no change in the reporting, recordkeeping and/or third party disclosure requirements. There is no change in the estimated number of respondents/responses, burden hours or annual costs.

Section 2.955 describes for each equipment device subject to verification, the responsible party, as shown in 47 CFR 2.909 shall maintain the records listed as follows:

(1) A record of the original design drawings and specifications and all changes that have been made that may affect compliance with the requirements of § 2.953.

(2) A record of the procedures used for production inspection and testing (if tests were performed) to insure the conformance required by § 2.953. (Statistical production line emission testing is not required.)

(3) A record of the measurements made on an appropriate test site that demonstrates compliance with the applicable regulations in this chapter. The record shall:

(i) Indicate the actual date all testing was performed;

(ii) State the name of the test laboratory, company, or individual performing the verification testing. The Commission may request additional information regarding the test site, the test equipment or the qualifications of the company or individual performing the verification tests;

(iii) Contain a description of how the device was actually tested, identifying the measurement procedure and test equipment that was used;

(iv) Contain a description of the equipment under test (EUT) and support equipment connected to, or installed within, the EUT;

(v) Identify the EUT and support equipment by trade name and model number and, if appropriate, by FCC Identifier and serial number;

(vi) Indicate the types and lengths of connecting cables used and how they were arranged or moved during testing;

(vii) Contain at least two drawings or photographs showing the test set-up for the highest line conducted emission and showing the test set-up for the highest radiated emission. These drawings or photographs must show enough detail to confirm other information contained in the test report. Any photographs used must be focused originals without glare or dark spots and must clearly show the test configuration used;

(viii) List all modifications, if any, made to the EUT by the testing company or individual to achieve compliance with the regulations in this chapter;

(ix) Include all of the data required to show compliance with the appropriate regulations in this chapter; and

(x) Contain, on the test report, the signature of the individual responsible for testing the product along with the name and signature of an official of the responsible party, as designated in § 2.909.

(4) For equipment subject to the provisions in part 15 of this chapter, the records shall indicate if the equipment was verified pursuant to the transition provisions contained in § 15.37 of this chapter.

(b) The records listed in paragraph (a) of this section shall be retained for two years after the manufacture of said equipment item has been permanently discontinued, or until the conclusion of an investigation or a proceeding if the manufacturer or importer is officially notified that an investigation or any other administrative proceeding involving his equipment has been instituted.

The Commission needs and requires the information under FCC Rules at 47 CFR Parts 15 and 18, that RF equipment manufacturers (respondents) ''self determine'' their responsibility for adherence to these rules, as guided by the following criteria:

(a) Whether the RF equipment device that is being marketed complies with the applicable Commission Rules; and

(b) If the operation of the equipment is consistent with the initially documented test results, as reported to the Commission.

The information collection is essential to controlling potential interference to radio communications.

(a) Companies that manufacture RF equipment are the anticipated respondents to this information collection.

(b) This respondent ''public'' generally remains the same, although the types of equipment devices that they manufacture may change in response to changing technologies and to new spectrum allocations made by the Commission.

(c) In addition, the Commission may establish new technical operating standards in response to these changing technologies and in allocating spectrum, which these RF equipment manufacturers must meet to receive their equipment authorization from the FCC.

(d) However, the process that RF equipment manufacturers must follow to verify their compliance, as mandated by 47 CFR Section 2.955 of FCC Rules, will not change despite new technical standards established for specific equipment.

This information collection, therefore, applies to a variety of equipment, which is currently manufactured, may be manufactured in the future, and that operates under varying technical standards.

Federal Communications Commission.

Marlene H. Dortch,

Secretary.

[FR Doc. E8–30130 Filed 12–18–08; 8:45 am]

BILLING CODE 6712–01–P

GENERAL SERVICES ADMINISTRATION

[OMB Control No. 3090–0280]

General Services Administration Acquisition Regulation; Information Collection; Tax Adjustment Clause 552.270–30

AGENCY: Office of the Chief Acquisition Officer, General Services Administration (GSA).

ACTION: Notice of request for comments regarding a renewal to an existing OMB clearance.

SUMMARY: Under the provisions of the Paperwork Reduction Act of 1995 (44 U.S.C. Chapter 35), the General Services Administration will be submitting to the Office of Management and Budget (OMB) a request to review and approve a renewal of a currently approved information collection requirement regarding tax adjustments under leasehold acquisitions. This collection requires contractors to submit information to the Government to substantiate an increase or decrease in real estate taxes under a leasehold acquisition so that the Government can make tax adjustments as necessary to the leasehold acquisition. Information collected under this authority is necessary to assess proper tax adjustments against each leasehold acquisition. The clearance currently expires on April 30, 2009.

Public comments are particularly invited on: Whether the proposed collection of information is necessary for the proper performance of the function of the agency, including whether the information shall have practical utility; the accuracy of the agency's estimate of the burden of the proposed information collection; ways to enhance the quality, utility, and clarity of the information to be collected; ways to minimize the burden of the information collection on respondents including through the use of automated collection techniques or other forms of information technology.

DATES: Submit comments on or before February 17, 2009.

FOR FURTHER INFORMATION CONTACT: Edward Chambers, Procurement Analyst, Contract Policy Division, GSA (202) 501–3221.

