roundtables, and will dispose of it following the roundtables. We may use your e-mail address to contact you with information about the roundtable. The FTC Act and other laws the Commission administers permit the collection of this contact information to consider and use for the above purposes. Under the Freedom of Information Act or other laws, we may be required to disclose the information you provide to outside organizations. For additional information, including routine uses permitted by the Privacy Act, *see* the Commission's privacy policy at *http://www.ftc.gov/ftc/privacy.shtm.*

B. Requests To Participate as a Panelist

The format will consist of a roundtable with participation by panelists selected by FTC staff. FTC staff will identify and invite persons with relevant expertise to participate in the roundtables. In addition, the FTC staff may invite other persons to participate who submit requests in response to the **Federal Register** notice. Persons seeking to participate as panelists in the roundtables must notify the FTC in writing of their interest in participating on or before March 28, 2011. Requests to participate filed in an electronic form should be submitted by e-mail to: *MotorVehicleRoundtables1@ftc.gov.* Emails should be captioned "Motor Vehicle Roundtables—Request to Participate, Project No. P104811."

A request to participate as a panelist filed in paper form should also include the reference "Motor Vehicle Roundtables, Project No. P104811" both in the text of the comment and on the envelope, and should be mailed or delivered to the following address: Federal Trade Commission, Office of the Secretary, Room H–113 (Annex V), 600 Pennsylvania Avenue, NW., Washington, DC 20580. The FTC is requesting that requests to participate filed in paper form be sent by courier or overnight service, if possible, because U.S. postal mail in the Washington, DC area and at the Commission is subject to delay due to heightened security precautions.

C. Comments

Interested parties are invited to submit written comments electronically or in paper form on the topics to be discussed at the roundtable. Submission of comments should be captioned "Motor Vehicle Roundtables—Comment, Project No. P104811." Please note that your comment—including your name and your state—will be placed on the public record of this proceeding, including on the publicly accessible FTC Web site, at *http://www.ftc.gov/os/publiccomments.shtm.*

Because comments will be made public, they should not include any sensitive personal information, such as any individual's Social Security number; date of birth; driver's license number or other state identification number, or foreign country equivalent; passport number; financial account number; or credit or debit card number. Comments also should not include any sensitive health information, such as medical records or other individually identifiable health information. In addition, comments should not include any "[t]rade secrets and commercial or financial information obtained from a person and privileged or confidential," as provided in Section 6(f) of the FTC Act, 15 U.S.C. 46(f), and Commission Rule 4.10(a)(2), 16 CFR 4.10(a)(2). Comments containing material for which confidential treatment is requested must be filed in paper form, must be clearly labeled "Confidential," and must comply with FTC Rule 4.9(c), 16 CFR 4.9(c).[20]

Because paper mail addressed to the FTC is subject to delay due to heightened security screening, please consider submitting your comments in electronic form. Comments filed in electronic form should be submitted at *https://ftcpublic.commentworks.com/ftc/motorvehicleroundtables1* following the instructions on the web-based form. If this Notice appears at *http://www.regulations.gov/search/index.jsp,* you may also file an electronic comment through that website. The Commission will consider all comments forwarded to it by regulations.gov. You may also visit the FTC Web site at *http://www.ftc.gov* to read the Notice and the news release describing it.

A comment filed in paper form should include the reference "Motor Vehicle Roundtables, Project No. 104811" both in the text of the comment and on the envelope, and should be mailed or delivered to the following address: Federal Trade Commission, Office of the Secretary, Room H–113 (Annex V), 600 Pennsylvania Avenue, NW., Washington, DC 20580. The FTC is requesting that comments filed in paper form be sent by courier or overnight service, if possible, because U.S. postal mail in the Washington, DC area and at the Commission is subject to delay due to heightened security precautions.

