

Dated: August 21, 2009.

Karen V. Gregory,
Secretary.

[FR Doc. E9-20608 Filed 8-25-09; 8:45 am]

BILLING CODE 6730-01-P

DEPARTMENT OF DEFENSE

GENERAL SERVICES ADMINISTRATION

NATIONAL AERONAUTICS AND SPACE ADMINISTRATION

[OMB Control No. 9000-0095]

Federal Acquisition Regulation; Submission for OMB Review; Commerce Patent Regulations

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

ACTION: Notice of request for comments regarding the reinstatement of a previously existing OMB clearance.

SUMMARY: Under the provisions of the Paperwork Reduction Act of 1995 (44 U.S.C. Chapter 35), the Federal Acquisition Regulation (FAR) Secretariat will be submitting to the Office of Management and Budget (OMB) a request to review and approve a reinstatement of a previously approved information collection requirement concerning Commerce Patent Regulations.

Public comments are particularly invited on: Whether this collection of information is necessary; 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 September 25, 2009.

ADDRESSES: Submit comments regarding this burden estimate or any other aspect of this collection of information, including suggestions for reducing this burden, to: General Services Administration (GSA), Regulatory Secretariat (VPR), 1800 F Street, NW., Room 4041, Washington, DC 20405. Please cite OMB Control No. 9000-0095, Commerce Patent Regulations, in all correspondence.

FOR FURTHER INFORMATION CONTACT: Mr. Ernest Woodson, Procurement Analyst, Contract Policy Division, GSA, (202) 501-3775 or e-mail ernest.woodson@gsa.gov.

SUPPLEMENTARY INFORMATION:

A. Purpose

As a result of the Department of Commerce (Commerce) publishing a final rule in the **Federal Register** implementing Pub. L. 98-620 (52 FR 8552, March 18, 1987), a revision to FAR Subpart 27.3 to implement the Commerce regulation was published in the **Federal Register** as an interim rule on June 12, 1989 (54 FR 25060). The final rule was published without change on June 21, 1990.

A Government contractor must report all subject inventions to the contracting officer, submit a disclosure of the invention, and identify any publication, or sale, or public use of the invention (52.227-11(c), 52.227-13(e)(1)). The contracting officer may modify 52.227-11(e) or otherwise supplement the clause to require contractors to submit periodic or interim and final reports listing subject inventions (27.303(b)(2)(i) and (ii)). In order to ensure that subject inventions are reported, the contractor is required to establish and maintain effective procedures for identifying and disclosing subject inventions (52.227-11, Alternate IV; 52.227-13(e)(1)). In addition, the contractor must require his employees, by written agreements, to disclose subject inventions (52.227-11(e)(2); 52.227-13(e)(4)). The contractor also has an obligation to utilize the subject invention, and agree to report, upon request, the utilization or efforts to utilize the subject invention (27.302(e); 52.227-11(f)).

B. Annual Reporting Burden

Respondents: 1,200.

Responses per Respondent: 9.75.

Total Responses: 11,700.

Hours per Response: 3.9.

Total Burden Hours: 45,630.

Obtaining Copies of Proposals:

Requesters may obtain a copy of the information collection documents from the General Services Administration (GSA), Regulatory Secretariat (VPR), 1800 F Street, NW., Room 4041, Washington, DC 20405, telephone (202) 501-4755. Please cite OMB Control No. 9000-0095, Commerce Patent Regulations, in all correspondence.

Dated: August 20, 2009.

Al Matera,

Director, Office of Acquisition Policy.

[FR Doc. E9-20517 Filed 8-25-09; 8:45 am]

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DEPARTMENT OF HEALTH AND HUMAN SERVICES

Meeting of the National Vaccine Advisory Committee

AGENCY: Department of Health and Human Services, Office of the Secretary, Office of Public Health and Science.

ACTION: Notice of meeting.

SUMMARY: As stipulated by the Federal Advisory Committee Act, the Department of Health and Human Services (HHS) is hereby giving notice that the National Vaccine Advisory Committee (NVAC) will hold a meeting. The meeting is open to the public. Pre-registration is required for both public attendance and comment. Individuals who wish to attend the meeting and/or participate in the public comment session should e-mail nvpo@hhs.gov, call 202-690-5566, or complete the on-line form on the NVAC Web site (<http://www.hhs.gov/nvpo/nvac/>) to register.

DATES: The meeting will be held on September 15, 2009, from 9 a.m. to 5:30 p.m. and on September 16, 2009, from 8:30 a.m. to 1:30 p.m.

ADDRESSES: Department of Health and Human Services; Hubert H. Humphrey Building, Room 800; 200 Independence Avenue, SW., Washington, DC 20201.

FOR FURTHER INFORMATION CONTACT: Ms. Andrea Krull, Public Health Advisor, National Vaccine Program Office, Department of Health and Human Services, Room 715-H Hubert H. Humphrey Building, 200 Independence Avenue, SW., Washington, DC 20201. Phone: (202) 690-5566; Fax: (202) 260-1165; e-mail: nvpo@hhs.gov.

SUPPLEMENTARY INFORMATION: Pursuant to Section 2101 of the Public Health Service Act (42 U.S.C. Section 300aa-1), the Secretary of Health and Human Services was mandated to establish the National Vaccine Program to achieve optimal prevention of human infectious diseases through immunization and to achieve optimal prevention against adverse reactions to vaccines. The National Vaccine Advisory Committee was established to provide advice and make recommendations to the Director of the National Vaccine Program, on matters related to the Program's responsibilities. The Assistant Secretary for Health serves as Director of the National Vaccine Program.

Topics to be discussed at the meeting include vaccine safety working group activity, the National Vaccine Plan, implementation plans for recent NVAC recommendations, financial considerations for adult immunizations,