

Secretary, Department of Health and Human Services, the Administrator, Alcohol, Drug Abuse, and Mental Health Administration, and the Director, National Institute on Drug Abuse, on the development of new initiatives and priorities and the efficient administration of drug abuse research, including prevention and treatment research, and research training. The Council also gives advice on policies and priorities for drug abuse grants and contracts, and reviews and makes final recommendations on grant applications.

Committee Name: National Advisory Mental Health Council

Date and Time: May 18-20: 9:00 a.m.

Place: May 18: National Institutes of Health, Building 31, Conference Room 10, 9000 Rockville Pike, Bethesda, MD 20892 May 19-20: Parklawn Building, Conference Rooms G and H, 5600 Fishers Lane, Rockville, MD 20857

Status of Meeting: OPEN—May 18: 9:00-5:00 p.m.; CLOSED—Otherwise

Contact: Rachel Driver, Room 9-105, Parklawn Building, 5600 Fishers Lane Rockville, MD 20857, (301) 443-3367.

Purpose: The National Advisory Mental Health Council advises the Secretary of Health and Human Services, the Administrator, Alcohol, Drug Abuse, and Mental Health Administration, and the Director, National Institute of Mental Health regarding policies and programs of the Department in the field of mental health. The Council reviews applications for grants-in-aid relating to research and training in the field of mental health and makes recommendations to the Secretary with respect to approval of applications for, and amount of, these grants.

Committee Name: Services Subcommittee of the Epidemiologic and Services Research Review Committee

Date and Time: May 27-29: 9:00 a.m.

Place: Sheraton Washington Hotel, 2660 Woodley Road, NW, Washington, DC 20008

Status of Meeting: OPEN—May 27: 9:00-10:00 a.m.; CLOSED—Otherwise

Contact: Gloria Yockelson, Room 9C-14, Parklawn Building 5600 Fishers Lane, Rockville, MD 20857, (301) 443-1367

Purpose: The Committee is charged with the initial review of applications for assistance from the National Institute of Mental Health for support of research and research training activities as they relate to mental health epidemiology, mental health services, with recommendations to the National Advisory Mental Health Council for final review.

Committee Name: Cognition, Emotion, and Personality Research Review Committee

Date and Time: May 28-30: 10:00 a.m.

Place: Dupont Plaza, 1500 New Hampshire Avenue, NW, Washington, DC 20036

Status of Meeting: OPEN—May 28: 9:00-10:00 a.m.; CLOSED—Otherwise

Contact: Shirley Maltz, Room 9C-26, Parklawn Building, 5600 Fishers Lane Rockville, MD 20857, (301) 443-3944

Purpose: The Committee is charged with the initial review of applications for assistance from the National Institute of Mental Health for support of research and research training activities relating to the fields of personality, cognition, emotion, and higher mental processes with recommendations to the National Advisory Mental Health Council for final review.

Substantive information may be obtained from the contact persons listed above. Summaries of the meetings and rosters of committee members may be obtained as follows: NIDA: Ms. Camilla Holland, Committee Management Officer, Room 16-22, Parklawn Building, 5600 Fishers Lane, Rockville, Maryland 20857, (301) 443-1622; NIMH: Ms. Joanna Kieffer, Committee Management Officer, Room 9-95, Parklawn Building, 5600 Fishers Lane, Rockville, MD 20857, (301) 443-4333.

Dated: April 9, 1987.

Estelle O. Brown,

Acting Committee Management Officer, Alcohol, Drug Abuse, and Mental Health Administration.

[FR Doc. 87-8290 Filed 4-13-87; 8:45 am]

BILLING CODE 4160-20-M

Food and Drug Administration

Advisory Committees; Meetings

AGENCY: Food and Drug Administration.

ACTION: Notice.

SUMMARY: This notice announces forthcoming meetings of public advisory committees of the Food and Drug Administration (FDA). This notice also summarizes the procedures for the meetings and methods by which interested persons may participate in open public hearings before FDA's advisory committees.

