

committee trade secret and/or confidential commercial information (5 U.S.C. 552b(c)(4)) regarding pending and future agency issues.

Persons attending FDA's advisory committee meetings are advised that the agency is not responsible for providing access to electrical outlets.

FDA welcomes the attendance of the public at its advisory committee meetings and will make every effort to accommodate persons with physical disabilities or special needs. If you require special accommodations due to a disability, please contact Shirley Meeks, Conference Management Staff, at 301-594-1283, ext. 105, at least 7 days in advance of the meeting.

Notice of this meeting is given under the Federal Advisory Committee Act (5 U.S.C. app. 2).

Dated: September 14, 2002.  
**Linda Arey Skladany,**  
*Senior Associate Commissioner for External Relations.*  
[FR Doc. 02-23949 Filed 9-19-02; 8:45 am]  
**BILLING CODE 4160-01-S**

**DEPARTMENT OF HEALTH AND HUMAN SERVICES**

**Health Resources and Services Administration**

**Agency Information Collection Activities: Submission for OMB Review; Comment Request**

Periodically, the Health Resources and Services Administration (HRSA) publishes abstracts of information collection requests under review by the Office of Management and Budget, in compliance with the Paperwork Reduction Act of 1995 (44 U.S.C. Chapter 35). To request a copy of the clearance requests submitted to OMB for review, call the HRSA Reports Clearance Office on (301) 443-1129.

The following request has been submitted to the Office of Management and Budget for review under the Paperwork Reduction Act of 1995:

**Proposed Project: The Sentinel Centers Network (SCN) Core Data Set—New**

HRSA's Bureau of Primary Health Care (BPHC) established the Sentinel Centers Network (SCN) to assist in addressing critical policy issues. Thirty-eight BPHC supported health centers and NHSC sites have been awarded

funds through sub-contracts in this first year of operation. These health centers were identified as having adequate infrastructure and commitment through the competitive contract process to serve as "laboratories" that will generate data for timely policy analyses and conducting projects on topics that have immediate policy impact.

A protocol for core data collection and retrieval, timelines, expectations, and evaluation of the Network sites is currently underway. It is expected that sites will submit these core data, or have these data extracted from their existing information systems periodically. These core data may include provider level, encounter level, and user level information regarding, for example, data on service delivery, utilization, payer sources, demographics, clinical diagnoses and outcomes, staffing, and costs. Since all data obtained from the participant sites will be extracted/compiled from existing information systems, and not through primary data collection, burden will therefore be minimized. In addition, each participant site will receive technical assistance both on site and via telephone to reduce burden as much as possible.

Estimated burden hours:

Type of respondent	Number of respondents	Responses per respondent	Total responses	Hours per responses	Total burden hours
Sites .....	38	4	152	8	1,216

Written comments and recommendations concerning the proposed information collection should be sent within 30 days of this notice to: John Morrall, Human Resources and Housing Branch, Office of Management and Budget, New Executive Office Building, Room 10235, Washington, DC 20503.

Dated: September 12, 2002.  
**Jane M. Harrison,**  
*Director, Division of Policy Review and Coordination.*  
[FR Doc. 02-23848 Filed 9-19-02; 8:45 am]  
**BILLING CODE 4165-15-P**

**DEPARTMENT OF HEALTH AND HUMAN SERVICES**

**National Institutes of Health**

**National Heart, Lung, and Blood Institute; Proposed Collection; Comment Request; Women's Health Initiative Observational Study**

*Summary:* 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, Office of the Director, the National Heart, Lung, and Blood Institute (NHLBI), the National Institutes of Health (NIH) will publish periodic summaries of proposed projects to be submitted to the Office of Management and Budget (OMB) for review and approval.

**Proposed Collection**

*Title:* Women's Health Initiative (WHI) Observational Study.  
*Type of Information Collection Request:* Revision OMB #0925-0414  
*Exp:* 05/03.

*Need for Use of Information Collection:* This study will be used by the NIH to evaluate risk factors for chronic disease among older women by developing and following a large cohort of postmenopausal women and relating subsequent disease development to baseline assessments of historical, physical, psychosocial, and physiologic characteristics. In addition, the observational study will complement the clinical trial (which has received clinical exemption) and provide additional information on the common causes of frailty, disability and death for postmenopausal women, namely, coronary heart disease, breast and colorectal cancer, and osteoporotic fractures.

*Frequency of Response:* On occasion.

*Affected Public:* Individuals and physicians.

*Type of Respondents:* Women, next-of-kin, and physician's office staff. The annual reporting burden is as follows: