A. Purpose

The General Services Administration (GSA) has various mission responsibilities related to the acquisition and provision of Federal Supply Service's (FSS's) Stock, Special Order, and Schedules Programs. These mission responsibilities generate requirements that are realized through the solicitation and award of various types of FSS 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 and measure success in meeting program objectives.

B. Annual Reporting Burden

Respondents: 5380.
Responses for Respondent: 1.
Total Responses: 5380.
Hours Per Response: .25.
Total Burden Hours. 1, 345.

Obtaining Copies of Proposals

Requester may obtain a copy of the proposal from the General Services Administration, Acquisition Policy (MVP), 1800 F Street, NW., Room 4035, Washington, DC 20405, telephone (202) 501–4744. Please cite OMB Control No. 3090–0248, Placement of Orders and Ordering Information, in all correspondence.

Dated: April 15, 2002.

Al Matera,

Director, Acquisition Policy Division.

[FR Doc. 02–9933 Filed 4–22–02; 8:45 am]

BILLING CODE 6820–34–M

DEPARTMENT OF HEALTH AND HUMAN SERVICES

Office of the Secretary

Agency Information Collection Activities; Proposed Collections; Comment Request

The Department of Health and Human Services, Office of the Secretary will periodically publish summaries of proposed information collections projects and solicit public comments in compliance with the requirements of Section 3506(c)(2)(A) of the Paperwork Reduction Act of 1995. To request more information on the project or to obtain a copy of the information collection plans and instruments, call the OS Reports Clearance Office at (202) 619–2118 or e-mail *Geerie.Jones@HHS.gov.*

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.

Proposed Project: ''National Study of Culturally and Linguistically Appropriate Services in Local Public Health Agencies''—New—The Office of Minority Health proposes to conduct a survey with a national sample of local health departments serving racially and ethnically diverse communities. The survey will provide data on the types of policies and practices that promote the delivery of culturally and linguistically appropriate services by local health departments, and the factors that facilitate and detract from the implementation of such policies and practices. The data collected will inform the Office of Minority Health about the current nature and extent of such services.

Respondents: Business or other for-profit, Non-profit organizations;
Number of Respondents: 150;
Response per Respondent: 3;
Average Burden per Response: 30 minutes;
Total Burden: 225 hours.

Send comments via e-mail to *Geerie.Jones@HHS.gov.* or mail to OS Reports Clearance Office, Room 503H, Hubert H. Humphrey Building, 200 Independence Avenue SW., Washington DC, 20201. Comments should be received within 60 days of this notice.

Dated: April 15, 2002.

Kerry Weems,

Acting, Deputy Assistant Secretary, Budget.

[FR Doc. 02–9873 Filed 4–22–02; 8:45 am]

BILLING CODE 4150–29–M

DEPARTMENT OF HEALTH AND HUMAN SERVICES

National Committee on Vital and Health Statistics: Meeting

Pursuant to the Federal Advisory Committee Act, the Department of Health and Human Services announces the following advisory committee meeting.

Name: National Committee on Vital and Health Statistics (NCVHS), HHS.

Time and Date: 12 noon–1 p.m. EDT—April 24, 2002.

Place: Conference Call, Participants' Information to be Announced.

Status: Open.

Purpose: During this telephone conference call, the Committee will discuss its comments to the Department on the Current Notice of Proposed Rule Making Covering proposed changes to the Health Insurance Portability and Accountability Act of 1996 (HIPAA) Privacy Rule.

Notice: This conference call is open to the public using a participants' dial-in telephone number and participants' code, but access may be limited by the number of available telephone lines. The number and code will be announced on the NCVHS website *http://www.ncvhs.hhs.gov/.*

CONTACT PERSON FOR MORE INFORMATION: Substantive program information as well as summaries of meetings and a roster of committee members may be obtained from Majorie S. Greenberg, Executive Secretary, NCVHS, National Center for Health Statistics, Centers for Disease Control and Prevention, Room 1100, Presidential Building, 6525 Belcrest Road, Hyattsville, Maryland 20782, telephone (301) 458–4245. Information also is available on the NCVHS home page of the HHS website: *http://www.ncvhs.hhs.gov/.*

Dated: April 17, 2002.

James Scanlon,

Director, Division of Data Policy, Office of the Assistant Secretary for Planning and Evaluation.

[FR Doc. 02–9874 Filed 4–22–02; 8:45 am]

BILLING CODE 4151–01–M

DEPARTMENT OF HEALTH AND HUMAN SERVICES

Centers for Disease Control and Prevention

[60Day–02–43]

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. CDC is requesting an emergency clearance from the Office of Management and Budget (OMB) to collect data under the Fertility Clinic Success Rate and Certification Act (FCSRCA) of 1992. Send comments to Anne O'Connor, CDC Assistant Reports Clearance Officer, 1600 Clifton

Road, MS D–24, Atlanta, GA 30333. Written comments should be received within 14 days of this notice. OMB is expected to act on the request of CDC within 21 days of publication of this notice.

