DEPARTMENT OF HEALTH AND HUMAN SERVICES

Administration for Children and Families

Proposed Information Collection Activity; 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), at the U.S. Department of Health and Human Services (HHS), proposes data collection activities as part of the Youth Empowerment IDEAS study. The goal of this project is to collect data that will inform educational topics and strategies for an optimal-health sexual risk avoidance (SRA) approach to reducing teen pregnancy and improving youth well-being. The project will identify strategies, skills, 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 OMB approval to collect survey information from a nationally-representative sample of youth and young adults age 14–24 and a nationally-representative sample of parents of teens ages 14–18.

DATES: *Comments due within 60 days of publication.* In compliance with the requirements of Section 3506(c)(2)(A) of the Paperwork Reduction Act of 1995, the Administration for Children and Families is soliciting public comment on the specific aspects of the information collection described above.

ADDRESSES: Copies of the proposed collection of information can be obtained and comments may be forwarded 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. Email address: *OPREinfocollection@acf.hhs.gov.* All requests should be identified by the title of the information collection.

SUPPLEMENTARY INFORMATION:

Description: We propose the following data collection instruments:

(1) *Parent 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. We will use both random-digit-dialing and a web survey.

(2) *Youth Survey:* We will administer a web survey in two parts. 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.

Respondents: A nationally representative sample of parents of teens ages 14–18 and a nationally representative sample of youth and young adults ages 14–24.

ANNUAL BURDEN ESTIMATES

Instrument	Total number of respondents	Annual number of respondents	Number of responses per respondent	Average burden hours per response	Annual burden hours
Parent Survey—Screener	7500	2500	1	.083	208
Parent Survey—Telephone Mode (RDD)	600	200	1	.500	100
Parent Survey—Web	900	300	1	.333	100
Part I Youth Web Survey	1500	500	1	.333	167
Part II Youth Web Survey	1200	400	1	.333	133

Estimated Total Annual Burden Hours: 708.

Comments: The Department specifically requests comments on (a) whether the proposed collection of information is necessary for the proper performance of the functions of the agency, including whether the information shall have practical utility; (b) the accuracy of the agency's estimate of the burden of the proposed collection of information; (c) the quality, utility, and clarity of the information to be collected; and (d) ways to minimize the burden of the collection of information on respondents, including through the use of automated collection techniques or other forms of information technology. Consideration will be given to comments and suggestions submitted within 60 days of this publication.

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

Mary B. Jones,

ACF/OPRE Certifying Officer.

[FR Doc. 2019–01566 Filed 2–7–19; 8:45 am]

BILLING CODE 4184–83–P

DEPARTMENT OF HEALTH AND HUMAN SERVICES

Administration for Community Living

Administration on Intellectual and Developmental Disabilities, President's Committee for People With Intellectual Disabilities

AGENCY: Administration for Community Living, HHS.

ACTION: Notice.

SUMMARY: The President's Committee for People with Intellectual Disabilities (PCPID) will host a webinar/conference call for its members to discuss the potential topics of the Committee's 2019 Report to the President. All the PCPID meetings, in any format, are open to the public. This virtual meeting will be conducted in a discussion format.

DATES:

Webinar/Conference Call: Monday, March 4, 2019 from 11:00 a.m. to 12:30 p.m. (EST).

Agenda: The Committee will discuss the preparation of the PCPID 2019 Report to the President, including its content and format, and related data collection and analysis required to complete the writing of the Report.

Additional Information: For further information and accommodations needs, please contact Ms. Allison Cruz, Director, Office of Innovation, 330 C Street, SW, Switzer Building, Room 1114, Washington, DC 20201. Telephone: 202–795–7334. Fax: 202–

795–7334. Email: *allison.cruz@acl.hhs.gov*

Supplemental Information: The purpose of this virtual meeting is to discuss the Committee's preparation of the 2019 Report to the President, including its content and format, and related data collection and analysis required to complete the writing of the Report.

Webinar/Conference Call: The webinar/conference call is scheduled for Monday, March 4, 2019 from 11:00 a.m. to 12:30 p.m. (EST) and may end early if discussions are finished.

Instructions to Participate in the Webinar/Conference Call on Monday, March 4, 2019: Please dial: (888) 949–2790; Pass Code: 1989852.

Background Information on the Committee: The PCPID acts in an advisory capacity to the President and the Secretary of Health and Human Services on a broad range of topics relating to programs, services and support for individuals with intellectual disabilities. The PCPID executive order stipulates that the Committee shall: (1) Provide such advice concerning intellectual disabilities as the President or the Secretary of Health and Human Services may request; and (2) provide advice to the President concerning the following for people with intellectual disabilities: (A) Expanding employment opportunities; (B) connecting people to services; (C) supporting families and caregivers; (D) strengthening the networks; and (E) protecting rights and preventing abuse.

