MITOR requires the merchant to make a refund and the consumer has paid by credit card, the Rule also requires the merchant to notify the consumer either that any charge to the consumer's charge account will be reversed or that the merchant will take no action that will result in a charge.

On January 19, 2016, the Commission sought comment on the information collection requirements in MTOR. See 81 FR 2860. The Commission received two comments but neither one addressed the issues raised by the public comment request. As required by OMB regulations, 5 CFR part 1320, the FTC is providing this second opportunity for public comment.

Likely Respondents: Businesses engaged in the sale of merchandise by mail or by telephone.

Estimated Annual Hours Burden: 1,953,840 hours.

Third Party Disclosure: [(33,267 established businesses × 50 hours) + (1,263 new entrants × 230 hours) = 1,953,840 hours.

Estimated Annual Cost Burden: $44,879,705, which is derived from 1,953,840 hours × $22.97/hour.[1]

Request for Comment

You can file a comment online or on paper. For the Commission to consider your comment, we must receive it on or before May 12, 2016. Write ''Mail, Internet, or Telephone Order Merchandise Trade Regulation Rule: FTC File No. R511929'' on your comment. Your comment—including your name and your state—will be placed on the public record of this proceeding, including, to the extent practicable, on the public Commission Web site, at *http://www.ftc.gov/os/publiccomments.shtm.* As a matter of discretion, the Commission tries to remove individuals' home contact information from comments before placing them on the Commission Web site.

Because your comment will be made public, you are solely responsible for making sure that your comment does not include any sensitive personal information, such as anyone'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. You are also solely responsible for making sure that your comment does not include any sensitive health information, like medical records or other individually identifiable health information. In addition, do not include any ''[t]rade secret or any commercial or financial information which is . . . privileged or confidential,'' as discussed in Section 6(f) of the FTC Act, 15 U.S.C. 46(f), and FTC Rule 4.10(a)(2), 16 CFR 4.10(a)(2). In particular, do not include competitively sensitive information such as costs, sales statistics, inventories, formulas, patterns, devices, manufacturing processes, or customer names.

If you want the Commission to give your comment confidential treatment, you must file it in paper form, with a request for confidential treatment, and you are required to follow the procedure explained in FTC Rule 4.9(c), 16 CFR 4.9(c). Your comment will be kept confidential only if the FTC General Counsel grants your request in accordance with the law and the public interest.

Postal mail addressed to the Commission is subject to delay due to heightened security screening. As a result, we encourage you to submit your comment online, or to send it to the Commission by courier or overnight service. To make sure that the Commission considers your online comment, you must file it at *https://ftcpublic.commentworks.com/ftc/mitorpra2,* by following the instructions on the Web-based form. If this Notice appears at *http://www.regulations.gov,* you also may file a comment through that Web site.

If you file your comment on paper, write ''Mail, Internet, or Telephone Order Merchandise Trade Regulation Rule: FTC File No. R511929'' on your comment and on the envelope, and mail or deliver it to the following address: Federal Trade Commission, Office of the Secretary, 600 Pennsylvania Avenue NW., Suite CC–5610 (Annex J), Washington, DC 20580, or deliver your comment to the following address: Federal Trade Commission, Office of the Secretary, Constitution Center, 400 7th Street SW., 5th Floor, Suite 5610 (Annex J), Washington, DC 20024. If possible, submit your paper comment to the Commission by courier or overnight service.

The FTC Act and other laws that 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 that it receives on or before May 12, 2016. You can find more information, including routine uses permitted by the Privacy Act, in the Commission's privacy policy, at *http://www.ftc.gov/ftc/privacy.shtm.*

Comments on the information collection requirements subject to review under the PRA should also be submitted to OMB. If sent by U.S. mail, address comments to: Office of Information and Regulatory Affairs, Office of Management and Budget, Attention: Desk Officer for the Federal Trade Commission, New Executive Office Building, Docket Library, Room 10102, 725 17th Street NW., Washington, DC 20503. Comments sent to OMB by U.S. postal mail, however, are subject to delays due to heightened security precautions. Thus, comments instead should be sent by facsimile to (202) 395–5167.

Christian S. White,

Deputy General Counsel.

[FR Doc. 2016–08369 Filed 4–11–16; 8:45 am]

BILLING CODE 6750–01–P

[1] This is the mean hourly income for workers in sales and related occupations according to the latest figures from the Department of Commerce's Bureau of Labor Statistics. *See* Table 1, National employment and wage data from the Occupational Employment Statistics survey by occupation, May 2015, at *http://www.bls.gov/news.release/ocwage.t01.htm.*

DEPARTMENT OF HEALTH AND HUMAN SERVICES

Centers for Disease Control and Prevention

Emergency Funding for Puerto Rico Department of Health, Zika Virus Outbreak

AGENCY: Centers for Disease Control and Prevention (CDC), Department of Health and Human Services (DHHS).

ACTION: This notice announces the Centers for Disease Control and Prevention's (CDC) intent to fund the Puerto Rico Department of Health with Prevention and Public Health Funds (PPHF).

PPHF 2016: Epidemiology and Laboratory Capacity Program—Emergency Funding for Puerto Rico Department of Health, Zika virus Outbreak for Infectious Diseases (ELC)—financed solely by Prevention and Public Health Funds.

FOA Number: CDC–RFA–CK14–140103CONTPPHF2016.

SUMMARY: The U.S. Centers for Disease Control and Prevention (CDC) is providing $3,700,000 in urgent funding through the Epidemiology and Laboratory Capacity for Infectious Diseases (ELC) Cooperative Agreement to the Puerto Rico Department of Health (PRDOH) to combat the current outbreak of Zika virus.

