

produce the indexes and statistical summaries required by the Rule, and thus, estimated very low capital or start-up costs.

The only additional cost imposed on IDSMs operating under the Rule that would not be incurred for other IDSMs is the annual audit requirement. According to representatives of each of the IDSMs currently operating under the Rule, the vast majority of costs associated with this requirement are the fees paid to the auditors and their staffs to perform the annual audit. Representatives of the IDSMs estimated a combined cost of \$300,000 for both IDSMs currently operating under the Rule

**Other non-labor costs:** \$29,000 in copying costs. This total is based on estimated copying costs of 7 cents per page and several conservative assumptions. Staff estimates that the average dispute-related file is 35 pages long and that a typical annual audit file is approximately 200 pages in length. As discussed above, staff assumes that twenty percent of consumers using an IDSM currently operating under the Rule (approximately 4,896 consumers) request copies of the records relating to their disputes.

Staff also estimates that a very small minority of consumers request a copy of the annual audit. This assumption is based on (1) the number of consumer requests actually received by the IDSMs in the past; and (2) the fact that the IDSMs' annual audits are available online. For example, annual audits are available on the FTC's web site, where consumers may view and or print pages as needed, at no cost to the IDSM. In addition, the Better Business Bureau makes available on its web site the annual audit of the BBB AUTO LINE. Therefore, staff conservatively estimates that only five percent of consumers using an IDSM covered by the Rule (approximately 1,224 consumers) will request a copy of the IDSM's audit report.

Thus, the total annual copying cost for dispute-related files is approximately \$11,995 (35 pages per file x \$.07 per page x 4,896 consumer requests) and the total annual copying cost for annual audit reports is approximately \$17,136 (200 pages per audit report x \$.07 per page x 1,224 consumer requests). Accordingly, the total cost attributed to copying under the Rule is approximately \$29,131 and the total non-labor cost under the Rule is approximately \$329,131 (\$300,000 for

auditor fees + \$29,131 for copying costs).

**William Blumenthal**

*General Counsel*

[FR Doc. E7-21399 Filed 10-30-07; 8:45 am]

[Billing Code: 6750 - 01-S]

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## GENERAL SERVICES ADMINISTRATION

### Temporary Duty and Relocation Travel of Employees to Areas Impacted by the Wildfires in California

**AGENCY:** Office of Governmentwide Policy, General Services Administration (GSA).

**ACTION:** Notice of Federal Travel Regulation (FTR) Bulletin 08-02.

**SUMMARY:** The General Services Administration (GSA) has issued FTR Bulletin 08-02. FTR Bulletin 08-02 informs agencies that certain provisions of the FTR governing the authorization of actual subsistence expenses for official travel (both TDY and relocation) are temporarily waived as a result of the Emergency Declaration signed by the President on October 23, 2007, in response to wildfires in parts of California. It is expected that finding lodging facilities and/or adequate meals in the affected areas may be difficult, and distances involved may be great resulting in increased costs for per diem expenses. FTR Bulletin 08-02 became effective on October 24, 2007 and will remain effective until January 24, 2008, unless extended or rescinded by GSA. This bulletin and all FTR bulletins are located at [gsa.gov/bulletin](http://gsa.gov/bulletin).

**FOR FURTHER INFORMATION CONTACT:** For clarification of content, contact Patrick McConnell, Office of Governmentwide Policy, General Services Administration, Washington, DC 20405, telephone (202) 501-2362, or by email at [patrick.mcconnell@gsa.gov](mailto:patrick.mcconnell@gsa.gov).

Dated: October 25, 2007.

**Russ Pentz,**

*Assistant Deputy Associate Administrator.*

[FR Doc. E7-21393 Filed 10-30-07; 8:45 am]

**BILLING CODE 6820-14-S**

## GENERAL SERVICES ADMINISTRATION

### Premium Fuel Purchases for Government Owned and Leased Vehicles Due to Market Shortages in Parts of California Affected by Wildfires

**AGENCY:** Office of Governmentwide Policy, General Services Administration (GSA).

**ACTION:** Notice of Federal Management Regulation (FMR) Bulletin B-16.

**SUMMARY:** The General Services Administration (GSA) has issued Bulletin B-16 which provides a deviation for executive agencies to purchase premium fuel for Government owned and leased vehicles when lower grade fuels are not available due to market shortages in parts of California affected by wildfires. FMR Bulletin B-16 became effective on October 24, 2007 and will remain effective until January 24, 2008, unless extended or rescinded by GSA. This bulletin and all FMR bulletins are located at [gsa.gov/bulletin](http://gsa.gov/bulletin).

**FOR FURTHER INFORMATION CONTACT:** For clarification of content, contact Janet Dobbs, Office of Governmentwide Policy, General Services Administration, Washington, DC 20405, telephone (202) 208-6601, or by email at [janet.dobbs@gsa.gov](mailto:janet.dobbs@gsa.gov).

Dated: October 25, 2007.