Proposed Project

Assisted Reproductive Technology (ART) Program Reporting System—New—National Center for Chronic Disease Prevention and Health Promotion (NCCDPHP), Centers for Disease Control and Prevention (CDC).

Background: Section 2(a) of Pub. L. 102–493 (known as the Fertility Clinic Success Rate and Certification Act of 1992 (FCSRCA), 42 U.S.C. 263a–1(a)) requires that each assisted reproductive technology (ART) program shall annually report to the Secretary through the Centers for Disease Control and Prevention—(1) pregnancy success rates achieved by such ART program, and (2) the identity of each embryo laboratory used by such ART program and whether the laboratory is certified or has applied for such certification under this act.

The Centers for Disease Control and Prevention (CDC) is seeking approval of a reporting system for Assisted Reproductive Technology (ART)Program from the Office of Management and Budget (OMB). This reporting system has been designed in collaboration with the Society for Reproductive Technology to comply with the requirements of the FCSRCA. The reporting system includes all ART cycles initiated by any of the approximately 400 ART programs in the United States, and covers the pregnancy outcome of each cycle, as well as a number of data items deemed important to explain variability in success rates across clinics and across individuals. Data is to be collected through computer software developed by SART in consultation with CDC.

In developing the definition of pregnancy success rates and the list of data items to be reported, CDC has consulted with representatives of SART, the American Society for Reproductive Medicine, and RESOLVE, the National Infertility Association (a national, nonprofit consumer organization), as well as a variety of individuals with expertise and interest in this field. The average annual cost to the respondent, including data entry labor and fees, is estimated to be $2,140.

Respondents	Number of respondents	Number of responses/respondent	Average burden/response (in hours)	Total burden (in hours)
ART Clinics	400	220	5/60	7,333
Total				7,333

Dated: April 16, 2002.

Nancy E. Cheal,

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

[FR Doc. 02–9843 Filed 4–22–02; 8:45 am]

BILLING CODE 4163–18–P

DEPARTMENT OF HEALTH AND HUMAN SERVICES

Food and Drug Administration

Advisory Committees; Filing of Annual Reports

AGENCY: Food and Drug Administration, HHS.

ACTION: Notice.

SUMMARY: The Food and Drug Administration (FDA) is announcing that, as required by the Federal Advisory Committee Act, the agency has filed with the Library of Congress the annual reports of those FDA advisory committees that held closed meetings.

ADDRESSES: Copies of the annual reports are available from the Dockets Management Branch (HFA–305), Food and Drug Administration, 5630 Fishers Lane, rm. 1061, Rockville, MD 20852, 301–827–6860.

FOR FURTHER INFORMATION CONTACT: Linda Ann Sherman, Advisory Committee Oversight and Management Staff (HF–4), Food and Drug Administration, 5600 Fishers Lane, Rockville, MD 20857, 301–827–1220.

SUPPLEMENTARY INFORMATION: Under section 13 of the Federal Advisory Committee Act (5 U.S.C. App. 2) and 21 CFR 14.60 (c), FDA has filed with the Library of Congress the annual reports for the following FDA advisory committees that held closed meetings during the period October 1, 1998, through September 30, 1999:

Center for Biologics Evaluation and Research:
- Allergenic Products Advisory Committee,
- Biological Response Modifiers Advisory Committee, and
- Vaccines and Related Biological Products Advisory Committee.

Center for Drug Evaluation and Research:
- Antiviral Drugs Advisory Committee,
- Arthritis Advisory Committee,
- Dermatologic and Ophthalmic Drugs Advisory Committee,
- Drug Abuse Advisory Committee, and
- Oncologic Drugs Advisory Committee.

Center for Devices and Radiological Health:
- Medical Devices Advisory Committee.

National Center for Toxicological Research:
- Science Advisory Board to the National Center for Toxicological Research.
- Science Board to the Food and Drug Administration.

Annual reports have also been filed for the following FDA advisory committees that held closed meetings during the period October 1, 1999, through September 30, 2000:

Center for Biologics Evaluation and Research:
- Biological Response Modifiers Advisory Committee,
- Blood Products Advisory Committee,
- Transmissible Spongiform Encephalopathies Advisory Committee, and
- Vaccines and Related Biological Products Advisory Committee.

Center for Drug Evaluation and Research:
- Antiviral Drugs Advisory Committee,
- Arthritis Advisory Committee, and
- Dermatologic and Ophthalmic Drugs Advisory Committee.

Center for Devices and Radiological Health:
- Medical Devices Advisory Committee.

National Center for Toxicological Research:
- Science Advisory Board to the National Center for Toxicological Research.

Annual reports are available for public inspection at: (1) The Library of Congress, Madison Bldg., Newspaper and Current Periodical Reading Room, 101 Independence Ave. SE., rm. 133, Washington, DC; and (2) the Dockets Management Branch (HFA–305), Food and Drug Administration, 5630 Fishers Lane, rm. 1061, Rockville, MD 20852, between 9 a.m. and 4 p.m., Monday through Friday.