information collection documents from the Regulatory Secretariat Division (MVCB), 1275 First Street NE., Washington, DC 20417, telephone (202) 501–4755. Please cite OMB Control No. 3090–0262, Identification of Products with Environmental Attributes, in all correspondence.

Dated: July 3, 2012.

Joseph A. Neurauter,

Director, Office of Acquisition Policy, Senior Procurement Executive.

[FR Doc. 2012–16898 Filed 7–10–12; 8:45 am]

BILLING CODE 6820–61–P

GENERAL SERVICES ADMINISTRATION

[Docket 2012–0001; Sequence 13; OMB Control NO. 3090–0283]

Office of the Chief Information Officer; Information Collection; Temporary Contractor Information Worksheet

AGENCY: Identity, Credential, and Access Management (ICAM) Division, Office of Enterprise Solutions (IA), Office of the Chief Information Officer (OCIO), General Services Administration (GSA).

ACTION: Notice of request for comments regarding an extension to an existing OMB clearance.

SUMMARY: Under the provisions of the Paperwork Reduction Act, the General Services Administration will be submitting to the Office of Management and Budget (OMB) a request to review and approve a previously approved information collection requirement regarding temporary contractor information worksheet.

GSA requires OMB approval for this collection to make determinations on granting unescorted physical access to GSA-controlled facilities. The approval is critical for GSA to continue to make physical access determinations for temporary contractors as a result of the American Recovery and Reinvestment Act of 2009 (Pub. L. 111–5).

Public comments are particularly invited on: Whether this collection of information is necessary and whether it will have practical utility; whether our estimate of the public burden of this collection of information is accurate, and based on valid assumptions and methodology; ways to enhance the quality, utility, and clarity of the information to be collected.

DATES: Submit comments on or before: September 10, 2012.

ADDRESSES: Submit comments identified by Information Collection 3090–0283, Temporary Contractor Information Sheet, by any of the following methods:

- *Regulations.gov: http://www.regulations.gov.* Submit comments via the Federal eRulemaking portal by searching the OMB control number. Select the link ''Submit a Comment'' that corresponds with ''Information Collection 3090–0283, Temporary Contractor Information Sheet''. Follow the instructions provided at the ''Submit a Comment'' screen. Please include your name, company name (if any), and ''Information Collection 3090–0283, Temporary Contractor Information Sheet'' on your attached document.
- *Fax:* 202–501–4067.
- *Mail:* General Services Administration, Regulatory Secretariat (MVCB), 1275 First Street, NE., Washington, DC 20417. Attn: Hada Flowers/IC 3090–0283, Temporary Contractor Information Sheet.

Instructions: Please submit comments only and cite Information Collection 3090–0283, Temporary Contractor Information Sheet, in all correspondence related to this collection. All comments received will be posted without change to *http://www.regulations.gov,* including any personal and/or business confidential information provided.

FOR FURTHER INFORMATION CONTACT: Mr. Phil Ahn, Director, OCIO Identity Credential and Access Management Division, GSA, telephone (202) 501–2447 or via email at *phil.ahn@gsa.gov.*

SUPPLEMENTARY INFORMATION:

A. Purpose

The U.S. Government conducts criminal checks to establish that applicants or incumbents working for the Government under contract may have unescorted access to GSA-controlled facilities. GSA uses the Temporary Contractor Information Worksheet and the FBI Form FD–258 Fingerprint Card to conduct a FBI National Criminal Information Check (NCIC) for each temporary contractor (working on contract for six months or less and require physical access only) on GSA contracts for American Recovery and Reinvestment Act of 2009 (Pub. L. 111–5) efforts to determine whether to grant unescorted access to GSA-controlled facilities. GSA will continue to make physical access determinations for temporary contractors due to the American Recovery and Reinvestment Act of 2009.

