

The USAspending.gov Web site, provides information, as collected from Federal agencies, to the public in accordance with the Federal Funding Accountability and Transparency Act of 2006 (Transparency Act).

USAspending.gov is a public-friendly Web site that provides details regarding each Federal award, such as: the name and location of the entity receiving the award, the amount of the award, funding agency for the award, *etc.* Additionally, the IT dashboard Web site, which is a part of USAspending.gov, provides details of Federal Information Technology (IT) investments and is based on data received from agency reports to the Office of Management and Budget (OMB). The ability to look at contracts, grants, loans, Information Technology investments, and other types of spending across many agencies, in greater detail, is a key ingredient to building public trust in government and credibility in the professionals who use these agreements. USAspending.gov visitors will be provided opportunities to provide feedback in the spirit of the President's open government and transparency initiative. Examples of feedback mechanisms are:

(1) A "Contact Us" entry page with an optional contact e-mail address for those visitors wishing to identify themselves on the USAspending.gov Web page,

(2) A "Contact Us" entry page with a contact e-mail address on the IT dashboard Web page; and

(3) A Collaborative Work Environment using wiki Web pages, e-mail discussion forum, message archive, shared file workspace, full text search capability, *etc.*

Additional feedback mechanisms may be placed in the future but additional details have not yet been defined regarding them. This information collection request for a generic clearance is a replacement of the emergency ICR approved by OMB. It is being submitted in order to fulfill the public feedback aspects of this important initiative.

Burden Statement: The annual public reporting and recordkeeping burden for this collection of information is estimated to average up to 500 hours per year. Burden means the total time, effort, or financial resources expended by persons to generate, maintain, retain, or disclose or provide information to or for a Federal agency. This includes the time needed to review instructions; develop, acquire, install, and utilize technology and systems for the purposes of collecting, validating, and verifying information, processing and maintaining information, and disclosing

and providing information; adjust the existing ways to comply with any previously applicable instructions and requirements which have subsequently changed; train personnel to be able to respond to a collection of information; search data sources; complete and review the collection of information; and transmit or otherwise disclose the information.

The estimated annual burden request is summarized here:

Affected entities: Anyone that chooses to visit USAspending.gov, including the IT Dashboard Web site.

Estimated total number of respondents: 5,000.

Frequency of responses: 105 per week.

Total Responses: 5000.

Average Burden Hours Per Response: 6 minutes.

Estimated total annual burden hours: 500 hours.

What Is the Next Step in the Process for This ICR?

GSA will consider the comments received and amend the ICR as appropriate. The final ICR package will then be submitted to OMB for review and approval pursuant to 5 CFR 1320.12.

Obtaining Copies of Proposals: Requesters may obtain a copy of the information collection documents from the General Services Administration, Regulatory Secretariat (MVPR), 1800 F Street, NW., Room 4041, Washington, DC 20405, telephone (202) 501-4755. Please cite OMB Control Number 3090-0285, USAspending/IT Dashboard Feedback Mechanisms, in all correspondence.

Dated: December 9, 2009.

Casey Coleman,

Chief Information Officer, General Services Administration.

[FR Doc. E9-29837 Filed 12-15-09; 8:45 am]

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DEPARTMENT OF DEFENSE

GENERAL SERVICES ADMINISTRATION

NATIONAL AERONAUTICS AND SPACE ADMINISTRATION

[OMB Control No. 9000-0173]

Submission for OMB Review; Limitations on Pass-Through Charges

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

ACTION: Notice of request for public comments regarding a new OMB information clearance.

SUMMARY: Under the provisions of the Paperwork Reduction Act of 1995 (44 U.S.C. Chapter 35), the Federal Acquisition Regulation (FAR) Regulatory Secretariat (MVPR) will be submitting to the Office of Management and Budget (OMB) a request to review and approve a new information collection requirement regarding Limitations on Pass-Through Charges.

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 January 15, 2010.

ADDRESSES: Submit comments regarding this burden estimate or any other aspect of this collection of information, including suggestions for reducing this burden to: FAR Desk Officer, OMB, Room 10102, NEOB, Washington, DC 20503, and a copy to the General Services Administration, Regulatory Secretariat (MVPR), General Services Administration, 1800 F Street, NW., Room 4041, Washington, DC 20405. Please cite OMB Control No. 9000-0173, Limitations on Pass-Through Charges, in all correspondence.

FOR FURTHER INFORMATION CONTACT: Mr. Edward Chambers, Procurement Analyst, Contract Policy Branch, at telephone (202) 501-3221 or via e-mail to Edward.chambers@gsa.gov.

SUPPLEMENTARY INFORMATION:

A. Purpose

To enable contracting officers to verify that pass-through charges are not excessive, the provision at 52.215-22 requires offerors submitting a proposal for a contract, task order, or delivery order to provide the following information with its proposal: (1) The percent of effort the offeror intends to perform and the percent expected to be performed by each subcontractor. (2) If the offeror intends to subcontract more than 70 percent of the total cost of work to be performed—(i) The amount of the

offeror's indirect costs and profit/fee applicable to the work to be performed by the subcontractor(s); and (ii) A description of the value added by the offeror as related to the work to be performed by the subcontractor(s). (3) If any subcontractor intends to subcontract to a lower-tier subcontractor more than 70 percent of the total cost of work to be performed under its subcontract—(i) The amount of the subcontractor's indirect costs and profit/fee applicable to the work to be performed by the lower-tier subcontractor(s); and (ii) A description of the value added by the subcontractor as related to the work to be performed by the lower-tier subcontractor(s).

