Dated: February 7, 2012.

Bernadette B. Wilson,

Senior Program Analyst. Executive Secretariat.

This Notice Issued February 7, 2012.

[FR Doc. 2012–3175 Filed 2–7–12; 4:15 pm]

BILLING CODE 6570–01–P

DEPARTMENT OF DEFENSE

GENERAL SERVICES ADMINISTRATION

NATIONAL AERONAUTICS AND SPACE ADMINISTRATION

[OMB Control No. 9000–0142; Docket 2011–0079; Sequence 19]

Federal Acquisition Regulation; Submission for OMB Review; Past Performance Information

AGENCIES: Department of Defense (DOD), General Services Administration (GSA), and National Aeronautics and Space Administration (NASA).

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

SUMMARY: Under the provisions of the Paperwork Reduction Act the Regulatory Secretariat will be submitting to the Office of Management and Budget (OMB) a request to review and approve an extension of a previously approved information collection requirement concerning past performance information. A notice was published in the **Federal Register** at 76 FR 67153, on October 31, 2011. No comments were received.

Public comments are particularly invited on: Whether this collection of information is necessary for the proper performance of functions of the FAR, 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; and ways in which we can minimize the burden of the collection of information on those who are to respond, through the use of appropriate technological collection techniques or other forms of information technology.

DATES: Submit comments on or before March 12, 2012.

ADDRESSES: Submit comments identified by Information Collection 9000–0142, Past Performance Information, by any of the following methods:

- *Regulations.gov: http://www.regulations.gov.* Submit comments via the Federal eRulemaking portal by inputting ''Information Collection 9000–0142, Past Performance Information,'' under the heading ''Enter Keyword or ID'' and selecting ''Search.'' Select the link ''Submit a Comment'' that corresponds with ''Information Collection 9000–0142, Past Performance Information.'' Follow the instructions provided at the ''Submit a Comment'' screen. Please include your name, company name (if any), and ''Information Collection 9000–0142, Past Performance Information,'' 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 9000–0142, Past Performance Information.

Instructions: Please submit comments only and cite Information Collection 9000–0142, Past Performance Information, 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. Curtis E. Glover, Sr., Procurement Analyst, Acquisition Policy Division, at GSA (202) 501–1448 or email *Curtis.glover@gsa.gov*.

SUPPLEMENTARY INFORMATION:

A. Purpose

Past performance information regarding a contractor's actions under previously awarded contracts is relevant information for future source selection purposes.

B. Annual Reporting Burden

Respondents: 150,000.
Responses per Respondent: 4.
Annual Responses: 600,000.
Hours per Response: 2.
Total Burden Hours: 1,200,000.
Obtaining Copies of Proposals: Requesters may obtain a copy of the information collection documents from the General Services Administration, Regulatory Secretariat (MVCB), 1275 First Street NE., Washington, DC 20417, telephone (202) 501–4755. Please cite OMB Control No. 9000–0142, Past Performance Information, in all correspondence.

Dated: February 3, 2012.

Laura Auletta,

Director, Office of Governmentwide Acquisition Policy, Office of Acquisition Policy, Office of Governmentwide Policy.

[FR Doc. 2012–3050 Filed 2–8–12; 8:45 am]

BILLING CODE 6820–EP–P

DEPARTMENT OF HEALTH AND HUMAN SERVICES

Meeting of the Secretary's Advisory Committee on Human Research Protections

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

ACTION: Notice.

SUMMARY: Pursuant to Section 10(a) of the Federal Advisory Committee Act, U.S.C. Appendix 2, notice is hereby given that the Secretary's Advisory Committee on Human Research Protections (SACHRP) will hold its twenty-seventh meeting. The meeting will be open to the public. Information about SACHRP and the meeting agenda will be posted on the SACHRP Web site at: *http://www.dhhs.gov/ohrp/sachrp/mtgings/index.html.*

DATES: The meeting will be held on Tuesday, February 28, 2012 from 8:30 a.m. until 5 p.m. and Wednesday, February 29, 2012 from 8:30 a.m. until 5:00 pm.

ADDRESSES: U.S. Department of Health & Human Services, 200 Independence Avenue SW., Hubert H. Humphrey Building, Room 705, Washington, DC 20201.

FOR FURTHER INFORMATION CONTACT: Jerry Menikoff, M.D., J.D., Director, Office for Human Research Protections (OHRP), or Julia Gorey, J.D., Executive Director, SACHRP; U.S. Department of Health and Human Services, 1101 Wootton Parkway, Suite 200, Rockville, Maryland 20852; 240–453–8141; fax: 240–453–6909; email address: *Julia.Gorey@hhs.gov*.

SUPPLEMENTARY INFORMATION: Under the authority of 42 U.S.C. 217a, Section 222 of the Public Health Service Act, as amended, SACHRP was established to provide expert advice and recommendations to the Secretary of Health and Human Services and the Assistant Secretary for Health on issues and topics pertaining to or associated with the protection of human research subjects.

