SUPPLEMENTARY INFORMATION: GSA published FTR Amendment 2008–04 in the **Federal Register** at 73 FR 35952 on June 25, 2008, specifying that GSA would no longer publish the RIT Allowance tables in Title 41 of the Code of Federal Regulations, Part 302–17, Appendices A through D (FTR prior to January 1, 2015—*www.gsa.gov/federaltravelregulation*—FTR and Related Files); instead, the tables would be available on a GSA Web site. FTR Bulletin 17–03: Relocation Allowances—Relocation Income Tax (RIT) Allowance Tables is now available and provides the annual changes to the RIT allowance tables necessary for calculating the amount of a transferee's increased tax burden due to his or her official permanent change of station. GSA published FTR Amendment 2014–01 in the **Federal Register** on August 21, 2014, (79 FR 49640), which eliminated the need for the Government-unique tax tables for relocations that began on January 1, 2015 and later. However, for relocations that began earlier than January 1, 2015, this bulletin is required to compute the employee's reimbursement for additional income taxes associated with the relocation. For relocations that began on or after January 1, 2015, transferees and agencies must use the tables published by the U.S. Internal Revenue Service (IRS), state, and local tax authorities, and follow the procedures in the FTR, Part 302–17. FTR Bulletin 17–03 and all other FTR Bulletins can be found at *www.gsa.gov/ftrbulletin.*

Giancarlo Brizzi,

Acting Associate Administrator, Office of Government-wide Policy, General Services Administration.

[FR Doc. 2017–09236 Filed 5–5–17; 8:45 am]

BILLING CODE 6820–14–P

DEPARTMENT OF HEALTH AND HUMAN SERVICES

Centers for Disease Control and Prevention Healthcare

Infection Control Practices Advisory Committee (HICPAC)

Correction: This notice was published in the **Federal Register** on April 14, 2017, Volume 82, Number 71, pages 17996–17997. The Status should read as follows:

Open to the public limited only by the availability of 200 telephone ports. To register for this call, please go to *www.cdc.gov/hicpac.* Time will be available for public comment.

Contact Person for More Information: Erin Stone, M.A., HICPAC, Division of Healthcare Quality Promotion, NCEZID, CDC, 1600 Clifton Road NE., Mailstop A–31, Atlanta, Georgia 30333; Email: *HICPAC@cdc.gov*.

The Director, Management Analysis and Services Office, has been delegated the authority to sign Federal Register notices pertaining to announcements of meetings and other committee management activities, for both the Centers for Disease Control and Prevention and the Agency for Toxic Substances and Disease Registry.

Elaine L. Baker,

Director, Management Analysis and Services Office, Centers for Disease Control and Prevention.

[FR Doc. 2017–09206 Filed 5–5–17; 8:45 am]

BILLING CODE 4163–18–P

DEPARTMENT OF HEALTH AND HUMAN SERVICES

Centers for Disease Control and Prevention

Board of Scientific Counselors, Office of Infectious Diseases (BSC, OID)

Correction: This notice was published in the **Federal Register** on April 12, 2017, Volume 82, Number 69, pages 17666–17667. The Status should read as follows:

Status: Open to the public limited only by the space and telephone ports available (The meeting room will accommodate up to 100 people and the telephone ports will accommodate up to 50 people). The toll-free dial-in number is 1–888–373–3590 with a pass code of 541544.

Contact Person For More Information: Robin Moseley, M.A.T., Designated Federal Officer, OID, CDC, 1600 Clifton Road NE., Mailstop D10, Atlanta, Georgia 30329, Telephone: (404) 639–4461.

The Director, Management Analysis and Services Office, has been delegated the authority to sign **Federal Register** notices pertaining to announcements of meetings and other committee management activities, for both the Centers for Disease Control and Prevention and the Agency for Toxic Substances and Disease Registry.

Elaine L. Baker,

Director, Management Analysis and Services Office, Centers for Disease Control and Prevention.

