

should be submitted to: Ed Springer, GSA Desk Officer, OMB, Room 10236, NEOB, Washington, DC 20503, and a copy to the General Services Administration, Regulatory Secretariat, 1800 F Street, NW., Room 4035, Washington, DC 20405.

SUPPLEMENTARY INFORMATION:

A. Purpose

The General Services Administration (GSA) has various mission responsibilities related to the acquisition and provision of supplies, transportation, ADP, telecommunications, real property management, and disposal of real and personal property. These mission responsibilities generate requirements that are realized through the solicitation and award of public contracts. Individual solicitations and resulting contracts may impose unique information collection/reporting requirements on contractors, not required by regulation, but necessary to evaluate particular program accomplishments and measure success in meeting special program objectives.

B. Annual Reporting Burden

Respondents: 126,870.

Responses Per Respondent: 1.36.

Total Responses: 172,500.

Hours Per Response: .4.

Total Burden Hours: 68,900.

Obtaining Copies of Proposals

Requester may obtain a copy of the proposal from the General Services Administration, Regulatory Secretariat (MVP), 1800 F Street, NW., Room 4035, Washington, DC 20405, telephone (202) 501-4755. Please cite OMB Control No. 3090-0163, Information Specific to a Contract or Contracting Action (Not Required by Regulation), in all correspondence.

David A. Drabkin,

Deputy Associate Administrator, Office of Acquisition Policy.

[FR Doc. 01-11529 Filed 5-7-01; 8:45 am]

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**GENERAL SERVICES
ADMINISTRATION**

[OMB Control No. 3090-0040]

**Submission for OMB Review;
Comment Request Entitled Application
for Shipping Instructions and Notice of
Availability**

AGENCY: Federal Supply Service, GSA.

ACTION: Notice of request for public comments regarding extension of a

currently approved OMB clearance (3090-0040).

SUMMARY: Under the provisions of the Paperwork Reduction Act of 1995 (44 U.S.C. Chapter 35), the Office of Acquisition Policy has submitted to the Office of Management and Budget (OMB) a request to review and approve an extension of a currently approved information collection requirement concerning Application for Shipping Instructions and Notice of Availability. A request for public comments was published at 65 FR 70356, November 22, 2000. No comments were received.

DATES: Comment Due Date: June 7, 2001.

FOR FURTHER INFORMATION CONTACT:

Shirley Jacks, Acquisition Operations & Electronic Commerce Center, Supply Management Division, (703) 305-7473.

ADDRESSES: Comments regarding this burden estimate or any other aspect of this collection of information, including suggestions for reducing this burden, should be submitted to: Edward Springer, GSA Desk Officer, Room 3235, NEOB, Washington, DC 20503, and to Stephanie Morris, General Services Administration (MVP), 100 F Street NW., Washington, DC 20405.

SUPPLEMENTARY INFORMATION:

A. Purpose

The GSA is requesting the Office of Management and Budget (OMB) to reinstate information collection, 3090-0040, concerning Application for Shipping Instructions and Notice of Availability. This information collection supports and justifies the markup of the six percent surcharge for the GSA export reimbursable program. It also is used to evaluate and obtain the best cube utilization of shipping vans and containers for export direct delivery shipments. The form contains data necessary to prepare Transportation Control and Movement Documents (TCMD) which are required when material enters the Defense Transportation System.

B. Annual Reporting Burden

Respondents: 360.

Annual responses: 3,000.

Average hours per response: 20 minutes.

Burden hours: 1,000.

Obtaining Copies of Proposals

Requester may obtain a copy of the proposal from the General Services Administration, Acquisition Policy Division (MVP) Room 4035, 1800 F Street, NW., Washington, DC 20405, telephone (202) 208-7312. Please cite

OMB control number 3090-0040, concerning Application for Shipping Instructions and Notice of Availability, in all correspondence.

David A. Drabkin,

Deputy Associate Administrator, Office of Acquisition Policy.

[FR Doc. 01-11530 Filed 5-7-01; 8:45 am]

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**DEPARTMENT OF HEALTH AND
HUMAN SERVICES**

**Centers for Disease Control and
Prevention**

[60Day-01-35]

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

In compliance with the requirement of Section 3506(c)(2)(A) of the Paperwork Reduction Act of 1995 for opportunity for public comment on proposed data collection projects, the Centers for Disease Control and Prevention (CDC) will publish periodic summaries of proposed projects. To request more information on the proposed projects or to obtain a copy of the data collection plans and instruments, call the CDC Reports Clearance Officer on (404) 639-7090.

Comments are invited on: (a) Whether the proposed collection of information is necessary for the proper performance of the functions of the agency, including whether the information shall have practical utility; (b) the accuracy of the agency's estimate of the burden of the proposed collection of information; (c) ways to enhance the quality, utility, and clarity of the information to be collected; and (d) ways to minimize the burden of the collection of information on respondents, including through the use of automated collection techniques or other forms of information technology. Send comments to Anne O'Connor, CDC Assistant Reports Clearance Officer, 1600 Clifton Road, MS-D24, Atlanta, GA 30333. Written comments should be received within 60 days of this notice.

