospice Visits in the Last Days of Life quality measure. *Form Number:* CMS–10390 (OMB control number: 0938–1153); *Frequency:* On Occasion; *Affected Public:* State, Local, or Tribal Governments, Private Sector (not-for-profit institutions); individuals or households; *Number of Respondents:* 4,688; *Total Annual Responses:* 1,328,417; *Total Annual Hours:* 636,312. (For policy questions regarding this collection contact Cindy Massuda at (410) 786–0652.)

Dated: August 10, 2020.

**William N. Parham, III,**

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

[FR Doc. 2020–17738 Filed 8–12–20; 8:45 am]

**BILLING CODE 4120–01–P**

## DEPARTMENT OF HEALTH AND HUMAN SERVICES

### Administration for Children and Families

### Submission for OMB Review; Youth Empowerment Information, Data Collection, and Exploration on Avoidance of Sex (IDEAS) (New Collection)

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

**ACTION:** Request for public comment.

**SUMMARY:** The Office of Planning, Research, and Evaluation (OPRE), Administration for Children and Families (ACF), U.S. Department of Health and Human Services (HHS), proposes survey data collection activities as part of the Youth Empowerment IDEAS study.

**DATES:** *Comments due within 30 days of publication.* OMB is required to make a decision concerning 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 directly to the following:

Office of Management and Budget, Paperwork Reduction Project, Email: *OIRA_SUBMISSION@OMB.EOP.GOV,* Attn: Desk Officer for the Administration for Children and Families.

Copies of the proposed collection may be obtained by emailing *OPREinfocollection@acf.hhs.gov.* Alternatively, copies can also be obtained by writing to the Administration for Children and Families, Office of Planning, Research and Evaluation, 330 C Street SW, Washington, DC 20201, Attn: OPRE Reports Clearance Officer. All requests, emailed or written, should be identified by the title of the information collection.

**SUPPLEMENTARY INFORMATION:**

*Description:* OPRE/ACF/HHS proposes data collection activities as part of the Youth Empowerment IDEAS study. The goal of this project is to collect descriptive data that will inform educational topics and strategies for adolescent pregnancy prevention and youth health and well-being. The project will identify messages and themes that are most likely to resonate with youth. The project will inform hypotheses on how to increase the effectiveness of sex education approaches so that more youth avoid the risks associated with teen sex and teen pregnancy rates are reduced. To support these efforts, we seek approval from the Office of Management and Budget to collect survey information from youth and young adults ages 14–24 and of parents of teens ages 14–18 using an online panel that is based on a probability-based sample of the U.S. population. We propose the following data collection instruments:

(1) *Parent Survey:* We will administer this as a web survey. Information collected through the Parent Survey will be used to report on demographics, the parent-child relationship, parents' attitudes and beliefs about youth sex education and sexual behaviors, and parental knowledge about youth sexual risk-taking.

(2) *Youth Survey:* We will administer a web survey in two parts to youth ages 14–18. Information collected on Part I of the survey will be used to report on demographics, the parent-child relationship, future aspirations, and attitudes and beliefs about youth sexual behavior. Information collected on Part II of the survey will include knowledge about sexual risk, experience with sex education, and sexual risk behaviors.

(3) *Young Adult Survey:* We will administer this to young adults ages 19–24 as a web survey. Topics align with the youth survey, but with slight wording changes to reflect the older population.

*Respondents:* The survey respondents are from an online panel of a probability-based sample of the U.S. population of parents of youth ages 14–18 and their youth ages 14–18 and of young adults ages 19–24.

ANNUAL BURDEN ESTIMATES

| Instrument | Number of respondents (total over request period) | Number of responses per respondent (total over request period) | Avg. burden per response (in hours) | Total burden (in hours) | Annual burden (in hours) |
|---|---|---|---|---|---|
| (1) Parent Survey | 1,550 | 1 | .333 | 516 | 172 |
| (2) Part I Youth Survey | 675 | 1 | .333 | 225 | 75 |
| (3) Part II Youth Survey | 590 | 1 | .333 | 197 | 66 |
| (4) Young Adult Survey | 775 | 1 | .583 | 452 | 151 |

*Estimated Total Annual Burden Hours:* 464.

(Authority: Sec. 510. [42 U.S.C. 710])

**John M. Sweet,**

*ACF/OPRE Certifying Officer.*

[FR Doc. 2020–17680 Filed 8–12–20; 8:45 am]

**BILLING CODE 4184–83–P**

## DEPARTMENT OF HEALTH AND HUMAN SERVICES

### Food and Drug Administration

**[Docket No. FDA–2020–N–1550]**

### New Drugs Regulatory Program Modernization: Implementation of the Integrated Assessment of Marketing Applications and Integrated Review Documentation; Public Workshop; Request for Comments

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

**ACTION:** Notice of public workshop; request for comments.

**SUMMARY:** The Food and Drug Administration (FDA or the Agency) is announcing the following public workshop entitled ''New Drugs Regulatory Program Modernization: Implementation of the Integrated Assessment of Marketing Applications and Integrated Review Documentation.'' The purpose of the public workshop is to seek public comments/feedback on the Integrated Review documentation generated by the new Integrated Assessment of marketing applications for new drug products developed as part of the New Drugs Regulatory Program Modernization. The Agency hopes to receive public feedback on how this Integrated Review documentation can continue supporting our stakeholders' needs. Please see information and examples relevant to the Integrated Review at *http://wcms-internet.fda.gov/drugs/news-events-human-drugs/integrated-assessment-marketing-applications-workshop-10302020-10302020.*

**DATES:** The public workshop will be held virtually and broadcast via webcast only on October 30, 2020, from 9 a.m. to 3 p.m. Registration to attend the meeting and other information can be found at *http://wcms-internet.fda.gov/drugs/news-events-human-drugs/integrated-assessment-marketing-applications-workshop-10302020-10302020.* The public meeting may be extended or may end early depending on the level of public participation. Submit either electronic or written comments on this public workshop by December 30, 2020. See the **SUPPLEMENTARY INFORMATION** section for registration date and information.

**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 December 30, 2020. The *https://www.regulations.gov* electronic filing system will accept comments until 11:59 p.m. Eastern Time at the end of December 30, 2020. 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–1550 for ''New Drugs Regulatory Program Modernization: Implementation of the Integrated Assessment of Marketing Applications and Integrated Review Documentation.'' 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., Monday through Friday, 240–402–7500.