ADDRESSES: Submit comments regarding this burden estimate or any other aspect of this collection of information, including suggestions for reducing this burden to the Regulatory Secretariat (VPR), General Services Administration, Room 4041 1800 F Street, NW., Washington, DC 20405. Please cite OMB Control No. 3090–0280, Tax Adjustment Clause 552.270–30, in all correspondence.

SUPPLEMENTARY INFORMATION:

A. Purpose

The General Services Administration (GSA) has various mission responsibilities related to the acquisition and provision supply, service, and leasehold acquisitions. These mission responsibilities generate requirements that are realized through the solicitation and award of various types of contracts. Individual solicitations and resulting contracts may impose unique information collection and reporting requirements on contractors, not required by regulation, but necessary to evaluate particular program accomplishments, measure success in meeting program objectives, or adjust acquisition requirements. Leasehold acquisitions provide for real estate tax adjustments due to changes in real estate taxes on land and buildings occupied by the Government. In a leasehold acquisition, the lessor shall provide the following information regarding real estate taxes: (1) Any notice which may affect the valuation of land and buildings covered by this lease for real estate tax purposes; (2) Any notice of a tax credit or tax refund related to land and buildings covered by this lease; and (3) Each tax bill related to land and building covered by this lease. The lessor is also required to provide the contracting officer a proper invoice including evidence of payment to receive the tax adjustment. Depending on the leasehold acquisition, the tax adjustment can result in either the lessor receiving a credit or the Government receiving a credit.

B. Annual Reporting Burden.

Respondents: 7041.
Responses Per Respondent: 1.
Total Responses: 7041.
Hours Per Response: 6.
Total Burden Hours: 42,246.
Obtaining copies of proposals: Requesters may obtain a copy of the information collection documents from the General Services Administration, Regulatory Secretariat (VPR), 1800 F Street, NW., Room 4041, Washington, DC 20405, telephone (202) 501–4755. Please cite OMB Control No. 3090–0280, Tax Adjustment Clause 552.270–30, in all correspondence.

Dated: December 10, 2008.

Al Matera,

Director,Contract Policy Division.

[FR Doc. E8–30016 Filed 12–18–08; 8:45 am]

BILLING CODE 6820–61–S

DEPARTMENT OF HEALTH AND HUMAN SERVICES

Notice of the Development of a Web-Based System Used To Request Meetings Regarding Medical Countermeasures to Naturally Occurring or Manmade Threats

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

ACTION: Notice.

SUMMARY: The Department of Health and Human Services (HHS) is announcing the availability of a new Web-based system, MedicalCountermeasures.gov. MedicalCountermeasures.gov will enable external stakeholders to request meetings with personnel from the organizations that comprise the Public Health Emergency Medical Countermeasures Enterprise (PHEMCE) regarding medical countermeasures to threats to public health, either naturally occurring or manmade. The goal of these meetings is to provide an opportunity for stakeholders to share information regarding medical countermeasures. The system can be accessed from the Web site *https://www.medicalcountermeasures.gov/RequestMeeting.aspx.*

During the BioShield Stakeholders Workshop, HHS Secretary Michael O. Leavitt announced that HHS would develop a Web based system ''through which those in industry and the research and development community can reach the people they need in the federal government, whether they're looking at a basic level of research or are focused on end-stage development.'' In fulfillment of this promise, HHS has developed MedicalCountermeasures.gov.

MedicalCountermeasures.gov enables external stakeholders to request a meeting with federal representatives from participating PHEMCE agencies regarding medical countermeasures they are developing for use in response to a public health emergency. The information will then be routed to personnel within the relevant PHEMCE agencies, which currently include: The National Institutes of Health (NIH), the Office of the Biomedical Advanced Research and Development Authority (BARDA), the Food and Drug Administration (FDA), and the Department of Veterans Affairs (VA). MedicalCountermeasures.gov also provides information on upcoming and past conferences; procurements and grants; regulatory information; and strategic plans from throughout the PHEMCE agencies.

FOR FURTHER INFORMATION CONTACT: Elizabeth Jarrett, M.A., Office of the Biomedical Advanced Research and Development, Office of the Assistant Secretary for Preparedness and Response, Department of Health and Human Services, 330 Independence Ave., SW., Room G640, Washington, DC 20201; phone: 202–260–1200; e-mail address: *BARDA@hhs.gov.*

Dated: December 9, 2008.

W. Craig Vanderwagen,

Assistant Secretary for Preparedness and Response, Department of Health and Human Services.

[FR Doc. E8–30150 Filed 12–18–08; 8:45 am]

BILLING CODE 4150–37–P

DEPARTMENT OF HEALTH AND HUMAN SERVICES

Opportunity for Cosponsorship of the Integrated Medical, Public Health, Preparedness, and Response Training Summit

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

ACTION: Notice.

SUMMARY: The Department of Health and Human Services' Office of the Assistant Secretary for Preparedness and Response (ASPR) and Office of the Surgeon General (OSG) announce the opportunity for both private sector and non-profit entities to cosponsor an annual training summit. The focus of this training is medical and public health preparedness and response during disasters and emergencies. Potential cosponsors must have a mutual interest in the subject matter, the capability to provide logistical and educational support, and be willing to participate substantively in the cosponsored activity.

DATES: To receive consideration, a request to participate as a cosponsor must be received by the close of business on February 2, 2009. Requests will meet the deadline if they are either (1) received on or before the deadline