Proposed Project

Assess Reporting Practices for Q fever in each State, and Collect Retrospective Information on Numbers of Q fever Cases Reported between 1978-1999—New—National Center for Infectious Diseases (NCID), Centers for Disease Control and Prevention (CDC). Q fever, caused by the rickettsial agent *Coxiella burnetii*, is considered a potential agent of bioterrorism. Q fever in humans was

made nationally notifiable in the United States in 1999, and data are collected passively by CDC through the National Electronic Telecommunications System for Surveillance (NETSS). At the time Q fever was made nationally notifiable, approximately half of states already considered it a reportable disease. However, current information on how many states have changed their reporting criteria since 1999 or what reporting methods are used are not available, making it difficult to interpret data reported to NETSS. Q fever reporting data is available by state for the years 1948–1977, and complete

annual NETSS data is available beginning in the year 2000. However, data for the years 1978–1999 have never been collected. These data are vitally important to establish baseline rates of infection, providing immediate benefits in understanding the geographic distribution of Q fever in the United States. Without these data, it will take several years to acquire useful NETSS data to make the same assessments.

The purpose of this study is to assess reporting practices for Q fever in each state, and to collect retrospective information on numbers of Q fever cases reported between 1978–1999.

Respondents include an epidemiologist and laboratorian in every state. The results will be used to evaluate Q fever reporting practices in each state, in order to better assess the usefulness of NETSS-reported data. Data will also be used to examine the epidemiology of cases reported between 1979–1999, including estimated incidence rates, geographic distribution maps, and demographic risk factors. The estimated cost to the public is between \$3405–\$6405. Time burden for response to Form A may vary, depending on whether there are cases of Q fever the state plans to report.

Respondents	Number of respondents	Number of responses per respondent	Avg. burden per respondent in hours	Total burden in hours
Part A: Epidemiologist	50	1	30–120/60	25–100
Part B: Laboratorians	50	1	15/60	13
Total	38–113

Dated: May 1, 2001.

Nancy Cheal,

Acting Associate Director for Policy, Planning and Evaluation, Centers for Disease Control and Prevention.

[FR Doc. 01–11498 Filed 5–7–01; 8:45 am]

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DEPARTMENT OF HEALTH AND HUMAN SERVICES

Centers for Disease Control and Prevention

[Program Announcement 01073]

Fetal Alcohol Syndrome Awareness and Education Project; Notice of Availability of Funds

A. Purpose

The Centers for Disease Control and Prevention (CDC) announces the availability of fiscal year (FY) 2001 funds for a cooperative agreement program for a Fetal Alcohol Syndrome (FAS) Awareness and Education Project. This program addresses the “Healthy People 2010” focus areas of Substance Abuse, and Maternal, Infant, and Child Health. The purpose of the program is to develop, implement, and evaluate materials to educate parents and professionals working in public health service agencies, social service agencies, and school systems about FAS and how to secure access to appropriate diagnostic and treatment services for affected children and their families.

B. Eligible Applicants

Applications may be submitted by public and private nonprofit organizations and by governments and their agencies; that is, universities, colleges, research institutions, hospitals, other public and private nonprofit organizations, State and local governments or their bona fide agents including Puerto Rico, Virgin Islands, the Commonwealth of Northern Mariana Islands, American Samoa, Guam, the Federated States of Micronesia, the Republic of the Marshall Islands, and the Republic of Palau, and federally recognized Indian tribal governments, Indian tribes, or Indian tribal organizations.

Note: Title 2 of the United States Code, Chapter 26, Section 1611 states that an organization described in section 501(c)(4) of the Internal Revenue Code of 1986 that engages in lobbying activities is not eligible to receive Federal funds constituting an award, grant, cooperative agreement, contract, loan or any other form.

C. Availability of Funds

Approximately \$500,000 is available in FY 2001 to fund approximately 2 to 4 awards. It is expected that the average award will be \$100,000, ranging from \$75,000 to \$200,000. It is expected that the awards will begin on or about September 30, 2001, and will be made for a 12-month budget period within a project period of up to three years. Funding estimates may change.

Continuation awards within an approved project period will be made on the basis of satisfactory progress as

evidenced by required reports and the availability of funds.

D. Program Requirements

In conducting activities to achieve the purpose of this program, the recipient will be responsible for the activities under 1. (Recipient Activities), and CDC will be responsible for the activities listed under 2. (CDC Activities).

1. Recipient Activities

a. Develop, implement and evaluate an educational curricula for training parents, public health service providers, and/or teachers in primary and secondary schools that includes, but is not limited to, the following:

1. A description of FAS, including salient physical, neurodevelopmental, and behavioral characteristics of children with FAS.

2. An overview of other prenatal alcohol-related conditions and disorders.

3. Essential components of the care of children with FAS and other prenatal alcohol-related effects from the family, medical, social, and educational perspective including diagnosis and treatment services for the child and support services for the family.

4. Instructions for parents and other client-advocates in how to research local and State resource providers to gain assistance in identifying appropriate diagnostic, treatment, and educational services.

5. Descriptions of family stresses experienced, factors that increase the risks of family disintegration, and approaches to enhancing family coping and survival.