record of this proceeding, including on the publicly accessible FTC Web site, at *http://www.ftc.gov/os/publiccomments.shtm.* Because comments will be made public, they should not include any sensitive personal information, such as any individual'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. Comments also should not include any sensitive health information, such as medical records or other individually identifiable health information. In addition, comments should not include ''trade 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 Federal Trade Commission Act (FTC Act), 15 U.S.C. 46(f), and FTC Rule 4.10(a)(2), 16 CFR 4.10(a)(2). Comments containing material for which confidential treatment is requested must be filed in paper form, must be clearly labeled ''Confidential,'' and must comply with FTC Rule 4.9(c).[8]

A comment filed in paper form should include the ''Pet Medications Workshop, Project No. P12–1201'' reference both in the text and on the envelope, and should be mailed or delivered to the following address: Federal Trade Commission, Office of the Secretary, Room H–113 (Annex X), 600 Pennsylvania Avenue NW., Washington, DC 20580. The FTC is requesting that any comment filed in paper form be sent by courier or overnight service, if possible, because U.S. postal mail in the Washington area and at the Commission is subject to delay due to heightened security precautions. 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, whether filed in paper or electronic form. Comments received will be available to the public on the FTC Web site, to the extent practicable, at *http://www.ftc.gov/os/publiccomments.shtm.* As a matter of discretion, the FTC makes every effort to remove home contact information for individuals from the public comments it receives before placing those comments on the FTC Web site. More information, including routine uses permitted by the Privacy Act, may be found in the FTC's privacy policy, at *http://www.ftc.gov/ftc/privacy.htm.*

By direction of the Commission.

Donald S. Clark,

Secretary.

[FR Doc. 2012–16594 Filed 7–6–12; 8:45 am]

BILLING CODE 6750–01–P

[8] The comment must be accompanied by an explicit request for confidential treatment, including the factual and legal basis for the request, and must identify the specific portions of the comment to be withheld from the public record. The request will be granted or denied by the Commission's General Counsel, consistent with applicable law and the public interest. *See* FTC Rule 4.9(c), 16 CFR 4.9(c).

GENERAL SERVICES ADMINISTRATION

[FMR Bulletin—PBS–2012–03; Docket 2012–0002; Sequence 11]

Federal Management Regulation; FMR Bulletin PBS–2012–03; Redesignations of Federal Buildings: Correction

AGENCY: Public Buildings Service (PBS), General Services Administration (GSA).

ACTION: Notice of a bulletin; correction.

SUMMARY: The U.S. General Services Administration published in the **Federal Register** of June 13, 2012, a bulletin announcing the designation and redesignation of three Federal buildings. Inadvertently, the two-letter State ''AL'' was incorrectly identified with the city of Anchorage. This document corrects the abbreviation of the State of Anchorage to ''AK''.

DATES: *Effective Date:* July 9, 2012.

FOR FURTHER INFORMATION CONTACT: U.S. General Services Administration, Public Buildings Service (PBS), 1800 F Street NW., Washington, DC 20405, telephone number: 202–501–1100.

SUPPLEMENTARY INFORMATION: The GSA published a document in the **Federal Register** of June 13, 2012, (77 FR 35393). Inadvertently, the two-letter State for the city Anchorage was identified incorrectly. This document corrects the abbreviation of the State for the city Anchorage to read ''AK''.

Correction

In FR Doc. 2012–14416 published in the **Federal Register** at 77 FR 35393, June 13, 2012 make the following correction:

On page 35393, in the table, first and second columns, second entries, remove ''Anchorage, AL'' and add ''Anchorage, AK'' in their places.

Dated: June 21, 2012.

Dan Tangherlini,

Acting Administrator of General Services.

[FR Doc. 2012–16712 Filed 7–6–12; 8:45 am]

BILLING CODE 6820–23–P

DEPARTMENT OF HEALTH AND HUMAN SERVICES

Meeting of the Scientific Advisory Committee on Alternative Toxicological Methods (SACATM)

AGENCY: Division of the National Toxicology Program (DNTP), National Institute of Environmental Health Sciences (NIEHS), National Institutes of Health (NIH), HHS.

