

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://ftcpublic.commentworks.com/ftc/RegsBEMZpra>, 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 "Regs BEMZ, PRA Comments, P084812" 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 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 April 9, 2012. 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>.

Willard K. Tom,
General Counsel.

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GENERAL SERVICES ADMINISTRATION

[Notice-FAS-2011-01; Docket No. 2011-0006; Sequence 22]

Providing Refurbishment Services to Federal Agencies

AGENCY: Federal Acquisition Service, U.S. General Services Administration.

ACTION: Notice.

SUMMARY: In accordance with the National Strategy for Electronics Stewardship recommendations (<http://www.gsa.gov/estewardship>), GSA is exploring whether and how best to make cost-effective refurbishment services available to Federal agencies to extend the useful life of non-functional

electronic equipment. GSA seeks to determine whether providing refurbishment as a service to Federal agencies fits into viable business models, what appropriate standards and certifications ought to be considered, and how best to build Federal contracts for such services.

DATES: *Effective date:* Submit comments on or before March 8, 2012.

ADDRESSES: Submit comments identified by "Notice-FAS-2011-01" by any of the following methods:

- *Regulations.gov:* <http://www.regulations.gov>. Submit comments via the Federal eRulemaking portal by inputting "Notice-FAS-2011-01" under the heading "Enter Keyword or ID" and selecting "Search." Select the link "Submit a Comment" that corresponds with "Notice-FAS-2011-01." Follow the instructions provided at the "Submit a Comment" screen. Please include your name, company name (if any), and "Notice-FAS-2011-01" on your attached document.

- *Fax:* (202) 501-4067.

- *Mail:* General Services Administration, Regulatory Secretariat (MVCB), ATTN: Hada Flowers, 1275 First Street, NE., 7th Floor, Washington, DC 20417.

Instructions: Please submit comments only and cite "Notice-FAS-2011-01", in all correspondence related to this case. 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:

Inquiries and clarifications may be sent to Mr. Chris Hoagland, Office of Governmentwide Policy, U.S. General Services Administration, christopher.hoagland@gsa.gov.

SUPPLEMENTARY INFORMATION:

I. Purpose and Information Sought

This notice is for data gathering and planning purposes only, does not constitute a solicitation, and is not to be construed as a commitment by the government to issue a solicitation, Blanket Purchase Agreement (BPA) or award a contract. The Government will not reimburse any respondent for any costs associated with information submitted in response to this notice.

GSA seeks to determine whether providing refurbishment as a service to Federal agencies fits into viable business models, what appropriate standards and certifications ought to be considered, and how best to build Federal contracts for such services. GSA is seeking this information through

voluntary responses to the following questions:

1. If you currently provide refurbishment as a service to customers, including Federal, state, or local government entities, describe the process of obtaining equipment and returning it to the customer, including the typical amount of time between pick-up and return.

2. Is there a minimum number of pieces of electronic equipment that must be provided (e.g., a pallet load, a truckload)?

3. Does providing refurbishment as a service (rather than refurbished equipment) fit into viable business models for computer refurbishment companies?

4. How do the fees you charge per refurbished item compare to the cost of new or used equipment?

5. Describe the process for disposing and recycling of failed equipment. Have all facilities in your recycling and disposal process been certified to safely recycle and manage electronics? If so, what certifications do they hold?

6. Who is responsible for disposition of equipment that cannot be refurbished, the customer or the provider of refurbishment services? Is there an additional fee for disposition of equipment that cannot be refurbished?

7. What certifications should the government require of firms offering refurbishment services, including those developed specifically for recycling facilities (e.g., R2 and e-Stewards)?

Dated: February 2, 2012.

Houston Taylor,

Assistant Commissioner, Office of Acquisition Management, Federal Acquisition Service, General Services Administration.