The FTC Act and other laws the Commission administers permit the collection of public comments to consider and use in this proceeding as appropriate. The Commission will consider all timely and responsive public comments it receives, whether filed in paper or electronic form. Comments received will be available to the public on the FTC website, to the extent practicable, at *http://www.ftc.gov/os/publics.htm.* As a matter of discretion, the Commission makes every effort to remove home contact information of individuals before their comments are placed on the FTC Web site. More information, including routine uses permitted by the Privacy Act, may be found in the FTC's privacy policy, at *http://www.ftc.gov/ftc/privacy.shtm.*

By direction of the Commission.

Donald S. Clark,

Secretary.

[FR Doc. 2011–5873 Filed 3–14–11; 8:45 am]

BILLING CODE 6750–01–P

[20] The comment must be accompanied by an explicit request for confidential treatment, including the factual and legal basis for the request, and must identify the specific portions of the comment to be withheld from the public record. The request will be granted or denied by the Commission's General Counsel, consistent with applicable law and the public interest. *See* 16 CFR 4.9(c).

DEPARTMENT OF HEALTH AND HUMAN SERVICES

Call for Comments on the Draft Report of the Adult Immunization Working Group to the National Vaccine Advisory Committee on Adult Immunization: Complex Challenges and Recommendations for Improvement; Correction

AGENCY: Department of Health and Human Services, Office of the Secretary, Office of the Assistant Secretary for Health, National Vaccine Program Office.

ACTION: Notice: correction.

SUMMARY: The Department of Health and Human Services published a notice in the **Federal Register** of March 4, 2011, announcing a call for comment on the draft report of the Adult Immunization Working Group to the National Vaccine Advisory Committee. It was announced that the draft report and recommendations could be found on the Web at *http://www.hhs.gov/nvpo/nvac/subgroups/adultimmunization.* The Web address where the draft report and recommendations can be found is *http://www.hhs.gov/nvpo/nvac/subgroups/adultimmunization.html.*

FOR FURTHER INFORMATION CONTACT: Ms. Lauren Wu, e-mail: *lauren.wu@hhs.gov,* phone: 202–690–1191.

Correction

In the **Federal Register** of March 4, 2011, Vol. 76, No. 43, on page 12118, in the first column, correct the **ADDRESSES** caption to read:

(1) The draft report and recommendations are available on the Web at *http://www.hhs.gov/nvpo/nvac/subgroups/adultimmunization.html.*

Dated: March 9, 2011.

Bruce Gellin,

Director, National Vaccine Program Office.

[FR Doc. 2011–5851 Filed 3–14–11; 8:45 am]

BILLING CODE 4150–44–P

DEPARTMENT OF HEALTH AND HUMAN SERVICES

Centers for Disease Control and Prevention

[60Day-11–11DE]

Proposed Data Collections Submitted for Public Comment and Recommendations

In compliance with the requirement of Section 3506(c)(2)(A) of the Paperwork Reduction Act of 1995 for opportunity for public comment on proposed data collection projects, the Centers for Disease Control and Prevention (CDC) will publish periodic summaries of proposed projects. To request more information on the proposed projects or to obtain a copy of the data collection plans and instruments, call 404–639–5960 and send comments to Carol E. Walker, CDC Acting Reports Clearance Officer, 1600 Clifton Road, MS–D74, Atlanta, GA 30333 or send an email to *omb@cdc.gov*.

Comments are invited on: (1) Whether the proposed collection of information is necessary for the proper performance of the functions of the agency, including whether the information will have practical utility; (2) the accuracy of the agency's estimate of the burden of the proposed collection of information, including the validity of the methodology and assumptions used; (3) the quality, utility, and clarity of the information to be collected; and (4) the burden of the collection of information on those who are to respond, including the use of appropriate automated, electronic, mechanical, or other technological collection techniques or other forms of information technology. Written comments should be received within 60 days of this notice.

Proposed Collection

Communication Research on Folic Acid to Support the Division of Birth Defects and Developmental Disabilities—New—National Center on Birth Defects and Developmental Disabilities (NCBDDD), Centers for Disease Control and Prevention (CDC).

