the General Services Administration, Regulatory Secretariat (MVCB), 1275 First Street NE., Washington, DC 20417, telephone (202) 501–4755. Please cite OMB Control No. 3090–0027, Contract Administration, Quality Assurance (GSAR Parts 542 and 546; GSA Form 1678, and GSA Form 308), in all correspondence.


Joseph A. Neurauter,
Director, Office of Acquisition Policy & Senior Procurement Executive.

[FR Doc. 2012–7422 Filed 3–27–12; 8:45 am]
BILLING CODE 6820–61–P

DEPARTMENT OF DEFENSE
GENERAL SERVICES ADMINISTRATION

NATIONAL AERONAUTICS AND SPACE ADMINISTRATION

[OMB Control No. 9000–0035; Docket 2012–0076; Sequence 1]

Federal Acquisition Regulation; Information Collection; Claims and Appeals

AGENCY: 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 claims and appeals.

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 May 29, 2012.

ADDRESSES: Submit comments identified by Information Collection 9000–0035, Claims and Appeals by any of the following methods:

• Regulations.gov: http://www.regulations.gov. Submit comments via the Federal eRulemaking portal by inputting “Information Collection 9000–0035, Claims and Appeals” under the heading “Enter Keyword or ID” and selecting “Search”. Select the link “Submit a Comment” that corresponds with “Information Collection 9000–0035, Claims and Appeals”. Follow the instructions provided at the “Submit a Comment” screen. Please include your name, company name (if any), and “Information Collection 9000–0035, Claims and Appeals” 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 9000–0035, Claims and Appeals.

Instructions: Please submit comments only and cite Information Collection 9000–0035, Claims and Appeals, 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 via email at anthony.robinson@gsa.gov.

SUPPLEMENTARY INFORMATION:

A. Purpose

It is the Government’s policy to try to resolve all contractual issues by mutual agreement at the contracting officer’s level without litigation. Reasonable efforts should be made to resolve controversies prior to submission of a contractor’s claim. The Contract Disputes Act of 1978 (41 U.S.C. 605) requires that claims exceeding $100,000 must be accompanied by a certification that (1) The claim is made in good faith; (2) supporting data are accurate and complete; and (3) the amount requested accurately reflects the contract adjustment for which the contractor believes the Government is liable. The information, as required by FAR clause 52.233–1, Disputes, is used by a contracting officer to decide or resolve the claim. Contractors may appeal the contracting officer’s decision by submitting written appeals to the appropriate officials.

B. Annual Reporting Burden

Respondents: 4,500.

Responses per Respondent: 3.

Annual Responses: 13,500.

Hours per Response: 1.

Total Burden Hours: 13,500.

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) 501–4753. Please cite OMB Control No. 9000–0035, Claims and Appeals, in all correspondence.


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

[FR Doc. 2012–7425 Filed 3–27–12; 8:45 am]
BILLING CODE 6820–EP–P

DEPARTMENT OF DEFENSE
GENERAL SERVICES ADMINISTRATION

NATIONAL AERONAUTICS AND SPACE ADMINISTRATION

[OMB Control No. 9000–0067; Docket 2012–0076; Sequence 10]

Federal Acquisition Regulation; Information Collection; Incentive Contracts

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 incentive contracts.

Public comments are particularly invited on: Whether this collection of information is necessary for the proper performance of functions of the Federal Acquisition Regulation (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.
collection techniques or other forms of information technology.

DATES: Submit comments on or before May 29, 2012.

ADDRESSES: Submit comments identified by Information Collection 9000–0067, Incentive Contracts, by any of the following methods:

• Regulations.gov: http://www.regulations.gov. Submit comments via the Federal eRulemaking portal by inputting “Information Collection 9000–0067, Incentive Contracts” under the heading “Enter Keyword or ID” and selecting “Search”. Select the link “Submit a Comment” that corresponds with “Information Collection 9000–0067, Incentive Contracts”. Follow the instructions provided at the “Submit a Comment” screen. Please include your name, company name (if any), and “Information Collection 9000–0067, Incentive Contracts” 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 9000–0067, Incentive Contracts.

Instructions: Please submit comments only and cite Information Collection 9000–0067, Incentive Contracts, 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. Michael O. Jackson, Procurement Analyst, Office of Acquisition Policy, GSA (202) 208–4949 or via email michael.o.jackson@gsa.gov.

SUPPLEMENTARY INFORMATION:

A. Purpose

In accordance with FAR 16.4, incentive contracts are normally used when a firm fixed-price contract is not appropriate and the required supplies or services can be acquired at lower costs, and sometimes with improved delivery or technical performance, by relating the amount of profit or fee payable under the contract to the contractor’s performance.

The information required periodically from the contractor, such as cost of work already performed, estimated costs of further performance necessary to complete all work, total contract price for supplies or services accepted by the Government for which final prices have been established, and estimated costs allocable to supplies or services accepted by the Government and for which final prices have not been established, is needed to negotiate the final prices of incentive-related items and services. Contractors are required to submit the information in accordance with several incentive fee FAR clauses: FAR 52.216–16, Incentive Price Revision—Firm Target; FAR 52.216–17, Incentive Price Revision—Successive Targets; and FAR 52.216–10, Incentive Fee.

The contracting officer evaluates the information received to determine the contractor’s performance in meeting the incentive target and the appropriate price revision, if any, for the items or services.

B. Annual Reporting Burden

Respondents: 3,000.

Responses Per Respondent: 1.

Annual Responses: 3,000.

Hours Per Response: 1.

Total Burden Hours: 3,000.

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 citeOMB Control No. 9000–0067, Incentive Contracts, in all correspondence.


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

[FR Doc. 2012–7416 Filed 3–27–12; 8:45 am]

BILLING CODE 6820–EP–P

DEPARTMENT OF HEALTH AND HUMAN SERVICES

[Document Identifier OS–0990–0260; 30-Day Notice]

Agency Information Collection Request; 30-Day Public Comment Request

AGENCY: Office of the Secretary, HHS.

In compliance with the requirement of section 3506(c)(2)(A) of the Paperwork Reduction Act of 1995, the Office of the Secretary (OS), Department of Health and Human Services, is publishing the following summary of a proposed collection 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.

To obtain copies of the supporting statement and any related forms for the proposed paperwork collections referenced above, email your request, including your address, phone number, OMB number, and OS document identifier, to Sherette.funcoleman@hhs.gov, or call the Reports Clearance Office on (202) 690–5683. Send written comments and recommendations for the proposed information collections within 30 days of this notice directly to the OS OMB Desk Officer; faxed to OMB at 202–395–5806.


Abstract: Section 491(a) of Public Law 99–158 states that the Secretary of HHS shall by regulation require that each entity applying for HHS support (e.g., a grant, contract, or cooperative agreement) to conduct research involving human subjects submit to HHS assurances satisfactory to the Secretary that it has established an institutional review board (IRB) to review the research in order to ensure protection of the rights and welfare of the human research subjects. IRBs are boards, committees, or groups formally designated by an entity to review, approve, and have continuing oversight of research involving human subjects.

Pursuant to the requirement of the Public Law 99–158, HHS promulgated regulations at 45 CFR part 46, subpart A, the basic HHS Policy for the Protection of Human Subjects. The June 18, 1991 adoption of the common Federal Policy (56 FR 28003) by 15 departments and agencies implements a recommendation of the President’s Commission for the Study of Ethical Problems in Medicine and Biomedical and Behavioral Research which was established on November 9, 1974, by Public Law 95–622. The Common Rule is based on HHS regulations at 45 CFR part 46, subpart A, the basic HHS Policy for the Protection of Human Subjects.