Safety Surveys with the exception of the inclusion of addressed based sampling (ABS) methods to explore the method as a possible alternative for new survey questions. ABS is sampling from address frames that are usually based, in part, on residential addresses in the U.S. Postal Service Computerized Delivery Sequence File. ABS is a cost effective method of sampling that provides much coverage of U.S. households for in-person, mail, telephone, and multimode surveys (including Web-based surveys.) The Food Safety Survey will continue to include cell phones in addition to landlines for the telephone interviews. A nationally representative sample of 4,000 adults will be selected at random to complete the survey. The survey will also include an oversample of Hispanics and Blacks to ensure a minimum of 400 each. Additionally, methods will be employed to test for the presence of response bias. Participation in the survey will be voluntary. Cognitive interviews and a pre-test will be conducted prior to fielding the survey.

FDA estimates the burden of this collection of information as follows:

TABLE 1—ESTIMATED ANNUAL REPORTING BURDEN [1]

Activity	Number of respondents	Number of responses per respondent	Total annual responses	Average burden per response	Total hours
Cognitive interview screener	75	1	75	0.083 (5 minutes)	6
Cognitive interview	9	1	9	1	9
Pretest screener	45	1	45	0.0167 (1 minute)	1
Pretest	18	1	18	0.33 (20 minutes)	6
Survey screener	10,000	1	10,000	0.0167 (1 minute)	167
Survey	4,000	1	4,000	0.33 (20 minutes)	1,320
Non-response survey screener	125	1	125	0.0167 (1 minute)	2
Non-response survey	50	1	50	0.167 (10 minutes)	8
Total [2]					1,519

[1] There are no capital costs or operating and maintenance costs associated with this collection of information.

FDA's burden estimate is based on the Agency's prior experience with the Food Safety Survey. FDA estimates that the burden hours for this information collection will remain the same since the last OMB approval.

Dated: June 27, 2017.

Anna K. Abram,

Deputy Commissioner for Policy, Planning, Legislation, and Analysis.

[FR Doc. 2017–13882 Filed 6–30–17; 8:45 am]

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DEPARTMENT OF HEALTH AND HUMAN SERVICES

Food and Drug Administration

[Docket No. FDA–2017–N–0001]

Advisory Committees; Filing of Closed Meeting Reports

AGENCY: Food and Drug Administration, HHS.

ACTION: Notice.

SUMMARY: The Food and Drug Administration (FDA) is announcing that, as required by the Federal Advisory Committee Act, the Agency has filed with the Library of Congress the annual reports of those FDA advisory committees that held closed meetings during fiscal year 2016.

ADDRESSES: Copies are available at the Dockets Management Staff (HFA–305), Food and Drug Administration, 5630 Fishers Lane, Rm. 1061, Rockville, MD 20852, 240–402–7500. You also may access the docket at *https://www.regulations.gov* for the annual reports of those FDA advisory committees that held closed meetings during fiscal year 2016. Insert the docket number found in brackets in the heading of this document at *https://www.regulations.gov* into the ''Search'' box, clear filter under Document Type (left side of screen), and check ''Supporting and Related Material,'' then Sort By Best Match (from the drop-down menu; top right side of screen), ''ID Number (Z–A)'' or Sort By Best Match (from the drop-down menu) ''Title (A–Z),'' also found in the heading of this document.

FOR FURTHER INFORMATION CONTACT: Michael Ortwerth, Director and Committee Management Officer, Advisory Committee and Oversight Management Staff, Food and Drug Administration, 10903 New Hampshire Ave., Silver Spring, MD 20993–0002, 301–796–8220.

SUPPLEMENTARY INFORMATION: Under section 10(d) of the Federal Advisory Committee Act (5 U.S.C. app.) and 21 CFR 14.60(d), FDA has filed with the Library of Congress the annual reports for the following FDA advisory committees that held closed meetings during the period October 1, 2015, through September 30, 2016:

Center for Biologics Evaluation and Research

Blood Products Advisory Committee
Vaccines and Related Biological Products Advisory Committee

National Center for Toxicological Research

Science Board to the National Center for Toxicological Research

Center for Drugs Evaluation and Research

Joint Meetings of the Anesthetic and Analgesic Drug Products Advisory Committee and the Drug Safety and Risk Management Advisory Committee

Annual Reports are available for public inspection between 9 a.m. and 4 p.m., Monday through Friday, at:

(1) The Library of Congress, Madison Building, Newspaper and Current Periodical Reading Room, 101 Independence Ave. SE., Rm. 133, Washington, DC; and

(2) Dockets Management Staff (HFA–305), Food and Drug Administration, 5630 Fishers Lane, Rm. 1061, Rockville, MD 20852.

Dated: June 27, 2017.

Anna K. Abram,

Deputy Commissioner for Policy, Planning, Legislation, and Analysis.

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