Item Number	Title of Standard	Reference Number and Date
51	Using Proficiency Testing (PT) to Improve the Clinical Laboratory; Approved Guideline	NCCLS GP27–A
52	Terminology and Definition for Use in NCCLS Documents; Approved Standard	NRSCL 8–A
53	Continuous Quality Improvement: Essential Management Approaches; Approved Guideline	NCCLS GP22–A
	OB–GYN/Gastroenterology	
16	Enteral Feeding Set Connectors and Adapters	ANSI/AAMI ID54 (1996)
17	Standard Specifications for Rubber Contraceptives (Male Condoms)	ASTM D3492–97
18	Electrosurgical Device	ANSI/AAMI HF–18 (1993)
	Ophthalmic	
14	Ophthalmics—Contact Lenses—Standard Terminology, Tolerances, Measurements, and Physicochemical Properties	ANSI Z80.20–1998
	Radiology	
44	Acoustic Output Measurement Standard for Diagnostic Ultrasound Equipment	AIUM (1998)
45	Standard for Real-Time Display of Thermal andMechanical Acoustic Output Indices on Diagnostic Ultrasound Equipment. Revision 1.	AIUM RTD (1998)
46	Acoustic Output Labeling Standard for Diagnostic Ultrasound Equipment: A Standard for How Manufacturers Should Specify Acoustic Output Data	AIUM AOL (1998)
47	Medical Electrical Equipment: Radionuclide Calibrators *B* Particular Methods for Describing Performance	IEC 61303 (1994–10)
48	Calibration and Usage of ''Dose Calibrator'' Ionization Chambers for the Assay of Radionuclides	ANSI N42.13 (1986)
49	Calibration and Usage of Ionization Chamber Systems for Assay of Radionuclides	IEC 61145 (1992–05)
	Software	
2	Standard for Developing Software Life Cycle Processes	IEEE 1074 (1997)
3	Industry Implementation of International Standard ISO/IEC 12207: 1995 (ISO/IEC 12207) Standard for Information Technology—Software Life Cycle Processes	IEEE/EIA 12207.0 (1996)
	Sterility	
37	Biological Evaluation of Medical Devices—Part 7: Ethylene Oxide Sterilization Residuals	ANSI/AAMI/ISO 10993–7 (1995)

Dated: June 30, 1999.

Linda S. Kahan,

Deputy Director for Regulations Policy.

[FR Doc. 99–17429 Filed 7–9–99; 8:45 am]

BILLING CODE 4160–01–F

DEPARTMENT OF HEALTH AND HUMAN SERVICES

Health Care Financing Administration

[Document Identifier: HCFA–R–289]

Emergency Clearance: Public Information Collection Requirements Submitted to the Office of Management and Budget (OMB)

AGENCY: Health Care Financing Administration, HHS.

In compliance with the requirement of section 3506(c)(2)(A) of the Paperwork Reduction Act of 1995, the Health Care Financing Administration (HCFA), Department of Health and Human Services, is publishing the following summary of proposed collections for public comment. Interested persons are invited to send comments regarding this burden estimate or any other aspect of this collection of information, including any of the following subjects: (1) The necessity and utility of the proposed information collection for the proper performance of the agency's functions; (2) the accuracy of the estimated burden; (3) ways to enhance the quality, utility, and clarity of the information to be collected; and (4) the use of automated collection techniques or other forms of information technology to

minimize the information collection burden.

We are, however, requesting an emergency review of the Information collections referenced below. In compliance with the requirement of section 3506(c)(2)(A) of the Paperwork Reduction Act of 1995, we have submitted to the Office of Management and Budget (OMB) the following requirements for emergency review. We are requesting an emergency review because of the unanticipated urgency to meet the enrollment cycle of the program, thereby reducing the burden on the demonstration sites. HCFA is supporting the demonstration within the current fiscal year, and as an agency priority. In addition, public harm may occur as the result of not evaluating and possibly providing alternative health care measures outlined in this demonstration.

HCFA is requesting OMB review and approval of this collection within three days, with a 180-day approval period. Written comments and recommendations will be accepted from the public if received by the individuals designated below within three days. During this 180-day period, we will publish a separate **Federal Register** notice announcing the initiation of an extensive 60-day agency review and public comment period on these requirements. We will submit the requirements for OMB review and an extension of this emergency approval.