Meetings: The following advisory committee meeting are announced:

Orthopedic and Rehabilitation Devices Panel

Date, time, and place. May 7, 1:30 p.m. and May 8, 2 p.m., North Auditorium, Health and Human Services Bldg., 330

Independence Ave. SW. (enter at C St. entrance), Washington, DC.

Type of meeting and contact person. Open public hearing, May 7, 1:30 p.m. to 2 p.m.; open committee discussion, 2 p.m. to 5 p.m.; closed committee deliberations, 5 p.m. to 6 p.m.; open public hearing, May 8, 2 p.m. to 2:30 p.m.; open committee discussion, 2:30 p.m. to 4 p.m.; closed committee deliberations, 4 p.m. to 4:30 p.m.; Nirmal K. Mishra, Center for Devices and Radiological Health (HFZ-410), Food and Drug Administration, 8757 Georgia Ave., Silver Spring, MD 20910, 301-427-7156.

General function of the committee. The committee reviews and evaluates available data on the safety and effectiveness of devices and makes recommendations for their regulation.

Agenda—Open public hearing. Interested person may present data, information, or views, orally or in writing, on issues pending before the committee. Those desiring to make formal presentations should notify the contact person before April 30, and submit a brief statement of the general nature of the evidence or arguments they wish to present, the names and addresses of proposed participants, and an indication of the approximate time required to make their comments.

Open committee discussion. The committee will discuss a premarket approval application (PMA) for a prosthetic ligament device and the 4-year followup data for an approved electrical bone growth stimulator. The committee will also have a brief discussion regarding guidelines for evaluation of prosthetic ligament devices.

Closed committee deliberations. The committee may review and/or discuss trade secret and/or confidential commercial information relevant to PMA's. This portion of the meeting will be closed to permit discussion of this information (5 U.S.C. 552(b)(4)).

Blood Products Advisory Committee

Date, time, and place. May 7 and 8, 8:30 a.m., May 7, Lister Hill Auditorium, Bldg. 38A, National Library of Medicine, National Institutes of Health, 8600 Rockville Pike, May 8, Rm. 121, Bldg. 29, Office of Biologics Research and Review, 8800 Rockville Pike, Bethesda, MD.

Type of meeting and contact person. Open public hearing, May 7, 8:30 a.m. to 9:30 a.m., unless public participation does not last that long; open committee discussion, 9:30 a.m. to 3 p.m.; closed committee deliberations, 3:30 p.m. to 5:30 p.m., May 8, 8:30 a.m. to 1:30 p.m.;

Clay Sisk, Center for Drugs and Biologics (HFN-32), Food and Drug Administration, 5600 Fishers Lane, Rockville, MD 20857, 301-443-5455.

General function of the committee.

The committee reviews and evaluates available data on the safety, effectiveness, and appropriate use of blood products intended for use in the diagnosis, prevention, or treatment of human diseases. The committee also reviews and evaluates the quality and relevance of FDA's research program which provides scientific support for the regulation of these products.

Agenda—Open public hearing.

Interested persons requesting to present data, information, or views, orally or in writing, on issues pending before the committee should communicate with the committee contact person.

Open committee discussion. The committee will have additional discussions on (1) blood and blood components drawn for autologous use but for which conversion of one or more components to homologous use may be contemplated, (2) specifications for in vitro diagnostic tests for antibody to hepatitis B core antigen, and (3) review of the intramural research programs of the Laboratory of Cell Biology, Division of Blood and Blood Products, Office of Biologics Research and Review, Center for Drugs and Biologics, Food and Drug Administration. The work of the individual scientists in the Laboratory will be presented.

Closed committee discussion. The committee will review part of the intramural research program in the Office of Biologics Research and Review. This session of the meeting will be closed to prevent disclosure of personal information concerning individuals associated with this research program, disclosure of which would constitute a clearly unwarranted invasion of personal privacy (5 U.S.C. 552b(c)(6)). The committee may also review trade secret and/or confidential commercial information relevant to pending license applications and investigational new drugs (IND's). This portion of the meeting will be closed to permit discussion of this information (5 U.S.C. 552b(c)(4)).

Ear, Nose, and Throat Devices Panel

Date, time, and place. May 18, 8:30 a.m. and May 19, 8 a.m., Rm. 503-529A, Hubert H. Humphrey Bldg., 200 Independence Ave. SW., Washington, DC.