Background and Brief Description

Since mandatory folic acid fortification of cereal grain products was mandated in 1998, rates of folic acid-preventable neural tube defects (NTDs) have declined. Disparities in rates remain, however, with NTD prevalence being highest among Hispanic women of childbearing age. Efforts to increase consumption of vitamin supplements containing folic acid among women in this ethnic group have been ongoing, however, due to differences in diet, many of these women have not benefitted from food fortification to the extent that other race/ethnic groups have. A performance goal for NCBDDD focuses specifically on the reduction of these disparities: Reduce health disparities in the occurrence of folic acid-preventable spina bifida and anencephaly by reducing the birth prevalence of these conditions. Moreover, Healthy People 2010 objectives refer to the reduction of NTD rates and increase of folic acid consumption for all women of childbearing age: (1) Reduce the occurrence of spina bifida and other NTDs; (2) Increase the proportion of pregnancies begun with an optimum folic acid level by increasing the consumption of at least 400 mcg of folic acid each day from fortified foods or dietary supplements by nonpregnant women aged 15 to 44 and increasing the median red blood cell folate level among nonpregnant women aged 15 to 44 years. The 2009 congressional omnibus appropriations language includes reference to reducing health disparities: "There is significant concern about disparity in the rates of folic acid intake and neural tube defects, particularly in the Hispanic population. Within the funds provided for folic acid, CDC is encouraged to provide increased funding to expand the folic acid education campaign to inform more women and healthcare providers about the benefits of folic acid * * *". Finally, CDC partners are working to develop a food additive petition that will be submitted for approval to the FDA. This petition would allow for the addition of folic acid to corn masa flour and corn masa flour products. Knowing the consumer attitudes toward this endeavor is important to the overall success of the effort. Although up to 70% of neural tube defects can be prevented if a woman consumes folic acid before and during the first weeks of pregnancy, many women are still unaware of folic acid until they are already pregnant. Because half of all pregnancies in the U.S. are unplanned, reaching women with the folic acid message prior to pregnancy is critical. NCBDDD currently has several folic acid educational brochures, tip sheets, and booklets available in both English and Spanish. Since 2000, over 12 million folic acid materials have been distributed. Providing our partners, health care providers, and the public with evidence-based information in a format that is easy to read and visually appealing is important to the mission of the Prevention Research team. We want to ensure that the materials we currently have available still meet the needs of the intended audience.

CDC, with contract support from Battelle Centers for Public Health Research and Evaluation, is conducting research to inform efforts to promote folic acid consumptions among women of child-bearing age through two closely-related data collection efforts: (1) Exploratory Research of Hispanic Women's Reactions to and Beliefs About Folic Acid Fortification of Corn Masa Flour, and (2) Exploratory Research of Childbearing Age Women's Folic Acid Awareness and Knowledge, and their Reactions to Existing CDC Folic Acid Educational Materials. The purpose of the first proposed primary data collection effort is to better understand consumer acceptance of fortifying corn masa flour, a staple product in many traditional Latino, and in particular Mexican, foods. The purpose of the second proposed primary data collection effort is to determine whether educational materials developed over 10 years ago to promote folic acid consumption continue to be appealing and resonate with the target audience today. To address these two purposes and support the folic acid education efforts of CDC, focus groups with the target audience are needed.

For the first data collection activity phase, participants will be English and Spanish-speaking women 18–44 years who self identify as Mexican or Mexican American, or Central American. Participants will be segmented into groups based on whether they consume corn masa flour less than 4 times per day or 4 or more times per day. The contractor will conduct sixteen (16) focus groups with five (5) participants in each focus group. It is estimated that 320 respondents will have to be screened in order to recruit 80 focus group participants. Each screening will take approximately 6 minutes. The estimated response burden for the screening process is 32 hours. The focus group session will be structured to