

**PLACE:** 999 E Street, NW., Washington, DC (Ninth Floor).

**STATUS:** This Meeting will be Open to the Public.

**ITEMS TO BE DISCUSSED:** Correction and Approval of the Minutes for the Meeting of June 15, 2011.

Draft Advisory Opinion 2011–11: Mr. Stephen Colbert.

Draft Advisory Opinion 2011–12: Majority PAC and House Majority PAC. Management and Administrative Matters.

Individuals who plan to attend and require special assistance, such as sign language interpretation or other reasonable accommodations, should contact Shawn Woodhead Werth, Commission Secretary and Clerk, at (202) 694–1040, at least 72 hours prior to the hearing date.

**PERSON TO CONTACT FOR INFORMATION:** Judith Ingram, Press Officer, Telephone: (202) 694–1220.

**Shawn Woodhead Werth,**

*Secretary and Clerk of the Commission.*

[FR Doc. 2011–16211 Filed 6–23–11; 4:15 pm]

**BILLING CODE 6715–01–M**

## DEPARTMENT OF DEFENSE

### General Services Administration;

### National Aeronautics and Space Administration

[OMB Control No. 9000–0018; Docket 2011–0079; Sequence 2]

### Federal Acquisition Regulation; Information Collection; Certification of Independent Price Determination and Parent Company and Identifying Data

**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 (44 U.S.C. chapter 35), the Regulatory Secretariat (MVCB) will be submitting to the Office of Management and Budget (OMB) a request to review and approve an extension of a currently approved information collection requirement concerning certification of independent price determination and parent company and identifying data.

*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 July 27, 2011.

**ADDRESSES:** Submit comments identified by Information Collection 9000–0018 by any of the following methods:

- *Regulations.gov:* <http://www.regulations.gov>. Submit comments via the Federal eRulemaking portal by inputting “Information Collection 9000–0018” under the heading “Enter Keyword or ID” and selecting “Search”. Select the link “Submit a Comment” that corresponds with “Information Collection 9000–0018”. Follow the instructions provided at the “Submit a Comment” screen. Please include your name, company name (if any), and “Information Collection 9000–0018” 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 9000–0018.

*Instructions:* Please submit comments only and cite Information Collection 9000–0018, 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. Anthony Robinson, Procurement Analyst, Contract Policy Branch, GSA (202) 501–2658 or e-mail [Anthony.robinson@gsa.gov](mailto:Anthony.robinson@gsa.gov).

### SUPPLEMENTARY INFORMATION:

#### A. Purpose

Agencies are required to report under 41 U.S.C. 252(d) and 10 U.S.C. 2305(d) suspected violations of the antitrust laws (*e.g.*, collusive bidding, identical bids, uniform estimating systems, *etc.*) to the Attorney General.

As a first step in assuring that Government contracts are not awarded to firms violating such laws, offerors on Government contracts must complete the certificate of independent price determination. An offer will not be considered for award where the

certificate has been deleted or modified. Deletions or modifications of the certificate and suspected false certificates are reported to the Attorney General.

### B. Annual Reporting Burden

*Respondents:* 64,250.

*Responses per Respondent:* 20.

*Total Responses:* 1,285,000.

*Hours per Response:* .01.

*Total Burden Hours:* 12,850.

*Obtaining Copies of Proposals:*

Requesters may obtain a copy of the information collection documents from the General Services Administration, Regulatory Secretariat Branch (MVCB), 1275 First Street, NE., Washington, DC 20417, telephone (202) 501–4755. Please cite OMB Control No. 9000–0018, Certification of Independent Price Determination and Parent Company and Identifying Data, in all correspondence.

Dated: June 9, 2011.