Type of meeting and contact person. Open public hearing, May 18, 8:30 a.m. to 9:30 a.m.; open committee discussion, 9:30 a.m. to 4:30 p.m.; closed committee deliberations, May 19, 8 a.m. to 9 a.m.;

open committee discussion, 9 a.m. to 4 p.m.; David A. Segerson, Center for Devices and Radiological Health (HFZ-470), Food and Drug Administration, 8757 Georgia Ave., Silver Spring, MD 20910, 301-427-8185.

General function of the committee.

The committee reviews and evaluates available data on the safety and effectiveness of devices and makes recommendations for their regulation.

Agenda—Open public hearing.

Interested persons may present data, information, or views, orally or in writing, on issues pending before the committee. Those desiring to make formal presentations should notify the contact person before May 4, and submit a brief statement of the general nature of the evidence or arguments they wish to present, the names and addresses of proposed participants, and an indication of the approximate time required to make their comments.

Open committee discussion. The committee will discuss a PMA for a cochlear implant for children.

Closed committee deliberations. The committee will discuss trade secret and/or confidential commercial information regarding the PMA. This portion of the meeting will be closed to permit discussion of this information (5 U.S.C. 552b(c)(4)).

Circulatory System Devices Panel

Date, time, and place. May 22, 8:30 a.m., Rm. 703-727A, Hubert H. Humphrey Bldg., 200 Independence Ave. SW., Washington, DC.

Type of meeting and contact person. Open public hearing, 8:30 a.m. to 9 a.m.; open committee discussion, 9 a.m. to 2 p.m.; closed committee deliberations, 2 p.m. to 4 p.m.; Keith Lusted, Center for Devices and Radiological Health (HFZ-450), Food and Drug Administration, 8757 Georgia Ave., Silver Spring, MD 20910, 301-427-7594.

General function of the committee. The committee reviews and evaluates available data on the safety and effectiveness of medical devices currently in use and makes recommendations for their regulation.

Agenda—Open public hearing.

Interested persons may present data, information, or views, orally or in writing, on issues pending before the committee. Those desiring to make formal presentations should notify the contact person before May 8, and submit a brief statement of the general nature of the evidence or arguments they wish to present, the names and addresses of proposed participants, and an indication of the approximate time required to make their comments.

Open committee discussion. The committee will discuss the labeling of pulse generator battery longevity and PMA's for prosthetic heart valves.

Closed committee deliberations. The committee may discuss trade secret and/or confidential commercial information regarding the heart valves listed above. This portion of the meeting will be closed to permit discussion of this information (5 U.S.C. 552b(c)(4)).

Ophthalmic Devices Panel

Date, time, and place. May 28 and 29, 9 a.m., Auditorium, Hubert H. Humphrey Bldg., 200 Independence Ave. SW., Washington, DC.

Type of meeting and contact person. Open public hearing, May 28, 9 a.m. to 10 a.m.; open committee discussion, 10 a.m. to 3 p.m.; closed committee deliberations, 3 p.m. to 4 p.m.; open public hearing, May 29, 9 a.m. to 10 a.m.; open committee discussion, 10 a.m. to 3 p.m.; closed committee deliberations, 3 p.m. to 4 p.m.; open committee discussion, 4 p.m. to 5 p.m.; Richard E. Lippman, Center for Devices and Radiological Health (HFZ-460), Food and Drug Administration, 8757 Georgia Ave., Silver Spring, MD 20910, 301-427-7320.

General function of the committee. The committee reviews and evaluates available data on the safety and effectiveness of devices currently in use and makes recommendations for their regulation. The committee also reviews data on new devices and makes recommendations regarding their safety and effectiveness and their suitability for marketing.

Agenda—Open public hearing. Interested persons may present data, information, or views, orally or in writing, on issues pending before the committee. Those desiring to make formal presentations should notify the contact person before May 11, and submit a brief statement of the general nature of the evidence or arguments they wish to present, the names and addresses of proposed participants, and an indication of the approximate time required to make their comments.