Project Description

Puerto Rico is experiencing an approximate doubling of confirmed Zika

virus cases per week—they are unique in the total number of cases, the level of local transmission, and the presence of the Zika-carrying vector. Currently, the PR DOH cannot sufficiently address necessary aspects of the outbreak response without additional support. In addition to equipment and supplies necessary for the increased testing for Zika virus, funds awarded to PRDOH will be used to support additional epidemiology and laboratory staff critical to the response efforts.

Prevention Fund Reporting Requirements: This award requires the grantee to complete projects or activities which are funded under the Prevention and Public Health Fund (PPHF) (Section 4002 of Pub. L. 111–148) and to report on use of PPHF funds provided through this award. Information from these reports will be made available to the public.

Grantees awarded a grant, cooperative agreement, or contract from such funds with a value of $25,000 or more shall produce reports on a semi-annual basis with a reporting cycle of January 1–June 30 and July 1–December 31; and email such reports to the CDC Web site (template and point of contact to be provided after award) no later than 20 calendar days after the end of each reporting period (*i.e.* July 20 and January 20, respectively). Grantee reports must reference the NoA number and title of the grant, and include a summary of the activities undertaken and identify any sub-awards (including the purpose of the award and the identity of each sub-recipient).

Responsibilities for Informing Sub-recipients: Grantees agree to separately identify each sub-recipient, document the execution date sub-award, date(s) of the disbursement of funds, the Federal award number, any special CFDA number assigned for PPHF fund purposes, and the amount of PPHF funds. When a grantee awards PPHF funds for an existing program, the information furnished to sub-recipients shall distinguish the sub-awards of incremental PPHF funds from regular sub-awards under the existing program.

DATES: Effective date is April 12, 2016.

ADDRESSES: Alvin Shultz, MSPH, Division of Preparedness and Emerging Infectious, National Center for Emerging and Zoonotic Infectious Diseases, Centers for Disease Control and Prevention, 1600 Clifton Road NE., Atlanta, GA 30333, Phone: 404–639–7028, E-Mail: *Ashultz@cdc.gov*.

FOR FURTHER INFORMATION CONTACT: Alvin Shultz, MSPH, Division of Preparedness and Emerging Infectious, National Center for Emerging and Zoonotic Infectious, Diseases Centers for Disease Control and Prevention, 1600 Clifton Road NE., Atlanta, GA 30333. Phone: 404–639–7028. E-Mail: *Ashultz@cdc.gov.*

Dated: March 25, 2016.

Terrance Perry,

Director, Office of Grants Services, Centers for Disease Control and Prevention.

[FR Doc. 2016–08318 Filed 4–11–16; 8:45 am]

BILLING CODE 4163–18–P

DEPARTMENT OF HEALTH AND HUMAN SERVICES

Centers for Disease Control and Prevention

[30 Day–16–0217]

Agency Forms Undergoing Paperwork Reduction Act Review

The Centers for Disease Control and Prevention (CDC) has submitted the following information collection request to the Office of Management and Budget (OMB) for review and approval in accordance with the Paperwork Reduction Act of 1995. The notice for the proposed information collection is published to obtain comments from the public and affected agencies.

Written comments and suggestions from the public and affected agencies concerning the proposed collection of information are encouraged. Your comments should address any of the following: (a) Evaluate 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; (b) Evaluate the accuracy of the agencies estimate of the burden of the proposed collection of information, including the validity of the methodology and assumptions used; (c) Enhance the quality, utility, and clarity of the information to be collected; (d) Minimize the burden of the collection of information on those who are to respond, including through the use of appropriate automated, electronic, mechanical, or other technological collection techniques or other forms of information technology, *e.g.,* permitting electronic submission of responses; and (e) Assess information collection costs.

To request additional information on the proposed project or to obtain a copy of the information collection plan and instruments, call (404) 639–7570 or send an email to *omb@cdc.gov.* Direct written comments and/or suggestions regarding the items contained in this notice to the Attention: CDC Desk Officer, Office of Management and Budget, Washington, DC 20503 or by fax to (202) 395–5806. Written comments should be received within 30 days of this notice.

Proposed Project

NCHS Vital Statistics Training Application (OMB Control No. 0920–0217, exp. 5/31/2016)—Revision—National Center for Health Statistics NCHS), Centers for Disease Control and Prevention (CDC).

Background and Brief Description

In the United States, legal authority for the registration of vital events, *i.e.,* births, deaths, marriages, divorces, fetal deaths, and induced terminations of pregnancy, resides individually with the States (as well as cities in the case of New York City and Washington, DC) and Puerto Rico, the Virgin Islands, Guam, American Samoa, and the Commonwealth of the Northern Mariana Islands. These governmental entities are the full legal proprietors of vital records and the information contained therein. As a result of this State authority, the collection of registration-based vital statistics at the national level, referred to as the U.S. National Vital Statistics System (NVSS), depends on a cooperative relationship between the States and the Federal government. This data collection, authorized by 42 U.S.C. 242k, has been carried out by NCHS since it was created in 1960.

NCHS assists in achieving the comparability needed for combining data from all States into national statistics, by conducting a training program for State and local vital statistics staff to assist in developing expertise in all aspects of vital registration and vital statistics. The training offered under this program includes courses for registration staff, statisticians, and coding specialists, all designed to bring about a high degree of uniformity and quality in the data provided by the States. This training program is authorized by 42 U.S.C. 242b, section 304(a).

NCHS notifies State and local vital registration officials, as well as Canadian counterparts, about upcoming training. Individual candidates for training then submit an application form including name, address, occupation, and other relevant information.

In this revision, the application for the Vital Statistics Training is being updated to capture additional logistical information. The proposed changes include the addition of two questions (1) to identify the training personnel as either State or locally-based and (2) to