The Office of Management and Budget (OMB) Guidance M–05–24 for Homeland Security Presidential Directive (HSPD) 12 authorizes Federal departments and agencies to ensure that temporary contractors have limited/controlled access to facilities and information systems. GSA Directive CIO P 2181.1 Homeland Security Presidential Directive-12 Personal Identity Verification and Credentialing (available at *http://www.gsa.gov/hspd12*) states that GSA temporary contractors must undergo a minimum of a FBI National Criminal Information Check (NCIC) to receive unescorted physical access. Temporary contractors' Social Security Number is needed to keep records accurate, because other people may have the same name and birth date. Executive Order 9397 Numbering System for Federal Accounts Relating to Individual Persons also allows Federal agencies to use this number to help identify individuals in agency records. GSA describes how information will be maintained in the Privacy Act system of record notice published in the **Federal Register** at 73 FR 35690 on June 24, 2008.

B. Annual Reporting Burden

Respondents: 1,250.
Responses per Respondent: 1.
Hours per Response: .25.
Total Burden Hours: 313.
Obtaining Copies of Proposals: Requesters may obtain a copy of the information collection documents from the General Services Administration, Regulatory Secretariat Division (MVCB), 1275 First Street, NE., Washington, DC 20417, telephone (202) 501–4755. Please cite OMB Control No. 3090–0283, Temporary Contractor Information Worksheet in all correspondence. The form can be downloaded from the GSA Forms Library at *http://www.gsa.gov/forms.* Type GSA 850 in the form search field.

Dated: June 28, 2012.

Casey Coleman,

Chief Information Officer.

[FR Doc. 2012–16899 Filed 7–10–12; 8:45 am]

BILLING CODE 6820–34–P

DEPARTMENT OF HEALTH AND HUMAN SERVICES

Meeting of the Presidential Advisory Council on HIV/AIDS

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

ACTION: Notice.

SUMMARY: As stipulated by the Federal Advisory Committee Act, the U.S. Department of Health and Human Service (DHHS) is hereby giving notice that the Presidential Advisory Council

on HIV/AIDS (PACHA) will hold a meeting. The meeting will be open to the public.

DATES: The meeting will be held July 25, 2012 from 3:00 p.m. to approximately 5:00 p.m. (EDT).

ADDRESSES: 801 K Street NW., Washington, DC, 20001.

FOR FURTHER INFORMATION CONTACT: Mr. Melvin Joppy, Committee Manager, Presidential Advisory Council on HIV/AIDS, Department of Health and Human Services, 200 Independence Avenue SW., Room 443H, Hubert H. Humphrey Building, Washington, DC 20201; (202) 690–5560. More detailed information about PACHA can be obtained by accessing the Council's Web site *www.aids.gov/pacha.*

SUPPLEMENTARY INFORMATION: PACHA was established by Executive Order 12963, dated June 14, 1995 as amended by Executive Order 13009, dated June 14, 1996. The Council was established to provide advice, information, and recommendations to the Secretary regarding programs and policies intended to promote effective prevention of HIV disease and AIDS. The functions of the Council are solely advisory in nature.

The Council consists of not more than 25 members. Council members are selected from prominent community leaders with particular expertise in, or knowledge of, matters concerning HIV and AIDS, public health, global health, philanthropy, marketing or business, as well as other national leaders held in high esteem from other sectors of society. Council members are appointed by the Secretary or designee, in consultation with the White House Office on National AIDS Policy. The agenda for the upcoming meeting will be posted on the Council's Web site at *www.aids.gov/pacha.*

Public attendance at the meeting is limited to space available. Individuals who plan to attend and need special assistance, such as sign language interpretation or other reasonable accommodations, should notify the designated contact person. Pre-registration for public attendance is advisable and can be accomplished by contacting the PACHA Committee Manager at *melvin.joppy@hhs.gov.* Members of the public will have the opportunity to provide comments at the meeting. Any individual who wishes to participate in the public comment session must register with Melvin Joppy at *melvin.joppy@hhs.gov;* registration for public comment will not be accepted by telephone. Public comment will be limited to two minutes per speaker. Any members of the public who wish to have printed material distributed to PACHA members at the meeting should submit, at a minimum, 1 copy of the materials to the Committee Manager, PACHA, no later than close of business Wednesday, July 18, 2012. Contact information for the PACHA Committee Manager is listed above.