Open committee discussion. On May 28 the committee will discuss general issues relating to approvals of PMA's for Nd:YAG lasers and intraocular lenses (IOL's), and may discuss specific PMA's for these devices. If discussion of all pertinent Nd:YAG laser or IOL issues is not completed, discussion will be continued the following day. On May 29 the committee will discuss PMA's for contact lenses and other devices and requirements for PMA approval.

Closed committee deliberations. The committee may discuss trade secret and/or confidential commercial information relevant to PMA's for IOL's, Nd:YAG lasers, contact lenses, or other ophthalmic devices. These portions of the meeting will be closed to permit discussion of this information (5 U.S.C. 552b(c)(4)).

General and Plastic Surgery Devices Panel

Date, time, and place. May 29, 9 a.m., Rm. 703-727A, Hubert H. Humphrey Bldg., 200 Independence Ave. SW., Washington, DC.

Type of meeting and contact person. Open public hearing, 9 a.m. to 10 a.m.; open committee discussion, 10 a.m. to 4 p.m.; closed committee deliberations, 4 p.m. to 5 p.m.; Paul F. Tilton, Center for Devices and Radiological Health (HFZ-410), Food and Drug Administration, 8757 Georgia Ave., Silver Spring, MD 20910, 301-427-7156.

General function of the committee. The committee reviews and evaluates available data on the safety and effectiveness of devices and makes recommendations for their regulation.

Agenda—Open public hearing. Interested persons may present data, information, or views, orally or in writing, on issues pending before the committee. Those desiring to make formal presentations should notify the contact person before May 8, and submit a brief statement of the general nature of the evidence or arguments they wish to present, the names and addresses of proposed participants, and an indication of the approximate time required to make their comments.

Open committee discussion. The committee will discuss a PMA for a gelatin matrix implant, a PMA for a collagen-hydroxylapatite implant, a reclassification petition for gut sutures, and, if time allows, the proposed classification regulation of wound dressings.

Closed committee deliberations. The committee will discuss trade secret and/or confidential commercial information regarding a PMA. This portion of the meeting will be closed to permit discussion of this information (5 U.S.C. 552b(c)(4)).

Each public advisory committee meeting listed above 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. The dates and times reserved for the separate 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 it practical, in accordance with the agenda published in this **Federal Register** notice. Charges 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 oral presentation at the hearing's conclusion, if time permits, at the chairperson's discretion.

Persons interested in specific agenda items to be discussed in open session may ascertain from the contact person the approximate time of discussion.

Details on the agenda, questions to be addressed by the committee, and a current list of committee members are available from the contact person before and after the meeting. Transcripts of the open portion of the meeting will be available from the Freedom of Information Office (HFW-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. 4-62, 5600 Fishers Lane, 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 will be available from the Freedom of Information Office (address above) beginning approximately 90 days after the meeting.

The Commissioner, with the concurrence of the Chief Counsel, has determined for the reasons stated that those portions of the advisory committee meetings so designated in this notice shall be closed. The Federal Advisory Committee Act (FACA), as amended by the Government in the Sunshine Act (Pub. L. 94-409), permits such closed advisory committee meetings in certain circumstances. Those portions of a meeting designated as closed, however, shall be closed for the shortest possible time, consistent with the intent of the cited statutes.

The FACA, as amended, provides that a portion of a meeting may be closed where the matter for discussion involves a trade secret; commercial or financial information that is privileged or confidential; information of a personal nature, disclosure of which would be a clearly unwarranted invasion of personal privacy; investigatory files compiled for law enforcement purposes; information the premature disclosure of which would be likely to significantly frustrate implementation of a proposed agency action; and information in certain other instances not generally relevant to FDA matters.

Examples of portions of FDA advisory committee meetings that ordinarily may be closed, where necessary and in accordance with FACA criteria, include the review, discussion, and evaluation of drafts of regulations or guidelines or similar preexisting internal agency documents, but only if their premature disclosure is likely to significantly frustrate implementation of proposed agency action; review of trade secrets and confidential commercial or financial information submitted to the agency; consideration of matters involving investigatory files compiled for law enforcement purposes; and review of matters, such as personnel records or individual patient records, where disclosure would constitute a clearly unwarranted invasion of personal privacy.

