imported into the United States under 21 U.S.C. 42.

Agenda—Open public hearing. Interested persons may present data, information, or views, orally or in writing, on issues pending before the committee.

Open board discussion. The board will discuss and select tea standards.

FDA public advisory committee meetings may have as many as four separable portions: (1) An open public hearing, (2) an open committee discussion, (3) a closed presentation of data, and (4) a closed committee deliberation. Every advisory committee meeting shall have an open public hearing portion. Whether or not it also includes any of the other three portions will depend upon the specific meeting involved. There are no closed portions for the meetings announced in this notice. The dates and times reserved for the open portions of each committee meeting are listed above.

The open public hearing portion of each meeting shall be at least 1 hour long unless public participation does not last that long. It is emphasized, however, that the 1 hour time limit for an open public hearing represents a minimum rather than a maximum time for public participation, and an open public hearing may last for whatever longer period the committee chairperson determines will facilitate the committee's work.

Public hearings are subject to FDA's guideline (subpart C of 21 CFR part 10) concerning the policy and procedures for electronic media coverage of FDA's public administrative proceedings, including hearings before public advisory committees under 21 CFR part 14. Under 21 CFR 10.205, representatives of the electronic media may be permitted, subject to certain limitations, to videotape, film, or otherwise record FDA's public administrative proceedings, including presentations by participants.

Meetings of advisory committees shall be conducted, insofar as is practical, in accordance with the agenda published in this **Federal Register** notice. Changes in the agenda will be announced at the beginning of the open portion of a meeting.

Any interested person who wishes to be assured of the right to make an oral presentation at the open public hearing portion of a meeting shall inform the contact person listed above, either orally or in writing, prior to the meeting. Any person attending the hearing who does not in advance of the meeting request an opportunity to speak will be allowed to make an oral presentation at the

hearing's conclusion, if time permits, at the chairperson's discretion.

The agenda, the questions to be addressed by the committee, and a current list of committee members will be available at the meeting location on the day of the meeting.

Transcripts of the open portion of the meeting may be requested in writing from the Freedom of Information Office (HFI-35), Food and Drug Administration, rm. 12A-16, 5600 Fishers Lane, Rockville, MD 20857, approximately 15 working days after the meeting, at a cost of 10 cents per page. The transcript may be viewed at the Dockets Management Branch (HFA-305), Food and Drug Administration, rm. 1-23, 12420 Parklawn Dr., Rockville, MD 20857, approximately 15 working days after the meeting, between the hours of 9 a.m. and 4 p.m., Monday through Friday. Summary minutes of the open portion of the meeting may be requested in writing from the Freedom of Information Office (address above) beginning approximately 90 days after the meeting.

This notice is issued under section 10(a)(1) and (2) of the Federal Advisory Committee Act (5 U.S.C. app. 2), and FDA's regulations (21 CFR part 14) on advisory committees.

The Commissioner approves the scheduling of meetings at locations outside of the Washington, DC, area on the basis of the criteria of 21 CFR 14.22 of FDA's regulations relating to public advisory committees.

Dated: January 13, 1995.

Linda A. Suydam,

Interim Deputy Commissioner for Operations. [FR Doc. 95–1552 Filed 1–19–95; 8:45 am]
BILLING CODE 4160–01–F

Health Care Financing Administration

Privacy Act of 1974; System of Records

AGENCY: Department of Health and Human Services (HHS), Health Care Financing Administration (HCFA).

ACTION: Notice to propose a name change, purpose change, and the addition of new routine uses for an existing system of records.

SUMMARY: HCFA is proposing to amend the system notice for the "Supplemental Medical Insurance" (SMI) Accounting Collection and Enrollment System (SPACE)," System No. 09–70–0505, by revising the system name, revising the purpose, and by adding new routine uses. Also, sections of this notice have been updated to reference current

addresses and appropriate HCFA components.

HCFA is proposing to change the system name to better reflect the current function of the SPACE system, which now processes Medicare premium billing information for both Part B, SMI, and Part A, HI. The proposed new name is "Supplementary Medical Insurance (SMI) and Hospital Insurance (HI) Premium Accounting, Collection and Enrollment System (SPACE)." Despite the amendment to the system name, the acronym SPACE, which refers to this system, will not be changed.

The purpose of this system of records is being updated to include beneficiaries whose HI benefit premiums are paid by a State Medicaid agency, the U.S. Office of Personnel Management (OPM), or a formal third party group (the latter defined in 42 CFR section 408.80 through section 408.92). The purpose originally only references those beneficiaries whose SMI was paid by these named parties.

HCFA is also proposing to add routine uses, which permit the disclosure of data without the prior written consent of an individual, when the use of a record is for a purpose which is compatible with the purpose for which the record was collected. The proposed new routine uses would permit the disclosure of information to the following parties: OPM, formal third party groups, contractors in connection with the maintenance of automated data processing (ADP) software, and an individual or organization for research. (SEE SUPPLEMENTARY INFORMATION)

EFFECTIVE DATES: HCFA filed an altered system report with the Chair of the House Committee on Government Operations, the Chair of the Senate Committee on Governmental Affairs, and the Administrator, Office of Information and Regulatory Affairs, Office of Management and Budget (OMB) on January 13, 1995. To ensure that all parties have adequate time in which to comment, the revised system of records, including routine uses, will become effective 40 days from the publication of this notice or from the date it is submitted to OMB and the Congress, whichever is later, unless HCFA receives comments which require alterations to this notice.

ADDRESSES: Please address comments to Richard A. DeMeo, HCFA Privacy Act Officer, Office of Customer Relations and Communications, Office of Beneficiary Services, Health Care Financing Administration, Room 2–H–4 East High Rise Building, 6325 Security Boulevard, Baltimore, Maryland 21207–