B. Annual Reporting Burden

Respondents: 25,380.

Responses per Respondent: 1.

Hours per Response: 147,515.

Total Burden Hours: 13,260.

Obtaining Copies of Proposals:

Requesters may obtain a copy of the information collection documents from the General Services Administration, Regulatory Secretariat (MVPR), 1800 F Street, NW., Room 4041, Washington, DC 20405, telephone (202) 501-4755. Please cite OMB Control No. 9000-0173, Limitations on Pass-Through Charges, in all correspondence.

Dated: December 10, 2009.

Al Matera,

Director, Acquisition Policy Division.

[FR Doc. E9-29876 Filed 12-15-09; 8:45 am]

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DEPARTMENT OF HEALTH AND HUMAN SERVICES

Office of the Secretary

Notice of Meeting: Secretary's Advisory Committee on Genetics, Health, and Society

Pursuant to Public Law 92-463, notice is hereby given of the twenty-first meeting of the Secretary's Advisory Committee on Genetics, Health, and Society (SACGHS), U.S. Public Health Service. The meeting will be held from 8:30 a.m. to approximately 5:30 p.m. on Thursday, February 4, 2010, and from 8 a.m. to approximately 3 p.m. on Friday, February 5, 2010, at the Omni Shoreham Hotel, 2500 Calvert Street, NW., Washington, DC 20008. The meeting will be open to the public with attendance limited to space available. The meeting also will be Web cast.

The main agenda items involve the review of a revised report on gene patents and licensing practices, the review of a public consultation draft

report on genetics education and training, and an information-gathering session on the mechanisms and policies related to genomic data sharing. Other agenda items include a preliminary discussion to help plan a future session on implications of an affordable genome; a report on activities of the Clinical Utility and Comparative Effectiveness Task Force; and updates from Federal agencies on activities related to the implementation of the Genetic Information Nondiscrimination Act, the coverage and reimbursement of genetic tests, the oversight of genetic testing, and the retention and use of residual dried blood spot specimens after newborn screening.

As always, the Committee welcomes hearing from anyone wishing to provide public comment on any issue related to genetics, health and society. Individuals who would like to provide public comment should notify the SACGHS Executive Secretary, Ms. Sarah Carr, by telephone at 301-496-9838 or e-mail at carrs@od.nih.gov. The SACGHS office is located at 6705 Rockledge Drive, Suite 750, Bethesda, MD 20892. Anyone planning to attend the meeting who needs special assistance, such as sign language interpretation or other reasonable accommodations, is also asked to contact the Executive Secretary.

Under authority of 42 U.S.C. 217a, Section 222 of the Public Health Service Act, as amended, the Department of Health and Human Services established SACGHS to serve as a public forum for deliberations on the broad range of human health and societal issues raised by the development and use of genetic and genomic technologies and, as warranted, to provide advice on these issues. The draft meeting agenda and other information about SACGHS, including information about access to the Web cast, will be available at the following Web site: http://oba.od.nih.gov/SACGHS/sacghs_meetings.html.

Dated: December 10, 2009.

Jennifer Spaeth,

Director, NIH Office of Federal Advisory Committee Policy.

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DEPARTMENT OF HEALTH AND HUMAN SERVICES

National Institutes of Health

National Center for Complementary and Alternative Medicine; Notice of Meeting

Pursuant to Section 10(d) of the Federal Advisory Committee Act, as amended (5 U.S.C. App.), notice is hereby given of the National Advisory Council for Complementary and Alternative Medicine (NACCAM) meeting.

The meeting will be open to the public as indicated below, with attendance 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 Contact Person listed below in advance of the meeting.

A portion of the meeting will be closed to the public in accordance with the provisions set forth in sections 552b(c)(4) and 552b(c)(6), Title 5 U.S.C., as amended. The grant applications and/or contract proposals and the discussion could disclose confidential trade secrets or commercial property such as patentable material, and personal information concerning individuals associated with the grant applications and/or contract proposals, the disclosure of which would constitute a clearly unwarranted invasion of personal privacy.

Name of Committee: National Advisory Council for Complementary and Alternative Medicine.

Date: February 5, 2010.

Closed: 8:30 a.m. to 10:30 a.m.

Agenda: To review and evaluate grant applications and/or proposals.

Open: 11 a.m. to 5 p.m.

Agenda: Opening remarks by the Director of the National Center for Complementary and Alternative Medicine, presentation of a new research initiative, and other business of the Council.

Place: National Institutes of Health, Neuroscience Building, 6001 Executive Boulevard, Conference Rooms C & D, Bethesda, MD 20892.

Contact Person: Martin H. Goldrosen, Ph.D., Executive Secretary, Director, Division of Extramural Activities, National Center for Complementary and Alternative Medicine, National Institutes of Health, 6707 Democracy Blvd., Suite 401, Bethesda, MD 20892, (301) 594-2014.

The public comments session is scheduled from 4:30-5 p.m., but could change depending on the actual time spent on each agenda item. Each speaker will be permitted 5 minutes for their presentation. Interested individuals and representatives of organizations are requested to notify Dr. Martin H. Goldrosen, National Center for