Examples of portions of FDA advisory committee meetings that ordinarily shall not be closed include the review, discussion, and evaluation of general preclinical and clinical test protocols and procedures for a class of drugs or devices; consideration of labeling requirements for a class of marketed drugs or devices; review of data and

information on specific investigational or marketed drugs and devices that have previously been made public; presentation of any other data or information that is not exempt from public disclosure pursuant to the FACA, as amended; and, notably deliberative sessions to formulate advice and recommendations to the agency on matters that do not independently justify closing.

This notice is issued under section 10(a)(1) and (2) of the Federal Advisory Committee Act (Pub. L. 92-463, 86 Stat. 770-776 (5 U.S.C. App. I)), and FDA's regulations (21 CFR Part 14) on advisory committees.

Dated: April 8, 1987.

Frank E. Young,

Commissioner of Food and Drugs.

[FR Doc. 87-8234 Filed 4-13-87; 8:45 am]

BILLING CODE 4160-01-M

Farmer's Union Grain Terminal Association; Withdrawal of Approval of NADA

AGENCY: Food and Drug Administration.
ACTION: Notice.

SUMMARY: The Food and Drug Administration (FDA) is withdrawing approval of a new animal drug application (NADA) held by the Farmer's Union Grain Terminal Association. The NADA provides for use of a Type A article containing 0.4- or 10-gram-per-pound tylosin for making Type C swine feeds. The firm requested the withdrawal of approval.

EFFECTIVE DATE: April 24, 1987.

FOR FURTHER INFORMATION CONTACT:

Mohammad I. Sharar, Center for Veterinary Medicine (HFV-214), Food and Drug Administration, 5600 Fishers Lane, Rockville, MD 20857, 301-443-3184.

SUPPLEMENTARY INFORMATION: Farmer's Union Grain Terminal Association, Feed Division, P.O. Box 1447, Sioux Falls, SD 57117, is the sponsor of NADA 96-160 originally approved May 2, 1974 (39 FR 15270). The NADA provides for use of a 0.4-gram-per-pound tylosin Type A article to make a Type C swine feed for use as in 21 CFR 558.625(f)(1)(vi)(a). Subsequently, the NADA was approved for use of a 10-gram-per-pound tylosin Type A article for the same purposes (45 FR 23686, April 8, 1980).

By letter dated January 20, 1987, the sponsor requested the withdrawal of approval because the product was not being manufactured or marketed.

Therefore, under the Federal Food, Drug, and Cosmetic Act (sec. 512(e), 82 Stat. 345-347 (21 U.S.C. 360(e))) and

under authority delegated to the Commissioner of Food and Drugs (21 CFR 5.10) and redelegated to the Center for Veterinary Medicine (21 CFR 5.84), and in accordance with § 514.115

Withdrawal of approval of application (21 CFR 514.115), notice is given that approval of NADA 96-160 and all supplements thereto is hereby withdrawn, effective April 24, 1987.

In a final rule published elsewhere in this issue of the *Federal Register*, FDA is removing those portions of the regulations that reflect this approval and is removing the firm from the list of sponsors of approved NADA's.

Dated: April 8, 1987.

Gerald B. Guest,

Director, Center for Veterinary Medicine.

[FR Doc. 87-8232 Filed 4-13-87; 8:45 am]

BILLING CODE 4160-01-M

Health Resources and Services Administration

Application Announcement for Grants for Two-Year Programs of Schools of Medicine or Osteopathy

The Bureau of Health Professions, Health Resources and Services Administration, announces that applications for Fiscal Year 1987 Grants for Two-Year Programs of Schools of Medicine or Osteopathy are now being accepted under the authority of section 788(a), of the Public Health Service Act, as amended by Pub. L. 99-129.

Section 788(a) authorizes the award of grants to maintain and improve schools which provide the first or last two years of education leading to the degree of doctor of medicine or osteopathy. Grants provided under this authority to schools that were in existence on September 30, 1985, may be used for construction and purchase of equipment.

