the final data collection and 58 hours for the 100 who participate in the pre-test.

C. Estimated Costs

The cost per respondent should be negligible. Participation is voluntary, and will not require any labor expenditures by respondents. There are no capital, start up, operation, maintenance, or other similar costs to the respondents.

D. 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 September 3, 2013. Write ''Fraud Susceptibility Internet Panel Study, FTC File No. P095500'' 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, like 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 obtained from any person and which is privileged or confidential,'' as provided 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 have to follow the procedure explained in FTC Rule 4.9(c), 16 CFR 4.9(c).[3] Your comment will be kept confidential only if the FTC General Counsel, in his or her sole discretion, 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 comments online. To make sure that the Commission considers your online comment, you must file it at *https://public.commentworks.com/ftc/fraudinternetpanelstudypra2,* by following the instructions on the web-based form. If this Notice appears at *http://www.regulations.gov/#!home,* you also may file a comment through that Web site.

If you file your comment on paper, write ''Fraud Susceptibility Internet Panel Study, FTC File No. P095500'' on your comment and on the envelope, and mail or deliver it to the following address: Federal Trade Commission, Office of the Secretary, Room H–113 (Annex J), 600 Pennsylvania Avenue NW., Washington, DC 20580. If possible, submit your paper comment to the Commission by courier or overnight service.

Visit the Commission Web site at *http://www.ftc.gov* to read this Notice and the news release describing it. 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 September 3, 2013. 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.htm.*

The FTC invites comments 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) ways to enhance the quality, utility, and clarity of the information to be collected; and (4) ways to minimize the burden of the collection of information.

Comments on the information collection requirements subject to review under the PRA should additionally be submitted to OMB. If sent by U.S. mail, they should be addressed 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.

[3] In particular, the written request for confidential treatment that accompanies the comment must include the factual and legal basis for the request, and must identify the specific portions of the comment to be withheld from the public record. *See* FTC Rule 4.9(c), 16 CFR 4.9(c).

David C. Shonka,

Principal Deputy General Counsel.

[FR Doc. 2013–18560 Filed 7–31–13; 8:45 am]

BILLING CODE 6750–01–P

DEPARTMENT OF HEALTH AND HUMAN SERVICES

Office of the Secretary

[Document Identifier HHS–OS–20165–60D]

Agency Information Collection Activities; Proposed Collection; Public Comment Request

AGENCY: Office of the Secretary, HHS.

ACTION: Notice.

SUMMARY: In compliance with section 3506(c)(2)(A) of the Paperwork Reduction Act of 1995, the Office of the Secretary (OS), Department of Health and Human Services, announces plans to submit an Information Collection Request (ICR), described below, to the Office of Management and Budget (OMB). The ICR is for revision of a previously-approved information collection assigned OMB control number 0937–0025, which expired on 08/31/2013. Prior to submitting that ICR to OMB, OS seeks comments from the public regarding the burden estimate below or any other aspect of the ICR.

DATES: Comments on the ICR must be received on or before September 30, 2013.

ADDRESSES: Submit your comments to *Information.CollectionClearance@hhs.gov* or by calling (202) 690–6162.

FOR FURTHER INFORMATION CONTACT: Information Collection Clearance staff, *Information.CollectionClearance@hhs.gov* or (202) 690–6162.

SUPPLEMENTARY INFORMATION: When submitting comments or requesting information, please include the document identifier HHS–OS–20165–60D for reference. Information Collection Request Title: Application for

Appointment as a Commissioned Officer in the PHS Commission Corps.

Abstract: The information collected will include personal information such as name, social security number, and date of birth. Other information will be responses to various questions regarding an applicants' qualifications to join the Commissioned Corps of the U.S. Public Health Service.

Need and Proposed Use of the Information: The Commissioned Corps of the U.S. Public Health Service has a need for the information in order to assess the qualifications of each applicant and make a determination whether the applicant meets the requirements to receive a commission. The information is used to make determinations on candidates/applicants seeking appointment to the Corps to assess their whether they are suitable for life in the uniformed services based upon a review of a variety of assessment factors including, but not limited to: Personal adjustment, employment history, character, suitability investigation clearance, and a candidate's prior history of service in one of the uniformed services. Their potential for leadership as a commissioned officer and their ability to deal effectively with people is evaluated.

Likely Respondents: Respondents would be applicants/candidates for a commission in the Commissioned Corps of the United States Public Health Service.

Burden Statement: The time expended by persons to generate, maintain, retain, disclose or provide the information requested. This includes the time needed to review instructions, to develop, acquire, install and utilize technology and systems for the purpose of collecting, validating and verifying information, processing and maintaining information, and disclosing and providing information, to train personnel and to be able to respond to a collection of information, to search data sources, to complete and review the collection of information, and to transmit or otherwise disclose the information. The total annual burden hours estimated for this ICR are summarized in the table below.

TOTAL ESTIMATED ANNUALIZED BURDEN—HOURS

Form name	Number of respondents	Number of responses per respondent	Average burden per response (in hours)	Total burden hours
Prequalification Review	8,000	1	15/60	2,000
PHS–50	1,000	1	1.0	1,000
PHS–1813	4,000	1	15/60	1,000
Addendum: Commissioned Corps Personal Statement	1,000	1	45/60	750
Total				4,750

The Office of the Secretary (OS), Department of Health and Human Services specifically requests comments on (1) the necessity and utility of the proposed information collection for the proper performance of the agency's functions, (2) the accuracy of the estimated burden, (3) ways to enhance the quality, utility, and clarity of the information to be collected, and (4) the use of automated collection techniques or other forms of information technology to minimize the information collection burden.

Darius Taylor,

Deputy Information Collection Clearance Officer.

[FR Doc. 2013–18459 Filed 7–31–13; 8:45 am]

BILLING CODE 4150–49–P

DEPARTMENT OF HEALTH AND HUMAN SERVICES

Solicitation of Written Comments on the Global Immunizations Working Group's Draft Report and Draft Recommendations for Enhancing the Work of the HHS National Vaccine Program in Global Immunizations for Consideration by the National Vaccine Advisory Committee

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

ACTION: Notice.

SUMMARY: The National Vaccine Advisory Committee (NVAC) was established in 1987 to comply with Title XXI of the Public Health Service Act (Pub. L. 99–660) (§ 2105) (42 U.S. Code 300aa–5 (PDF–78 KB)). Its purpose is to advise and make recommendations to the Director of the National Vaccine Program on matters related to program responsibilities. The Assistant Secretary for Health (ASH) has been designated by the Secretary of Health and Human Services (HHS) as the Director of the National Vaccine Program.

The ASH charged the NVAC with reviewing the role of HHS in global vaccination, the effects of global vaccination on global populations, the effects of global vaccination on U.S. populations, and recommending how HHS can best continue to contribute, consistent with its newly established Global Health Strategy and Goal 5 of the National Vaccine Plan. The NVAC was also asked to make recommendations on how to best communicate this information to decision makers and the general public to ensure continued sufficient resources for the global vaccination efforts. The NVAC established the Global Immunizations Working Group to assist in addressing these charges.

A draft report and draft recommendations have been developed by the working group for consideration by the NVAC and will be deliberated on by the NVAC when developing NVAC's final recommendations to the ASH. The National Vaccine Program Office (NVPO) is soliciting public comment on the draft report and draft recommendations from a variety of stakeholders, including the general public, for consideration by the NVAC as they develop their final recommendations to the ASH. It is anticipated that the draft report and draft recommendations, as revised with consideration given to public comment