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

Status: Open.

Purpose: The agenda for Tuesday, January 27th, will be devoted to final reports on standards for five domains that were prepared as part of the Consolidated Health Informatics Initiative (CHI) and a related letter on CHI recommendations to the Secretary will be finalized. The afternoon will focus on issues related to the implementation of the HIPAA Security Rule.

The morning of the second day will include an update on implementation of HIPAA transactions and code sets provisions; the development of a draft letter to the Secretary concerning the Claims Attachment Standard; and a session on dental data standards issues. The afternoon will be devoted to Subcommittee planning of future activities around E-prescriptions.

FOR FURTHER INFORMATION CONTACT: Substantive program information as well as summaries of meetings and a roster of Committee members may be obtained from Maria Friedman, Health Insurance Specialist, Security and Standards Group, Centers for Medical and Medicaid Services, MS: C5–24–04, 7500 Security Boulevard, Baltimore, MD 21244–1850, telephone: (410) 786–6333 or Marjorie S. Greenberg, Executive Secretary, NCVHS, National Center for Health Statistics, Centers for Disease Control and Prevention, Room 1100, Presidential Building, 3311 Toledo Road, Hyattsville, Maryland 20782, telephone: (301) 458–4245. Information also is available on the NCVHS home page of the HHS Web site: *http://www.ncvhs.hhs.gov/* where an agenda for the meeting will be posted when available. Should you require reasonable accommodation, please contact the CDC Office of Equal Employment Opportunity on (301) 458–4EEO (4336) as soon as possible.

Dated: December 24, 2003.

James Scanlon,

Acting Deputy Assistant Secretary for Science and Data Policy, Office of the Assistant Secretary for Planning and Evaluation.

[FR Doc. 04–396 Filed 1–8–04; 8:45 am]

BILLING CODE 4151–05–M

DEPARTMENT OF HEALTH AND HUMAN SERVICES

Administration on Aging

Agency Information Collection Activities; Proposed Collection; Comment Request; AoA Nutrition and Physical Activity Campaign

AGENCY: Administration on Aging, HHS.

ACTION: Notice.

SUMMARY: The Administration on Aging (AoA) is announcing an opportunity for public comment on the proposed collection of certain information by the agency. Under the Paperwork Reduction Act of 1995 (the PRA), Federal agencies are required to publish notice in the **Federal Register** concerning each proposed collection of information, including each proposed extension of an existing collection of information, and to allow 60 days for public comment in response to the notice. This notice solicits comments on the information collection requirements relating to organizations that wish to enroll as a partner with AoA in a campaign to create awareness and make nutrition and physical activity programs available to older Americans. The requested information includes providing general information about the entity, its programs, and counts of populations served.

DATES: Submit written or electronic comments on the collection of information by March 9, 2004.

ADDRESSES: Submit electronic comments on the collection of information to: *Kathleen.Loughrey@aoa.gov.*

Submit written comments on the collection of information to Administration on Aging, Washington, DC 20201.

FOR FURTHER INFORMATION CONTACT: Kathleen Loughrey, U.S. Department of Health and Human Services, Administration on Aging, Washington, DC 20201.

SUPPLEMENTARY INFORMATION: Under the PRA (44 U.S.C. 3501–3520), Federal agencies must obtain approval from the Office of Management and Budget (OMB) for each collection of information they conduct or sponsor. ''Collection of information'' is defined in 44 U.S.C. 3502(3) and 5 CFR 1320.3(c) and includes agency request or requirements that members of the public submit reports, keep records, or provide information to a third party. Section 3506(c)(2)(A) of the PRA (44 U.S.C. 3506(c)(2)(A)) requires Federal agencies to provide a 60-day notice in the **Federal Register** concerning each proposed collection of information, including each proposed extension of an existing collection of information, before submitting the collection to OMB for approval. To comply with this requirement, AoA is publishing notice of the proposed collection of information set forth in this document. With respect to the following collection of information, AoA invites comments on: (1) Whether the proposed collection of information is necessary for the proper performance of AoA's functions, including whether the information will have practical utility; (2) the accuracy of AoA's estimate of the burden of the proposed collection of information, including the validity of the methodology and assumptions used; (3) ways to enhance the quality, utility, and clarity of the information to be collected; and (4) ways to minimize the burden of the collection of information on respondents, including through the use of automated collection techniques when appropriate, and other forms of information technology.

Describe Collection of Information

AoA estimates the burden of this collection of information as follows: AoA estimates a total of no more than 500 hours will be required to collect this information. This estimate is based on these assumptions: AoA estimates that 2,000 organizations will complete an entry form to become a campaign partner. Completion of each entry form will require a total of 15 minutes per organization including five minutes to answer questions, five minutes to insert a program description, and five minutes to look up data from existing program records.

Dated: January 2, 2004.

Josefina G. Carbonell,

Assistant Secretary for Aging.

[FR Doc. 04–471 Filed 1–8–04; 8:45 am]

BILLING CODE 4154–01–P

DEPARTMENT OF HEALTH AND HUMAN SERVICES

Food and Drug Administration

[Docket No. 2003N–0311]

Agency Information Collection Activities; Announcement of Office of Management and Budget Approval; Medical Device User Fee and Modernization Act Small Business Qualification Certification (Form FDA 3602)

AGENCY: Food and Drug Administration, HHS.

ACTION: Notice.

SUMMARY: The Food and Drug Administration (FDA) is announcing that a collection of information entitled ''Medical Device User Fee and Modernization Act Small Business Qualification Certification (Form FDA 3602)'' has been approved by the Office of Management and Budget (OMB) under the Paperwork Reduction Act of 1995.

FOR FURTHER INFORMATION CONTACT: Peggy Robbins, Office of Management Programs (HFA–250), Food and Drug Administration, 5600 Fishers Lane, Rockville, MD 20857, 301–827–1223.

SUPPLEMENTARY INFORMATION: In the **Federal Register** of October 10, 2003 (68