ACTION: Meeting announcement and request for comments.

SUMMARY: Pursuant to section 10(a) of the Federal Advisory Committee Act, as amended (5 U.S.C. appendix 2), notice is hereby given of a meeting of SACATM on September 5–6, 2012, at the Rodbell Auditorium, Rall Building at the NIEHS, 111 T.W. Alexander Drive, Research Triangle Park, NC 27709. The meeting is open to the public with attendance limited only by the space available. The meeting will be webcast through a link at (*http://www.niehs.nih.gov/news/video/live*). SACATM advises the Interagency Coordinating Committee on the Validation of Alternative Methods (ICCVAM), the NTP Interagency Center for the Evaluation of Alternative Toxicological Methods (NICEATM), and the Director of the NIEHS and NTP regarding statutorily mandated duties of ICCVAM and activities of NICEATM.

DATES: The SACATM meeting will be held on September 5–6, 2012. The meeting is tentatively scheduled from 8:30 a.m. Eastern Daylight Time to 5:30 p.m. on September 5 and 8:30 a.m. until adjournment on September 6. All individuals who plan to attend are encouraged to register online at the NTP Web site (*http://ntp.niehs.nih.gov/go/32822*) by August 29, 2012. In order to facilitate planning, persons wishing to make an oral presentation are asked to notify Dr. Lori White, NTP Designated Federal Officer, via online registration, phone, or email by August 29, 2012 (see **ADDRESSES** below). Written comments should also be received by August 29, 2012, to enable review by SACATM and NIEHS/DNTP staff before the meeting. TTY users should contact the Federal TTY Relay Service at 800–877–8339. Requests should be made at least 5 business days in advance of the event.

ADDRESSES: The SACATM meeting will be held at the Rodbell Auditorium, Rall Building at the NIEHS, 111 T.W. Alexander Drive, Research Triangle Park, NC 27709. Public comments and other correspondence should be directed to Dr. Lori White (Office of Liaison, Policy and Review, DNTP, NIEHS, P.O. Box 12233, MD K2–03,

Research Triangle Park, NC 27709; telephone: 919–541–9834 or email: *whiteld@niehs.nih.gov*). Courier address: NIEHS, 530 Davis Drive, Room 2136, Morrisville, NC 27560.

SUPPLEMENTARY INFORMATION:

Preliminary Agenda and Other Meeting Information

A preliminary agenda, roster of SACATM members, background materials, public comments, and any additional information, when available, will be posted on the SACATM meeting Web site (*http://ntp.niehs.nih.gov/go/32822*) or available upon request (see **ADDRESSES** above). Following the meeting, summary minutes will be prepared and available on the SACATM Web site or upon request.

Request for Comments

Both written and oral public input on the agenda topics is invited. Written comments received in response to this notice will be posted on the NTP Web site. Persons submitting written comments should include their name, affiliation (if applicable), and sponsoring organization (if any) with the document. Time is allotted during the meeting for presentation of oral comments and each organization is allowed one time slot per public comment period. At least 7 minutes will be allotted for each speaker, and if time permits, may be extended up to 10 minutes at the discretion of the chair. Registration for oral comments will also be available on-site, although time allowed for presentation by on-site registrants may be less than for pre-registered speakers and will be determined by the number of persons who register at the meeting. In addition to in-person oral comments at the meeting, public comments can be presented by teleconference line. There will be 50 lines for this call; availability will be on a first-come, first-served basis. The available lines will be open from 8:00 a.m. until 5:30 p.m. on September 5 and 8:30 a.m. to adjournment on September 6, although public comments will be received only during the formal public comment periods, which will be indicated on the preliminary agenda. The access number for the teleconference line will be provided to registrants by email prior to the meeting.