To be eligible for a grant under this authority, the applicant must be a public or nonprofit, private school providing the first or last two years of education leading to the degree of doctor of medicine or osteopathy and be accredited or be operated jointly with a school that is accredited by a recognized body or bodies approved for such purpose by the Secretary of Education.

Application materials will be sent only upon request. Requests for application materials and questions regarding grants policy should be directed to: Grants Management Officer (D31), Bureau of Health Professions, Health Resources and Services Administration, Parklawn Building, Room 8C-22, 5600 Fishers Lane, Rockville, Maryland 20857, Telephone (301) 443-6880.

The deadline for receipt of applications is May 29, 1987.

Applications shall be considered as meeting the deadline if they are either:

(1) Received on or before the deadline date, or

(2) Postmarked on or before the deadline date, and received in time for submission to the independent review group.

A legibly dated receipt from a commercial carrier or U.S. Postal Service will be accepted in lieu of a postmark. Private metered postmarks shall not be acceptable as proof of timely mailing.

Approximately \$500,000 is available in Fiscal Year 1987 for competing awards. Authorization for the current fiscal year is provided under a Continuing Resolution (Public Laws 99-500 and 591).

Should additional programmatic information be requested, please contact: Multidisciplinary Resources Development Branch, Division of Medicine, Bureau of Health Professions, Health Resources and Services Administration, Parklawn Building, Room 4C-16, 5600 Fishers Lane, Rockville, Maryland 20857, Telephone (301) 443-6817.

The standard application form and specific instructions for this program have been approved by the Office of Management and Budget under the Paperwork Reduction Act. The OMB clearance number is 0915-0060.

This program is not yet listed in the Catalog of Federal Domestic Assistance, therefore, no catalog number has been assigned. Applications submitted in response to this announcement that request construction assistance are subject to the intergovernmental review under provisions of Executive Order 12372, as supplemented by 42 CFR Part 100, Intergovernmental Review of Federal Programs. Applications submitted for program support only are not subject to intergovernmental review under these provisions.

Dated: April 8, 1987.

David N. Sundwall,

Assistant Surgeon General.

[FR Doc. 87-8291 Filed 4-13-87; 8:45 am]

BILLING CODE 4160-15-M

Social Security Administration

Statement of Organization, Functions and Delegations of Authority

Part S of the Statement of Organization, Functions and Delegations of Authority for the Department of Health and Human Service (DHHS) covers the Social Security

Administration (SSA). Notice is given that Chapter SB as published in the *Federal Register* on January 4, 1983 and amended on August 15, 1983 is amended to establish the Division of Integration and Environmental Testing (DIET) and to realign other division-level functional responsibilities in the Office of System Operations (OSO). Chapter SU as published in the *Federal Register* on January 4, 1983 and amended on August 15, 1983; May 18, 1984; October 17, 1984; September 3, 1985 and November 8, 1985 is amended to reflect the abolishment of the Office of Systems Engineering in the Office of System Integration (OSI). These changes result from a major consolidation and realignment of systems functional responsibilities which is intended to improve organizational effectiveness.

The new material and changes are as follows:

Section SB.10 The Office of System Operations—(Organization):

D. The Office of Computer Processing Operations (SBP).

Add:

5. Division of Integration and Environmental Testing (SBP6).

Section SB.20 The Office of System Operations—(Functions):

D. The Office of Computer Processing Operations (SBP).

Add:

In Line 14, after "... information data.", add "The Office of Computer Processing Operations (OCPO) plans, directs and controls the integration, testing and, in conjunction with OSI, OSR and OIS, the production release of new or revised hardware and systems software. It develops policy and standards for integration testing, validation and acceptance of Automated Data Processing/Telecommunications systems software."

