

TABLE I—Continued

| Populations or areas to be served           | Number of competing grants to be awarded | FY99 funding | Appl. due date | Grant funding date |
|---|--|--------------|----------------|--------------------|
| Republic of Palau .....                     | 1  | 57,971       | 03/01/00       | 07/01/00           |
| Federated States of Micronesia .....        | 1  | 220,564      | 03/01/00       | 07/01/00           |
| Gila River, Arizona .....                   | 1  | 172,582      | 03/01/00       | 07/01/00           |
| Region X:                                   |  |              |                |                    |
| Columbia, Willamette Counties, Oregon ..... | 1  | 561,485      | 03/01/00       | 07/01/00           |
| Idaho .....                                 | 1  | 961,979      | 03/01/00       | 07/01/00           |
| Seattle, Washington .....                   | 1  | 123,800      | 05/30/00       | 09/30/00           |
| Total .....                                 | 32                                       | 83,748,838   | .....          | .....              |

Dated: August 27, 1999.

**Denese O. Shervington,**

*Deputy Assistant Secretary for Population Affairs.*

[FR Doc. 99-22819 Filed 9-1-99; 8:45 am]

BILLING CODE 4160-17-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).

*Times and Dates:* 9:00 a.m.–5:30 p.m., September 27, 1999; 10:15 a.m.–3:30 p.m., September 28, 1999.

*Place:* Conference Room 705A, Hubert H. Humphrey Building, 200 Independence Avenue SW, Washington, DC 20201.

*Status:* Open.

*Purpose:* The meeting will focus on a variety of health data policy and privacy issues. Department officials will update the Committee on recent activities of the HHS Data Council and the status of HHS activities in implementing the administrative simplification provisions of Pub. L. 104-191, the Health Insurance Portability and Accountability Act of 1996 (HIPAA). The Committee also will discuss its forthcoming report on Medicaid managed care data issues, as well as a report to the Secretary concerning standards for computer-based patient records, and plans for a November, 1999 workshop at the National Academy of Sciences on Health Statistics for the 21st Century. The Committee also will receive briefings on quality of care data issues and the revision of the U.S. Standard Certificates for Live Birth and Death for 2002. In addition, Subcommittee breakout sessions and reports to the full Committee are planned.

All topics are tentative and subject to change. Prior to the meeting, please check the NCVHS web site, where a detailed agenda will be posted when available.

*Contact Person for More Information:*

Substantive information as well as summaries of NCVHS meetings and a roster of committee members may be obtained by visiting the NCVHS website (<http://aspe.os.dhhs.gov/ncvhs>) where an agenda for the meeting will be posted when available. Additional information may be obtained by calling James Scanlon, NCVHS Executive Staff Director, Office of the Assistant Secretary for Planning and Evaluation, DHHS, Room 440-D, Humphrey Building, 200 Independence Avenue SW., Washington, DC 20201, telephone (202) 690-7100, or Marjorie S. Greenberg, Executive Secretary, NCVHS, NCHS, CDC, Room 1100, Presidential Building, 6525 Belcrest Road, Hyattsville, Maryland 20782, telephone 301/436-7050.

**Note:** In the interest of security, the Department has instituted stringent procedures for entrance to the Hubert H. Humphrey Building by non-government employees. Thus, individuals without a government identification card may need to have the guard call for an escort to the meeting room.

Dated: August 26, 1999.

**James Scanlon,**

*Director, Division of Data Policy.*

[FR Doc. 99-22821 Filed 9-1-99; 8:45 am]

BILLING CODE 4151-04-M

**DEPARTMENT OF HEALTH AND HUMAN SERVICES**

**Centers for Disease Control and Prevention**

[INFO-99-28]

**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 for other forms of information technology. Send comments to Seleda Perryman, 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**

Translating Research Into Action for Diabetes (TRIAD)—New—The National Center for Chronic Disease Prevention and Health Promotion (NCCDPHP), Division of Diabetes Translation. Diabetes exerts a huge public health burden, and there are several efficacious interventions to combat the effects of this disease. Yet, the quality of care and quality of life among people with diabetes remain sub-optimal in the United States. The Centers for Disease Control and Prevention, Division of Diabetes Translation, intends to conduct a multi-center study called Translating Research Into Action for Diabetes (TRIAD). This study will assess quality of diabetes care and identify effective means of applying existing knowledge to improve care and quality of life. Data provided from TRIAD will be critical to the Division of Diabetes Translation's ongoing efforts to reduce the burden of diabetes.

Data will be collected through cooperative agreements with six