The meeting will open February 28 with remarks from SACHRP Chair Dr. Barbara Bierer and OHRP Director Dr. Jerry Menikoff, followed by a summary report from the Presidential Commission for the Study of Bioethical Issues on that group's recent report *Moral Science: Protecting Participants in Human Subjects Research.* This will be followed by a summary of public comment from OHRP on the ANPRM *Human Subjects Research Protections:*

Enhancing Protections for Research Subjects and Reducing Burden, Delay and Ambiguity for Investigators. The afternoon will also include a presentation from the Subpart A Subcommittee (SAS) to inform SACHRP of recent work. SAS is charged with developing recommendations for consideration by SACHRP regarding the application of subpart A of 45 CFR part 46 in the current research environment; this subcommittee was established by SACHRP in October 2006.

On February 29, SACHRP will hear recommendations from the Subcommittee on Harmonization (SOH). SOH was established by SACHRP at its July 2009 meeting, and is charged with identifying and prioritizing areas in which regulations and/or guidelines for human subjects research adopted by various agencies or offices within HHS would benefit from harmonization, consistency, clarity, simplification and/or coordination. Following the SOH report, SACHRP will hear a discussion on the IRB use of component analysis, utilizing speakers from the FDA and academia.

Public Comment will be heard on both days.

Public attendance at the meeting is limited to space available. Individuals who plan to attend the meeting and need special assistance, such as sign language interpretation or other reasonable accommodations, should notify the designated contact persons. Members of the public will have the opportunity to provide comments on both days of the meeting. Public comment will be limited to five minutes per speaker. Any members of the public who wish to have printed materials distributed to SACHRP members for this scheduled meeting should submit materials to the Executive Director, SACHRP, prior to the close of business February 23, 2012.

Dated: February 3, 2012.

Jerry Menikoff,

Director, Office for Human Research Protections, Executive Secretary, Secretary's Advisory Committee on Human Research Protections.

[FR Doc. 2012–2958 Filed 2–8–12; 8:45 am]

BILLING CODE 4150–36–P

DEPARTMENT OF HEALTH AND HUMAN SERVICES

Agency for Toxic Substances and Disease Registry

[ATSDR–273]

Notice of Development of Set 25 Toxicological Profiles

AGENCY: Agency for Toxic Substances and Disease Registry (ATSDR), Department of Health and Human Services (DHHS).

ACTION: Notice.

SUMMARY: This notice announces the development of Set 25 Toxicological Profiles, which will consist of four updated profiles. ATSDR will make these profiles available to the public on or about October 17, 2012 and will solicit public comments at that time for a 90-day period. Electronic access to these documents will be available at the ATSDR Web site: *http://www.atsdr.cdc.gov/toxprofiles/index.asp.*

Set 25 Toxicological Profiles

The following toxicological profiles are now being developed:

	Name	CAS
1	Hexachlorobenzene (UPDATE)	118–74–1
2	Endosulfan (UPDATE)	115–29–7
	Endosulfan sulfate	1031–07–8
	Endosulfan-alpha	95–99–98
	Endosulfan-beta	33213–65–9
3	1,1-Dichloroethane (UPDATE)	75–34–3
4	Dinitrotoluenes (DNT) (UPDATE):	
	2,3-DNT	602–01–7
	2,4-DNT	121–14–2
	2,5-DNT	619–15–8
	2,6-DNT	606–20–2
	3,4-DNT	610–39–9
	3,5-DNT	618–85–9

SUPPLEMENTARY INFORMATION: The Superfund Amendments and Reauthorization Act of 1986 (SARA) (42 U.S.C. 9601 *et seq.*) amended the Comprehensive Environmental Response, Compensation, and Liability Act of 1980 (CERCLA or Superfund) (42 U.S.C. 9601 *et seq.*) by establishing certain requirements for ATSDR and the U.S. Environmental Protection Agency (EPA) with regard to hazardous substances that are most commonly found at facilities on the CERCLA National Priorities List (NPL). Among these statutory requirements is a mandate for the Administrator of ATSDR to prepare toxicological profiles for each substance included on the Priority List of Hazardous Substances (*www.atsdr.cdc.gov/SPL*). This list names 275 hazardous substances that pose the most significant potential threat to human health as determined by ATSDR and EPA. The availability of the revised list of the 275 priority substances was announced in the **Federal Register** on November 3, 2011 (76 FR 68193). For prior versions of the list of substances, see **Federal Register** notices dated April 17, 1987 (52 FR 12866); October 20, 1988 (53 FR 41280); October 26, 1989 (54 FR 43619); October 17, 1990 (55 FR 42067); October 17, 1991 (56 FR 52166); October 28, 1992 (57 FR 48801); February 28, 1994 (59 FR 9486); April 29, 1996 (61 FR 18744; November 17, 1997 (62 FR 61332); October 21, 1999 (64 FR 56792); October 25, 2001 (66 FR 54014); November 7, 2003 (68 FR 63098); December 7, 2005 (70 FR 70284); and March 6, 2008 (73 FR 12178).

Notice of the availability of drafts of these four updated toxicological profiles for public review and comment will be published in the **Federal Register** on or about October 17, 2012, with notice of a 90-day public comment period for each profile, starting from the actual release date. Following the close of the comment period, chemical-specific comments will be addressed, and, where appropriate, changes will be incorporated into each profile.

FOR FURTHER INFORMATION CONTACT: Commander Jessilynn B. Taylor, Division of Toxicology and Environmental Medicine, Agency for Toxic Substances and Disease Registry, 1600 Clifton Road NE., Mail Stop F–62,