[FR Doc. 2017–09207 Filed 5–5–17; 8:45 am]

BILLING CODE 4163–18–P

DEPARTMENT OF HEALTH AND HUMAN SERVICES

Centers for Disease Control and Prevention

National Center for Chronic Disease Prevention and Health Promotion, Interagency Committee on Smoking and Health (ICSH)

In accordance with section 10(a) (2) of the Federal Advisory Committee Act (Pub. L. 92–463), the Centers for Disease Control and Prevention (CDC) announces the following meeting of the aforementioned committee.

Time and Date: 9:00 a.m.–4:00 p.m., EDT, May 31, 2017.

Place: U.S. Department of Health and Human Services, Hubert H. Humphrey Building, the Great Hall, located at 200 Independence Avenue SW., Washington, DC 20201, Telephone: (202) 245–0552. This meeting is also accessible by teleconference.

Login information for teleconference is as follows:

Toll Free Phone#: (800) 593–8961.

Participant Passcode: 3435645.

Participants can join the visual portion only for this event directly at: *https://webconf.cdc.gov/zqe0/3d9qzwzb.*

If you are offered the option to join audio, please select ''don't join audio'' and use the Toll Free number listed above.

Status: Open to the public, limited only by the space and telephone lines available. Time will also be available for public comment. To register for this meeting please email the contact person below (see Contact Person for More Information). If you will require a sign language interpreter, or have other special needs, please notify the contact person by 4:30 p.m., EDT, on May 18, 2017.

Purpose: The Interagency Committee on Smoking and Health shall provide advice and guidance to the Secretary, Department of Health and Human Services (HHS), regarding: (a) Coordination of research, educational programs, and other activities within the Department that relate to the effect of smoking on human health and on coordination of these activities, with similar activities of other Federal and private agencies; and (b) establishment and maintenance of liaisons with appropriate private entities, other Federal agencies, and State and local public agencies, regarding activities relating to the effect of cigarette smoking on human health.

Matters for Discussion: The topic of the meeting is ''Increasing the Impact of Evidence-Based Tobacco Treatment'' and the objective of the meeting is to identify federal actions to increase the reach and effectiveness of efforts to help smokers quit.

Agenda items are subject to change as priorities dictate.

Contact Person for More Information: Monica L. Swann, Management and Program Analyst, National Center for Chronic Disease Prevention and Health Promotion, CDC, 395 E. Street SW., Washington, DC 20024, Telephone: (202) 245–0552; email: *mswann@cdc.gov.*

The Director, Management Analysis and Services Office has been delegated the authority to sign **Federal Register** notices pertaining to announcements of meetings and other committee management activities, for both the Centers for Disease Control and Prevention and the Agency for Toxic Substances and Disease Registry.

Elaine L. Baker,

Director, Management Analysis and Services Office, Centers for Disease Control and Prevention.

[FR Doc. 2017–09208 Filed 5–5–17; 8:45 am]

BILLING CODE 4163–18–P

DEPARTMENT OF HEALTH AND HUMAN SERVICES

Office of the Secretary

[Document Identifier: HHS–OS–0990–0379]

60-Day Notice Template for Extension of Generic Clearance for the Collection of Qualitative Feedback on Agency Service Delivery

AGENCY: U.S. Department of Health and Human Services (HHS).

ACTION: Notice and request for comments. Office of the National Coordinator for Health Information Technology is requesting OMB approval for an extension by OMB.

SUMMARY: Department of Health and Human Services, The Office of the Secretary (OS), as part of its continuing effort to reduce paperwork and respondent burden, invites the general public to take this opportunity to comment on the ''Generic Clearance for the Collection of Qualitative Feedback on Agency Service Delivery'' for approval under the Paperwork Reduction Act (PRA). This collection was developed as part of a Federal Government-wide effort to streamline the process for seeking feedback from the public on service delivery. This notice announces our intent to submit this collection to OMB for approval and solicits comments on specific aspects for the proposed information collection.