Dated: February 4, 2018.

Julie Hocker,

Commissioner, Administration on Disabilities (AoD).

[FR Doc. 2019–01698 Filed 2–7–19; 8:45 am]

BILLING CODE 4154–01–P

DEPARTMENT OF HEALTH AND HUMAN SERVICES

Food and Drug Administration

[Docket No. FDA–2018–N–3918]

Request for Nomination From Industry Organizations Interested in Participating in the Selection Process for Nonvoting Industry Representatives and Request for Nominations for Nonvoting Industry Representatives on Public Advisory Committees

AGENCY: Food and Drug Administration, HHS.

ACTION: Notice.

SUMMARY: The Food and Drug Administration (FDA or the Agency) is requesting that any industry organizations interested in participating in the selection of nonvoting industry representatives to serve on its public advisory committees for the Center for Drug Evaluation and Research (CDER) notify FDA in writing. FDA is also requesting nominations for nonvoting industry representatives to serve on CDER's public advisory committees. A nominee may either be self-nominated or nominated by an organization to serve as a nonvoting industry representative. Nominations will be accepted for vacancies which become available on November 1, 2019, for the 4-year term of November 1, 2019 to October 31, 2023.

DATES: Any industry organization interested in participating in the selection of an appropriate nonvoting member to represent industry interests must send a letter stating that interest to FDA by March 11, 2019, (see sections I and II of this document for further details). Concurrently, nomination materials for prospective candidates should be sent to FDA by March 11, 2019.

ADDRESSES: All statements of interest from industry organizations interested in participating in the selection process of nonvoting industry representative nominations should be sent to Cicely Reese (see **FOR FURTHER INFORMATION CONTACT**). All nominations for nonvoting industry representatives may be submitted electronically by accessing the FDA Advisory Committee Membership Nomination Portal: *https://www.accessdata.fda.gov/scripts/FACTRSPortal/FACTRS/index.cfm* or by mail to: Division of Advisory Committee and Consultant Management, Food and Drug Administration, 10903 New Hampshire Ave., Bldg. 31, Rm. 2417, Silver Spring, MD 20993–0002. Information about becoming a member of an FDA advisory committee can also be obtained by visiting FDA's website at: *http://www.fda.gov/AdvisoryCommittees/default.htm.*

FOR FURTHER INFORMATION CONTACT: Cicely Reese, Center for Drug Evaluation and Research, Food and Drug Administration, 10903 New Hampshire Ave., Bldg. 31, Rm. 2417, Silver Spring, MD 20993–0002, 301–796–9001, Fax: 301–847–8533, email: *Cicely.Reese@fda.hhs.gov.*

SUPPLEMENTARY INFORMATION: The Agency intends to add a nonvoting industry representative to the following advisory committees:

I. CDER Advisory Committees

A. Anesthetic and Analgesic Drug Products Advisory Committee

Reviews and evaluates available data concerning the safety and effectiveness of marketed and investigational human drug products for use in anesthesiology and surgery.

B. Antimicrobial Drugs Advisory Committee

Reviews and evaluates available data concerning the safety and effectiveness of marketed and investigational human drug products for use in the treatment of infectious diseases and disorders.

C. Arthritis Advisory Committee

Reviews and evaluates available data concerning the safety and effectiveness of marketed and investigational human drug products for use in the treatment of arthritis, rheumatism, and related diseases.

D. Bone, Reproductive, and Urologic Drugs Advisory Committee

Reviews and evaluates available data on the safety and effectiveness of marketed and investigational human drugs for use in the practice of obstetrics, gynecology, urology, and related specialties.

E. Cardiovascular and Renal Drugs Advisory Committee

Reviews and evaluates available data on the safety and effectiveness of marketed and investigational human drug products for use in the treatment of cardiovascular and renal disorders.

F. Dermatologic and Ophthalmic Drugs Advisory Committee

Reviews and evaluates available data concerning the safety and effectiveness of marketed and investigational human drug products for use in the treatment of dermatologic and ophthalmic disorders.

G. Drug Safety and Risk Management Advisory Committee

Reviews and evaluates information on risk management, risk communication, and quantitative evaluation of spontaneous reports for drugs for human use.

H. Endocrinologic and Metabolic Drugs Advisory Committee

Reviews and evaluates available data concerning the safety and effectiveness of marketed and investigational human drug products for use in the treatment of endocrine and metabolic disorders.