Delete:

1. The Division of Production Systems Operations (SBP1) in its entirety.

2. The Division of Computer Operations Production Control (SBP2) in its entirety.

3. The Division of Computer Operations Systems Software (SBP3) in its entirety.

4. The Division of Telecommunications Systems Operations (SBP5) in its entirety.

Add:

1. The Division of Production Systems Operations (SBP1):

a. Operates the centralized OSO computer facility, which includes computer systems hardware and associated peripheral equipment.

b. Directs the continuous operations of SSA's host telecommunications

computers in support of SSA-designed networks.

c. Schedules day-to-day workflow for the ADP facility within plans and priorities established by OCPO's Division of Computer Operations Production Control.

d. Controls the flow of materials into ADP production jobs. Reviews production results for accuracy and completeness.

e. Analyzes equipment problems, isolates malfunctions and oversees correction actions by SSA or vendor personnel.

f. Schedules and assures preventive maintenance of all equipment under the operational control of OCPO.

g. Develops and maintains a centralized integrated control center for use in monitoring the operating systems utilization, network control facilities and environmental status.

2. The Division of Computer Operations Production Control (SBP2):

a. Manages the production workload of OSO and administers effective resource utilization.

b. Designs, develops, implements and operates production control ADP systems which supervise library controls, automates the scheduling and allocates the production workload.

c. Manages and directs automated magnetic media processes and directs the activity of the magnetic tape library function.

d. Participates in the design reviews of proposed application systems to assure operational support and control aspects are being considered. Analyzes applications systems to assure compliance with systems standards. Approves applications systems for production status and incorporates them into the production library.

e. Assembles input material for ADP production jobs and delivers them to the Division of Production Systems Operations.

f. Expedites processing of critical jobs, operations and corrections.

g. Provides liaison with the users on status of production jobs and/or associated problems as required.

h. Maintains the integrity, manages and performs required recovery of all operational data, data media, tape and direct access for systems.

i. Maintains and enhances a transaction system for the control of a high-volume tape library.

j. Analyzes performance of the ADP production processes and recommends and implements improvements. Destroys sensitive material in compliance with provisions of the Privacy Act and SSA procedures.

3. The Division of Computer Operations Systems Software (SBP3):

a. Directs the analysis, design, development, implementation and maintenance of computer operating systems and utility software in support of programmatic and management information workloads for SSA's central data processing center and field components.

b. Directs the design, development, testing and continuing support of specialized data communications control software used to support SSA's data communications systems.

c. Directs the design, development, implementation and maintenance of information systems software in support of the central data processing center's problem, change and configuration management systems.

d. Supports the utility software and user activities in the areas of computer graphics, small computers and nonimpact printers.

e. Supports the user liaison and systems development activities of other OSO components in the resolution of technical and operational problems.

4. The Division of Telecommunications System Operations (SBP5):

a. Directs the operations of SSA's telecommunications network facilities for the transmission of program and management data over SSA-established networks.

b. Manages traffic flow between the telecommunications complex and other SSA computers. Monitors telecommunications operations, analyzes equipment problems and effects proper maintenance and repair.

c. Directs the implementation of new or revised operating policies and procedures. Recommends new procedures and appraises telecommunications operating instructions, centrally and in the field.

d. Establishes and enforces standards for controlling workflow and for assuring the integrity of data processed through the various data communications operations.

e. Acts as liaison with common carriers and network equipment vendors to maintain operational effectiveness of equipment.

f. Directs the operational performance evaluation of SSA's data communications systems.

g. Provides technical expertise and assistance on data communications procurements and other SSA systems modernization projects.

h. Directs the design, development and implementation of software to gather and report statistical information

on the functioning of telecommunications networks.

Distributes the information to other SSA components to report on network performance and equipment utilization.

i. Designs and implements security software for SSA's telecommunications network and ensures that related procedures are followed by technical personnel.

j. Manages the installation, removal and relocation of local and remote telecommunications facilities, assuring compliance with governing Federal regulations.

k. Maintains an integrated control center, centrally and at the remote network nodes, to provide a point of contact of field offices reporting equipment or operational problems.

l. Conducts ongoing analyses of network configurations and workloads and initiates changes to the network topology to optimize cost/performance.

m. Develops standards and procedures for applications developers in interfacing to SSA's data communications network. Evaluates requested or proposed applications for impact on network resources.

n. Maintains and controls an inventory of all remote data communications equipment which assesses SSA's telecommunications networks, and the history and status of equipment outages for all SSA owned or leased data communications equipment.

o. Assigns, maintains and provides to data communications systems users, telecommunications site routing codes and terminal identifiers consistent with Governmentwide network addressing conventions.

p. Conducts investigations and analysis of system problems affecting local and remote users of the data communications networks, and provides liaison with regional staffs in identifying and correcting chronic problems and trends.