DATES: Comments on the ICR must be received on or before July 7, 2017.

ADDRESSES: Submit comments by one of the following methods:

- *Web site: www.regulations.gov.* Direct comments to Docket ID OMB–2010–0021.
- *Email: Information.CollectionClearance@hhs.gov.*
- *Phone:* (202) 795–7714.

Comments submitted in response to this notice may be made available to the public through relevant Web sites. For this reason, please do not include in your comments information of a confidential nature, such as sensitive personal information or proprietary information. If you send an email comment, your email address will be automatically captured and included as part of the comment that is placed in the public docket and made available on the Internet. Please note that responses to this public comment request containing any routine notice about the confidentiality of the communication will be treated as public comments that may be made available to the public notwithstanding the inclusion of the routine notice.

FOR FURTHER INFORMATION CONTACT: Sherrette Funn, *Sherrrette.funn@HHS.GOV* or (202) 795–7714.

SUPPLEMENTARY INFORMATION:

Title: Generic Clearance for the Collection of Qualitative Feedback on Agency Service Delivery.

Abstract: The proposed information collection activity provides a means to garner qualitative customer and stakeholder feedback in an efficient, timely manner, in accordance with the Administration's commitment to improving service delivery. By qualitative feedback we mean information that provides useful insights on perceptions and opinions, but are not statistical surveys that yield quantitative results that can be generalized to the population of study. This feedback will provide insights into customer or stakeholder perceptions, experiences and expectations, provide an early warning of issues with service, or focus attention on areas where communication, training or changes in operations might improve delivery of products or services. These collections will allow for ongoing, collaborative and actionable communications between the Agency and its customers and stakeholders. It will also allow feedback to contribute directly to the improvement of program management.

The solicitation of feedback will target areas such as: Timeliness, appropriateness, accuracy of information, courtesy, efficiency of service delivery, and resolution of issues with service delivery. Responses will be assessed to plan and inform efforts to improve or maintain the quality of service offered to the public. If this information is not collected, vital feedback from customers and stakeholders on the Agency's services will be unavailable.

The Agency will only submit a collection for approval under this generic clearance if it meets the following conditions:

- The collections are voluntary;
- The collections are low-burden for respondents (based on considerations of total burden hours, total number of respondents, or burden-hours per respondent) and are low-cost for both the respondents and the Federal Government;
- The collections are non-controversial and do not raise issues of concern to other Federal agencies;
- Any collection is targeted to the solicitation of opinions from respondents who have experience with the program or may have experience with the program in the near future;
- Personally identifiable information (PII) is collected only to the extent necessary and is not retained;
- Information gathered will be used only internally for general service improvement and program management purposes and is not intended for release outside of the agency;
- Information gathered will not be used for the purpose of substantially informing influential policy decisions; and
- Information gathered will yield qualitative information; the collections will not be designed or expected to yield statistically reliable results or used as though the results are generalizable to the population of study.

Feedback collected under this generic clearance provides useful information, but it does not yield data that can be generalized to the overall population. This type of generic clearance for qualitative information will not be used for quantitative information collections that are designed to yield reliably actionable results, such as monitoring trends over time or documenting program performance. Such data uses require more rigorous designs that address: The target population to which generalizations will be made, the sampling frame, the sample design (including stratification and clustering), the precision requirements or power calculations that justify the proposed sample size, the expected response rate, methods for assessing potential non-response bias, the protocols for data collection, and any testing procedures that were or will be undertaken prior to fielding the study. Depending on the degree of influence the results are likely to have, such collections may still be eligible for submission for other generic mechanisms that are designed to yield quantitative results.

As a general matter, information collections will not result in any new system of records containing privacy information and will not ask questions of a sensitive nature, such as sexual behavior and attitudes, religious beliefs,