[FR Doc. 2012-2767 Filed 2-6-12; 8:45 am]

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DEPARTMENT OF HEALTH AND HUMAN SERVICES

Meeting of the Presidential Advisory Council on HIV/AIDS

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

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 Tuesday, February 28, 2012 and Wednesday, February 29, 2012. The meeting will be held from 9:30 a.m. to approximately 5 p.m. on Tuesday, February 28, 2012, and from 10 a.m. to approximately 3 p.m. on Wednesday, February 29, 2012.

ADDRESSES: South Court Auditorium, Eisenhower Executive Office Building, Pennsylvania Avenue and 17th Street NW., Washington, DC.

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.pacha.gov.

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.pacha.gov.

This meeting of the PACHA will be on White House property, thus, each person must be screened and cleared by the U.S. Secret Service. Pre-registration for public attendance is mandatory. Please contact: Melvin Joppy, Office of HIV/AIDS Policy (202) 690-5560 or melvin.joppy@hhs.gov. The second day of the meeting, February 29, will be held in a different room within the same venue; space will be limited. For this reason, members of the public will be accommodated on a first come, first serve basis for this portion of the meeting. Mr. Joppy will need your full name, social security number, date of birth, country of origin, gender, and city

and state of residence to process public access attendance. Registration must be submitted by close of business Tuesday, February 21, 2012.

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. It is requested that any members of the public who wish to have printed material distributed to PACHA members at the meeting submit, at a minimum, two copies of the materials to the Committee Manager, PACHA, no later than close of business Tuesday, February 21, 2012. Contact information for the PACHA Committee Manager is listed above.

Dated: February 1, 2012.

Christopher Bates,

Executive Director, Presidential Advisory Council on HIV/AIDS.

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DEPARTMENT OF HEALTH AND HUMAN SERVICES

Centers for Medicare & Medicaid Services

[Document Identifier: CMS-10421]

Agency Information Collection Activities: Proposed Collection; Comment Request

AGENCY: Centers for Medicare & Medicaid Services, HHS.

In compliance with the requirement of section 3506(c)(2)(A) of the Paperwork Reduction Act of 1995, the Centers for Medicare & Medicaid Services (CMS) is publishing the following summary of proposed collections for public comment. Interested persons are invited to send comments regarding this burden estimate or any other aspect of this collection of information, including any of the following subjects: (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.

1. *Type of Information Collection Request:* New collection; *Title of Information Collection:* Fee-for-Service Recovery Audit Prepayment Review Demonstration and Prior Authorization Demonstration; *Use:* The Centers for Medicare & Medicaid Services (CMS) is requesting the Office of Management and Budget (OMB) approval of the collections required for two demonstrations of prepayment review and prior authorization. The first demonstration would allow Medicare Recovery Auditors to review claims on a pre-payment basis in certain States. The second demonstration would establish a prior authorization program for Power Mobility Device claims in certain States.

For the Recovery Audit Prepayment Review Demonstration, CMS and its agents will request additional documentation, including medical records, to support submitted claims. As discussed in more detail in Chapter 3 of the Program Integrity Manual, additional documentation includes any medical documentation, beyond what is included on the face of the claim that supports the item or service that is billed. For Medicare to consider coverage and payment for any item or service, the information submitted by the provider or supplier (e.g., claims) must be supported by the documentation in the patient's medical records. When conducting complex medical review, the contractor specifies documentation they require in accordance with Medicare's rules and policies. In addition, providers and suppliers may supply additional documentation not explicitly listed by the contractor. This supporting information may be requested by CMS and its agents on a routine basis in instances where diagnoses on a claim do not clearly indicate medical necessity, or if there is a suspicion of fraud.

For the Prior Authorization of Power Mobility Devices (PMDs) Demonstration, CMS will pilot prior authorization for Power Mobility Devices. Prior authorization will allow the applicable documentation that supports a claim to be submitted before the item is delivered. For prior authorization, relevant documentation for review is submitted before the item is delivered or the service is rendered. CMS will conduct this demonstration in California, Florida, Illinois, Michigan, New York, North Carolina and Texas based on beneficiary address as reported to the Social Security Administration and recorded in the Common Working File (CWF). For the demonstration, a prior authorization request can be completed by the (ordering) physician