During the data collection period, researchers will follow a subset of 900 infants until 2-years of age. A parent of each of these infants will answer a questionnaire at 6, 9, 12, 18, and 24 months, as well as have other clinical assessments performed to examine developmental delays.

CDC will use study results to guide recommendations made by both INS and CDC to prevent ZIKV infection; to improve counseling of patients about risks to themselves, their pregnancies, their partners, and their infants; and to help agencies prepare to provide services to affected children and families. Participation in this study is voluntary and there are no costs to participants other than their time.

The total burden hours are 14,210.

ESTIMATED ANNUALIZED BURDEN HOURS

Type of respondents	Form name	Number of respondents	Number of responses per respondent	Average burden per response (in hours)
Pregnant Women	Pregnant Women Eligibility Questionnaire	600	1	5/60
	Pregnant Women Enrollment Questionnaire	500	1	35/60
	Adult Symptoms Questionnaire	500	15	10/60
	Pregnant Women Follow-up Questionnaire	500	8	15/60
	Infant Symptoms Questionnaire	2,250	14	10/60
	Parent-Child Eligibility Questionnaire	1,000	1	5/60
	Parent-Child Enrollment Questionnaire	900	1	20/60
	Parent-Child Follow-up Questionnaire	900	4	15/60
	Ages and Stages Questionnaire: 2 and 6 Month Visits.	2,250	2	15/60
	Ages and Stages Questionnaire: 12 and 24 Month Visits.	900	2	15/60
	Bayley Scales of Infant and Toddler Development.	900	3	30/60
	Strengths and Difficulties Questionnaire	900	1	5/60
	Peabody Developmental Motor Scales	900	1	30/60
	Parenting Stress Index IV	900	5	10/60
	Center for Epidemiologic Studies Depression Scale.	900	5	5/60
	Test of Nonverbal Intelligence	900	1	20/60
Male partners	Male Partner Eligibility Questionnaire	150	1	5/60
	Male Enrollment Questionnaire	125	1	25/60
	Adult Symptoms Questionnaire	125	7	10/60

Leroy A. Richardson,

Chief, Information Collection Review Office, Office of Scientific Integrity, Office of the Associate Director for Science, Office of the Director, Centers for Disease Control and Prevention.

[FR Doc. 2017–25495 Filed 11–24–17; 8:45 am]

BILLING CODE 4163–18–P

DEPARTMENT OF HEALTH AND HUMAN SERVICES

Administration for Children and Families

Submission for OMB Review; Comment Request

Title: Personal Responsibility Education Program (PREP) Multi-Component Evaluation Extension.

OMB No.: 0970–0398.

Description: The Family and Youth Services Bureau (FYSB) and the Office of Planning, Research, Evaluation (OPRE) in the Administration for Children and Families (ACF) are requesting an extension without change of a currently approved information collection (OMB No. 0970–0398). The purpose of the extension is to complete the ongoing follow-up data collection for the Personal Responsibility Education Program (PREP) Multi-Component Evaluation, which was designed to document how PREP programs are designed and implemented in the field, collect performance measure data for PREP programs, and assess the effectiveness of selected PREP-funded programs.

The PREP Multi-Component Evaluation contains three components: A Design and Implementation Study, a Performance Analysis Study, and an Impact and In-Depth Implementation Study. Data collection related to the Design and Implementation Study is complete; data collection related to the Performance Analysis Study will be complete in late summer 2017. This notice is specific to data collection activities for the Impact and In-Depth Implementation Study, which is being conducted in four sites. The proposed extension is necessary to complete ongoing follow-up data collection. The resulting data will be used in a rigorous program impact analysis to assess the effectiveness of each program in reducing teen sexual activity and associated risk behaviors.

Respondents: Youth participants who agreed to participate in the study upon enrollment in the four impact study sites.

ANNUAL BURDEN ESTIMATES

Instrument	Total/annual number of respondents	Number of responses per respondents	Average burden hours per response	Total/annual burden hours
Second follow-up survey	325	1	.75	244

Estimated Total/Annual Burden Hours: 244.

Additional Information: Copies of the proposed collection may 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 should be identified by the title of the information collection. Email address: *OPREinfocollection@acf.hhs.gov.*

OMB Comment: 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. 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.

Mary Jones,

ACF/OPRE Certifying Officer.

[FR Doc. 2017–25444 Filed 11–24–17; 8:45 am]

BILLING CODE 4184–37–P

DEPARTMENT OF HEALTH AND HUMAN SERVICES

Food and Drug Administration

[Docket Nos. FDA–2010–N–0601, FDA–2010–N–0598, FDA–2010–N–0600, FDA–2007–N–0037, FDA–2010–N–0597, FDA–2011–N–0017, and FDA–2016–N–2496]

Agency Information Collection Activities; Announcement of Office of Management and Budget Approvals

AGENCY: Food and Drug Administration, HHS.

ACTION: Notice.

SUMMARY: The Food and Drug Administration (FDA) is publishing a list of information collections that have been approved by the Office of Management and Budget (OMB) under the Paperwork Reduction Act of 1995.

FOR FURTHER INFORMATION CONTACT: Ila S. Mizrachi, Office of Operations, Food and Drug Administration, Three White Flint North, 10A–12M, 11601 Landsdown St., North Bethesda, MD 20852, 301–796–7726, *PRAStaff@fda.hhs.gov.*

SUPPLEMENTARY INFORMATION: The following is a list of FDA information collections recently approved by OMB under section 3507 of the Paperwork Reduction Act of 1995 (44 U.S.C. 3507). The OMB control number and expiration date of OMB approval for each information collection are shown in Table 1. Copies of the supporting statements for the information collections are available on the internet at *https://www.reginfo.gov/public/do/PRAMain.* An Agency may not conduct or sponsor, and a person is not required to respond to, a collection of information unless it displays a currently valid OMB control number.

TABLE 1—LIST OF INFORMATION COLLECTIONS APPROVED BY OMB

Title of collection	OMB control No.	Date approval expires
Current Good Manufacturing Practice for Medicated Feeds	0910–0152	8/31/2020
Current Good Manufacturing Practice for Type A Medicated Articles	0910–0154	8/31/2020
Animal Drug User Fee Cover Sheet, Form FDA 3546	0910–0539	8/31/2020
Animal Drug User Fee Waivers and Reductions	0910–0540	8/31/2020
Index of Legally Marketed Unappropriated New Animal Drugs for Minor Species	0910–0620	8/31/2020
Voluntary National Retail Food Regulatory Program Standards	0910–0621	8/31/2020
Impact Trade Auxiliary Communication System	0910–0842	8/31/2020

Dated: November 20, 2017.

Leslie Kux,

Associate Commissioner for Policy.

[FR Doc. 2017–25452 Filed 11–24–17; 8:45 am]

BILLING CODE 4164–01–P

DEPARTMENT OF HEALTH AND HUMAN SERVICES

Food and Drug Administration

[Docket No. FDA–2010–N–0161]

Agency Information Collection Activities; Proposed Collection; Comment Request; Export of Food and Drug Administration-Regulated Products: Export Certificates

AGENCY: Food and Drug Administration, HHS.

ACTION: Notice.

SUMMARY: The Food and Drug Administration (FDA or Agency) 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 export certificates for the export of FDA-regulated products.

DATES: Submit either electronic or written comments on the collection of information by January 26, 2018.

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 January 26, 2018. The *https://www.regulations.gov* electronic filing system will accept comments until midnight Eastern Time at the end of January 26, 2018. 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