Type of Information Collection Request: New Collection;

Title of Information Collection: Medicare Lifestyle Modification Program Demonstration;

Form No.: HCFA–R–289;

Use: The Health Care Financing Administration (HCFA) through its Office of Clinical Standards and Quality (OCSQ) is planning to conduct a new demonstration to test the feasibility and cost effectiveness of cardiovascular lifestyle modification. This demonstration will focus on Medicare provider sponsored, lifestyle modification programs designed to reverse, reduce, or ameliorate the indications of cardiovascular disease (CAD) of Medicare beneficiaries at risk for invasive treatment procedures. This demonstration will test the feasibility and cost effectiveness of providing payment for cardiovascular lifestyle modification program services to Medicare beneficiaries. In addition, the demonstration will test the use of contractual agreements for administration, claims processing and payment, and routine monitoring of quality of care.

Frequency: On occasion, Monthly, and Quarterly;

Affected Public: Individuals or Households, and Not-for-profit institutions;

Number of Respondents: 12;

Total Annual Responses: 4,500;

Total Annual Hours: 750.

To obtain copies of the supporting statement and any related forms for the proposed paperwork collections referenced above, access HCFA's Web Site address at http://www.hcfa.gov/regs/prdact95.htm, or E-mail your request, including your address, phone number, to Paperwork@hcfa.gov, or call the Reports Clearance Office on (410) 786–1326.

Interested persons are invited to send comments regarding the burden or any other aspect of these collections of Information requirements. However, as noted above, comments on these Information collection and recordkeeping requirements must be mailed and/or faxed to the designees referenced below, within three days:

Health Care Financing Administration, Office of Information Services, Security and Standards Group, Division of HCFA Enterprise Standards, Attention: Dawn Willinghan, Room N2–14–26, 7500 Security Boulevard, Baltimore, Maryland 21244–1850

and

Office of Information and Regulatory Affairs, Office of Management and Budget, Room 10235, New Executive Office Building, Washington, DC 20503, Fax Number: (202) 395–6974 or (202) 395–5167. Attn: Allison Herron Eydt, HCFA Desk Officer.

Dated: July 1, 1999.

John P. Burke III,

HCFA Reports Clearance Officer, HCFA Office of Information Services, Security and Standards Group, Division of HCFA Enterprise Standards.

[FR Doc. 99–17559 Filed 7–9–99; 8:45 am]

BILLING CODE 4120–03–P

DEPARTMENT OF HEALTH AND HUMAN SERVICES

National Institutes of Health

National Center for Complementary and Alternative Medicine; Notice of Closed Meeting

Pursuant to section 10(d) of the Federal Advisory Committee Act, as amended (5 U.S.C. Appendix 2), notice is hereby given of the following meeting.

The meeting will be closed to the public in accordance with the provisions set forth in sections 552b(c)(4) and 552b(c)(6), Title 5 U.S.C., as amended. The grant applications and the discussions could disclose confidential trade secrets or commercial property such as patentable material, and personal information concerning individuals associated with the grant applications, the disclosure of which would constitute a clearly unwarranted invasion of personal privacy.

Name of Committee: National Center for Complementary and Alternative Medicine Special Emphasis Panel.

Date: July 22, 1999.

Time: 8:30 am to 5 pm

Agenda: To review and evaluate cooperative agreement applications.

Place: Marriott Pooks Hill, 5151 Pooks Hill Road, Bethesda, MD 20814.

Contact Person: Eugene G. Hayunga, Ph.D., Scientific Review Administrator, National Institutes of Health, NCCAM, Building 31, Room 5B50, 9000 Rockville Pike, Bethesda, MD 20892, 301–594–2014, hayungae@od.nih.gov.

This notice is being published less than 15 days prior to the meeting due to the timing limitations imposed by the review and funding cycle.

Dated: July 6, 1999.

LaVerne Y. Stringfield,

Director, Office of Federal Advisory Committee Policy.

[FR Doc. 99–17592 Filed 7–9–99; 8:45 am]

BILLING CODE 4140–01–M

DEPARTMENT OF HEALTH AND HUMAN SERVICES

National Institutes of Health

National Institute on Drug Abuse; Notice of Closed Meetings

Pursuant to section 10(d) of the Federal Advisory Committee Act, as amended (5 U.S.C. Appendix 2), notice is hereby given of the following meetings.

The meetings will be closed to the public in accordance with the provisions set forth in sections 552b(c)(4) and 552b(c)(6), Title 5 U.S.C., as amended. The contract proposals and the discussions could disclose confidential trade secrets or commercial property such as patentable material, and personal information concerning individuals associated with the contract proposals, the disclosure of which would constitute a clearly unwarranted invasion of personal privacy.

Name of Committee: National Institute on Drug Abuse Special Emphasis Panel, ''Clinical Data Management and Support Center.''

Date: July 13, 1999.

Time: 9:00 am to 5:00 pm.