**Millisa Gary,**

*Acting Director, Federal Acquisition Policy Division.*

[FR Doc. 2011–16054 Filed 6–24–11; 8:45 am]

**BILLING CODE 6820–EP–P**

## DEPARTMENT OF DEFENSE

### GENERAL SERVICES ADMINISTRATION

### NATIONAL AERONAUTICS AND SPACE ADMINISTRATION

[OMB Control No. 9000–0164; Docket 2011–0079; Sequence 20]

### Federal Acquisition Regulation; Submission for OMB Review; Contractor Business Ethics Compliance Program and Disclosure Requirements

**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 (44 U.S.C. chapter 35), the Regulatory Secretariat (MVCB) 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 contractor business ethics compliance program and disclosure requirements.

*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 August 26, 2011.

**ADDRESSES:** Submit comments identified by Information Collection 9000-0164, Contractor Business Ethics Compliance Program and Disclosure Requirements, by any of the following methods:

- *Regulations.gov:* <http://www.regulations.gov>. Submit comments via the Federal eRulemaking portal by inputting "Information Collection 9000-0164, Contractor Business Ethics Compliance Program and Disclosure Requirements", under the heading "Enter Keyword or ID" and selecting "Search". Select the link "Submit a Comment" that corresponds with "Information Collection 9000-0164, Contractor Business Ethics Compliance Program and Disclosure Requirements". Follow the instructions provided at the "Submit a Comment" screen. Please include your name, company name (if any), and "Information Collection 9000-0164, Contractor Business Ethics Compliance Program and Disclosure Requirements", 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 9000-0164, Contractor Business Ethics Compliance Program and Disclosure Requirements.

*Instructions:* Please submit comments only and cite Information Collection 9000-0164, Contractor Business Ethics Compliance Program and Disclosure Requirements, 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. Anthony Robinson, Procurement Analyst, Acquisition Policy Division, GSA (202) 501-2658 or e-mail [Anthony.Robinson@gsa.gov](mailto:Anthony.Robinson@gsa.gov).

**SUPPLEMENTARY INFORMATION:**

#### A. Purpose

The collection applies to the FAR requirements for a contractor code of business ethics and conduct, an internal control system, and disclosure to the Government of certain violations of criminal law, violations of the civil False Claims Act, or significant overpayments.

#### B. Annual Reporting Burden

*Respondents:* 284.

*Responses per Respondent:* 1.

*Total Responses:* 284.

*Hours per Response:* 60.

*Total Burden Hours:* 17,040.

*Obtaining Copies of Proposals:*

Requesters may obtain a copy of the information collection documents from the General Services Administration, Regulatory Secretariat Branch (MVCB), 1275 First Street, NE., Washington, DC 20417, telephone (202) 501-4755. Please cite OMB Control No. 9000-0164, Contractor Business Ethics Compliance Program and Disclosure Requirements, in all correspondence.

Dated: June 8, 2011.

**Millisa Gary,**

*Acting Director, Office of Governmentwide Acquisition Policy.*

[FR Doc. 2011-16058 Filed 6-24-11; 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-fifth 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/mtgtings/index.html>.

**DATES:** The meeting will be held on Tuesday, July 19, 2011 from 8:30 a.m. until 5 p.m. and Wednesday, July 20, 2011 from 8:30 a.m. until 5 p.m.

**ADDRESSES:** U.S. Department of Health and Human Services, 200 Independence Avenue, SW., Hubert H. Humphrey Building, Room 800, 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; *e-mail address:* [Julia.Gorey@hhs.gov](mailto: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.

On July 19, 2011, SACHRP will hear a presentation by the Executive Director of the Presidential Commission for the Study of Bioethical Issues focusing on the work of the Commission; this will be followed by SACHRP discussion. After lunch, SACHRP will hear the report of the Subpart A Subcommittee (SAS). 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.

Recommendations to be discussed focus on the return of research results to subject, internet-based research, and improvements to the informed consent process.

On July 20, 2011, the morning will open with a report from the Subcommittee on Harmonization (SOH). The 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. In the afternoon, SACHRP will hear a panel of speakers discussing consequences and processes surrounding scientific misconduct and fraud.

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