**Russ Pentz,**

*Assistant Deputy Associate Administrator.*

[FR Doc. E7-21418 Filed 10-30-07; 8:45 am]

**BILLING CODE 6820-14-S**

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## DEPARTMENT OF HEALTH AND HUMAN SERVICES

### Centers for Disease Control and Prevention

[30Day-08-07AV]

#### Agency Forms Undergoing Paperwork Reduction Act Review

The Centers for Disease Control and Prevention (CDC) publishes a list of information collection requests under review by the Office of Management and Budget (OMB) in compliance with the Paperwork Reduction Act (44 U.S.C. Chapter 35). To request a copy of these requests, call the CDC Reports Clearance Officer at (404) 639-5960 or send an e-mail to [omb@cdc.gov](mailto:omb@cdc.gov). Send written comments to CDC Desk Officer, Office of Management and Budget, Washington, DC or by fax to (202) 395-6974. Written comments should be received within 30 days of this notice.

**Proposed Project**

Academic Centers of Excellence on Youth Violence Prevention Program Information System—New—National Center for Injury Prevention and Control (NCIPC), Centers for Disease Control and Prevention.

*Background and Brief Description*

Eight Academic Centers of Excellence on Youth Violence Prevention (ACEs) and two Urban Partnerships—Academic Centers of Excellence on Youth Violence Prevention (U-PACEs) are currently funded through CDC to foster and promote a stable, visible, long term strategy to address the complex problem of youth violence. The centers work with community members and many educational, justice and social work partners to develop action plans, partnerships, and priorities to prevent youth violence in a local community.

In addition, one ACE Coordinating Center is funded to initiate, foster, and support coordinated efforts, including the development and dissemination of activities and products in youth violence research and practice, among the ACEs, UPACEs, and CDC. It also aims to facilitate increased collaboration among organizations working to prevent youth violence to support the sustainability of youth violence prevention programs.

The Academic Centers of Excellence on Youth Violence Prevention Program Information System will collect, in electronic format: (a) Data needed to measure progress toward, or achievement of, performance indicators and other outcomes and (b) information on Academic Centers of Excellence on Youth Violence Prevention that is currently being collected in various electronic and paper documents. The clerical staff or Program Managers (n=11) will complete the majority of the

system. The principal investigators (n=11) will review the information in the system and add details related to study design and outcomes of projects, as necessary.

An Internet-based information system will allow CDC to monitor, and report on, ACE activities more efficiently. Data reported to CDC through the ACE information system will be used by CDC to identify training and technical assistance needs, monitor compliance with cooperative agreement requirements, evaluate the progress made in achieving center-specific goals, and obtain information needed to respond to Congressional and other inquiries regarding program activities and effectiveness.

There are no costs to respondents except their time to enter data into the Information System.

The total estimated annualized burden hours are 161.

**ESTIMATED ANNUALIZED BURDEN**

Respondents	Number of respondents	Number of responses per respondent	Average burden per response (in hrs.)
Clerical .....	11	2	320/60
Directors/Principal Investigators .....	11	2	120/60

Dated: October 23, 2007.

**Maryam I. Daneshvar,**

*Acting Reports Clearance Officer, Centers for Disease Control and Prevention.*

[FR Doc. E7-21415 Filed 10-30-07; 8:45 am]

**BILLING CODE 4163-18-P**

**DEPARTMENT OF HEALTH AND HUMAN SERVICES**

**Centers for Disease Control and Prevention**

[30Day-08-0199]

**Proposed Data Collections Submitted for Public Comment and Recommendations**

The Centers for Disease Control and Prevention (CDC) publishes a list of information collection requests under review by the Office of Management and Budget (OMB) in compliance with the Paperwork Reduction Act (44 U.S.C. Chapter 35). To request a copy of these requests, call the CDC Reports Clearance Officer at (404) 639-5960 or send an e-mail to *omb@cdc.gov*. Send written comments to CDC Desk Officer, Office of Management and Budget, Washington, DC or by fax to (202) 395-6974. Written

comments should be received within 30 days of this notice.

*Proposed Project:* Importation of Etiologic Agents, Hosts, and Vectors of Human Disease (42 CFR Part 71.54)—(OMB Control No. 0920-0199)—Extension—Office of the Director (OD), CDC. The Foreign Quarantine Regulations (42 CFR Part 71) set forth provisions to prevent the introduction, transmission, and spread of communicable disease from foreign countries into the United States. Subpart F—Importations—contains provisions for importation of etiologic agents, hosts, and vectors (42 CFR 71.54), requiring persons that import or distribute after importation these materials to obtain a permit issued by the CDC. This request is for the information collection requirements contained in 42 CFR 71.54 for issuance of permits by CDC to importers or distributors after importation of etiologic agents, hosts, or vectors of human disease.

CDC is requesting continued OMB approval to collect this information through the use of two separate forms. On an annual basis, approximately 2,300 laboratory facilities complete these forms to receive permits issued by

CDC. These forms are: (1) Application for Permit to Import or Transport Etiologic Agents, Hosts, or Vectors of Human Disease and (2) Application for Permit to Import or Transport Live Bats.

The Application for Permit to Import or Transport Etiologic Agents, Hosts, or Vectors of Human Disease will be used by laboratory facilities, such as those operated by government agencies, universities, research institutions, and zoologic exhibitions, and also by importers of nonhuman primate trophy materials, such as hunters or taxidermists, to request permits for the importation and subsequent distribution after importation of etiologic agents, hosts, or vectors of human disease. The Application for Permit to Import or Transport Etiologic Agents, Hosts, or Vectors of Human Disease requests applicant and sender contact information; description of material for importation; facility isolation and containment information; and personnel qualifications. Estimated average time to complete this form is 20 minutes.

The Application for Permit to Import or Transport Live Bats will be used by laboratory facilities such as those operated by government agencies, universities, research institutions, and