Dated: July 5, 2012.

B. Kaye Hayes,

Executive Director, Presidential Advisory Council on HIV/AIDS.

[FR Doc. 2012–16907 Filed 7–10–12; 8:45 am]

BILLING CODE 4150–43–P

DEPARTMENT OF HEALTH AND HUMAN SERVICES

Agency for Healthcare Research and Quality

Agency Information Collection Activities: Proposed Collection; Comment Request

AGENCY: Agency for Healthcare Research and Quality, HHS.

ACTION: Notice.

SUMMARY: This notice announces the intention of the Agency for Healthcare Research and Quality (AHRQ) to request that the Office of Management and Budget (OMB) approve the proposed information collection project: ''American Recovery and Reinvestment Act ''Developing a Registry of Registries''.'' In accordance with the Paperwork Reduction Act, 44 U.S.C. 3501–3521, AHRQ invites the public to comment on this proposed information collection.

This proposed information collection was previously published in the **Federal Register** on February 23rd, 2012 and allowed 60 days for public comment. Several comments were received. The purpose of this notice is to allow an additional 30 days for public comment.

DATES: Comments on this notice must be received by August 10, 2012.

ADDRESSES: Written comments should be submitted to: AHRQ's OMB Desk Officer by fax at (202) 395–6974 (attention: AHRQ's desk officer) or by email at *OIRA_submission@omb.eop.gov* (attention: AHRQ's desk officer).

Copies of the proposed collection plans, data collection instruments, and specific details on the estimated burden can be obtained from the AHRQ Reports Clearance Officer.

FOR FURTHER INFORMATION CONTACT: Doris Lefkowitz, AHRQ Reports Clearance Officer, (301) 427–1477, or by email at *doris.lefkowitz@AHRQ.hhs.gov.*

SUPPLEMENTARY INFORMATION:

Proposed Project

American Recovery and Reinvestment Act ''Developing a Registry of Registries''

The Food and Drug Administration Modernization Act of 1997, Public Law 105–115, provided for the creation of a Clinical Trials Data Bank, known as *ClinicalTrials.gov.* Since its launch in 2000, the ClinicalTrials.gov system has registered over 90,500 trials. The large volume of studies currently listed in ClinicalTrials.gov and the high usage numbers suggest that the system has been successful at improving access to information about clinical studies. However, while ClinicalTrials.gov supports the listing of observational studies, such listing is not required.

Patient registries are a distinct type of observational study. Patient registries may be designed for many purposes, such as to observe the natural history of disease, examine comparative effectiveness, or fulfill post-approval commitments. Patient registries have specific characteristics that are not currently captured on ClinicalTrials.gov. To date, some registry sponsors have attempted to leverage the observational study model to post patient registry-type records on ClinicalTrials.gov. However, stakeholders have noted that the system does not fully meet their needs.

Patient registries have received significant attention and funding in recent years. Similar to controlled interventional studies, patient registries represent some burden to patients (e.g., time to complete patient reported outcome measures, risk of loss of privacy), who often participate voluntarily in hopes of improving knowledge about a disease or condition. Patient registries also represent a substantial investment of health research resources. Despite these factors, registration of patient registries in ClinicalTrials.gov is not currently required, presenting the potential for duplication of efforts and insufficient dissemination of findings that are not published in the peer-reviewed literature. To ensure that resources are used in the most efficient manner, registries need to be listed in a manner similar to that of trials in ClinicalTrials.gov.

By creating a central point of collection for information about all patient registries in the United States, the Registry of Patient Registries (RoPR) helps to further AHRQ's goals by making information regarding quality, appropriateness, and effectiveness of health services (and patient registries in