Persons registering to make oral comments are asked to do so through the online registration form (*http://ntp.niehs.nih.gov/go/32822*) and to send a copy of their statement to Dr. White (see **ADDRESSES** above) by August 29, 2012, to enable review by SACATM, NICEATM–ICCVAM, and NIEHS/DNTP staff prior to the meeting. Written statements can supplement and may expand the oral presentation. If registering on-site and reading from written text, please bring 40 copies of the statement for distribution and to supplement the record.

Background Information on ICCVAM, NICEATM, and SACATM

ICCVAM is an interagency committee composed of representatives from 15 Federal regulatory and research agencies that require, use, generate, or disseminate toxicological and safety testing information. ICCVAM conducts technical evaluations of new, revised, and alternative safety testing methods with regulatory applicability and promotes the scientific validation and regulatory acceptance of toxicological and safety testing methods that more accurately assess the safety and hazards of chemicals and products and that reduce, refine (decrease or eliminate pain and distress), or replace animal use. The ICCVAM Authorization Act of 2000 (42 U.S.C. 285*l*–3) established ICCVAM as a permanent interagency committee of the NIEHS under NICEATM. NICEATM administers ICCVAM, provides scientific and operational support for ICCVAM-related activities, and conducts independent validation studies to assess the usefulness and limitations of new, revised, and alternative test methods and strategies. NICEATM and ICCVAM work collaboratively to evaluate new and improved test methods and strategies applicable to the needs of U.S. Federal agencies. NICEATM and ICCVAM welcome the public nomination of new, revised, and alternative test methods and strategies for validation studies and technical evaluations. Additional information about ICCVAM and NICEATM can be found on the NICEATM–ICCVAM Web site *(http://iccvam.niehs.nih.gov).*

SACATM was established in response to the ICCVAM Authorization Act [Section 285*l*–3(d)] and is composed of scientists from the public and private sectors. SACATM advises ICCVAM, NICEATM, and the Director of the NIEHS and NTP regarding statutorily mandated duties of ICCVAM and activities of NICEATM. SACATM provides advice on priorities and activities related to the development, validation, scientific review, regulatory acceptance, implementation, and national and international harmonization of new, revised, and alternative toxicological test methods. Additional information about SACATM, including the charter, roster, and records of past meetings, can be found at *http://ntp.niehs.nih.gov/go/167.*

Dated: June 27, 2012.

John R. Bucher,

Associate Director, National Toxicology Program.

[FR Doc. 2012–16675 Filed 7–6–12; 8:45 am]

BILLING CODE 4140–01–P

DEPARTMENT OF HEALTH AND HUMAN SERVICES

Centers for Disease Control and Prevention

[30Day-12–0856]

Agency Forms Undergoing Paperwork Reduction Act Review

The Centers for Disease Control and Prevention (CDC) publishes a list of information collection requests under review by the Office of Management and Budget (OMB) in compliance with the Paperwork Reduction Act (44 U.S.C. chapter 35). To request a copy of these requests, call the CDC Reports Clearance Officer at (404) 639–7570 or send an email to *omb@cdc.gov.* Send written comments to CDC Desk Officer, Office of Management and Budget, Washington, DC or by fax to (202) 395–5806. Written comments should be received within 30 days of this notice.

Proposed Project

National Quitline Data Warehouse (OMB No. 0920–0856, exp. 7/31/2012)—Revision—National Center for Chronic Disease Prevention and Health Promotion (NCCDPHP), Centers for Disease Control and Prevention (CDC).

Background and Brief Description

Tobacco use remains the leading preventable cause of disease and death in the United States, resulting in approximately 440,000 deaths annually and contributing to $92 billion annually in lost worker productivity. Although the prevalence of current smoking among adults decreased significantly since its peak in the 1960s, overall smoking prevalence among U.S. adults has remained virtually unchanged during the past five years. Large disparities in smoking prevalence continue to exist among members of racial/ethnic minority groups and individuals of low socioeconomic status.

The National Tobacco Control Program (NTCP) was established by CDC to help reduce tobacco-related disease, disability, and death. The NTCP provides funding for state quitlines, which provide telephone-based tobacco cessation services to help tobacco users