Add:

5. The Division of Integration and Environmental Testing (SBP6):

a. Directs and controls all activities with the release of new or enhanced versions of host programmatic and telecommunications-related software. Enforces software acceptance and certifications standards. Directs the initial staging of program modules to be tested, including generation of executable code.

b. Develops and maintains extensive test data bases for use in the acceptance, integration and environmental testing processes. Develops and incorporates the use of software simulators and emulators in software acceptance testing.

c. Directs the integration testing of new or enhanced communications host software, remote network/terminal and microprocessor software and network communications software. Participates in the movement and/or migration of software systems and associated data files between complexes and processing components.

d. Directs environmental testing to ensure that new or enhanced software is compatible with changing hardware configurations. Directs the integration of new or enhanced SSA programmatic software. Administers the generation of finalized testing results for evaluation. Directs software performance evaluations, parallel testing, timing studies, inter/intrasystem relationship and testing trend analysis.

e. Responsible for administering ADP hardware integration and acceptance testing.

f. Provides the checks and balances on SSA's ADP systems and equipment procurements for complying with contractual performance requirements throughout the life cycle of the procurement.

E. The Office of System Support and Planning (SBL):

Add:

In Line 7, after "... OSO.", add "The Office develops an ADP Capacity Plan to perform capacity allocation, monitor its usage and determine future requirements, based upon future workload information."

In Line 11, after "... resources.", add "It participates in the development of policies and procedures for the acquisition of ADP Computer and Telecommunications equipment, software and services, in compliance with SSA, HHS, the Office of Management and Budget and the General Services Administration policies and regulations."

Delete:

1. The Division of Operational Capacity Performance Management (SBL) in its entirety.

Add:

1. The Division of Operational Capacity Performance Management (SBL):

a. Evaluates computer performance and monitors resource utilization to ensure that the ADP and telecommunications systems are utilized effectively and efficiently. Analyses systems capacity as it relates to utilization and service objectives and prepares recommendations for upper management. As directed, performs similar functions for other SSA components including the program service centers.

b. Ensures that sufficient ADP capacity is available to process present and future workloads, coordinating decisions on target systems for new/modified workloads and system configuration changes.

c. Monitors the OSO service delivery to ensure that systems performance objectives as defined in the User Service Agreements are being met. Provides recommendations to enhance delivered service as necessary, and ensure that data bases are efficiently implemented.

d. Provides advice and services to other OSO components in the use of computer performance evaluation tools and the interpretation of reports/data resulting from evaluation and utilization studies.

e. Uses operations research tools (e.g., simulation and mathematical models and statistical analyses) to investigate operational efficiency problems and develop relationships between transaction volumes, resource utilization and resulting service delivery.

f. Schedules, arranges, conducts and reports on structured systems' effectiveness reviews to compare OSO service commitments with delivered levels of performance and contributes towards planning and enhancement of existing systems.

g. Coordinates, assists in the development/maintenance, and monitors all Systems service level agreements. Represents OSO in the User Service Agreement negotiations.

h. Performs a wide range of user coordination and problem resolution functions. Gathers and disseminates timely information, related to operational problems, errors and changes that affect the users.

2. The Division of Standards and Control (SBL2):

Delete:

g. Performs ongoing analysis of equipment configurations and projects requirements based on technological enhancements, workload changes and growth.

h. Prepares comprehensive reports and technical justifications for management's use.

Add:

g. Formulates an OSO-wide Systems Plan and assigns responsibility among major OSO components for various parts of the Plan. Work with OSO components to evaluate their proposed systems objectives in terms of technical feasibility, availability of OSO resources and systems costs. Identifies the major OSO activities and resources needed to support these objectives.

h. Directs and coordinates the OSO activities associated with operational planning and ADP Systems Planning.

i. Coordinates and directs the development of the total OSO technical manpower, equipment and other special costs for the SSA budget process and justifies these on the basis of the Operational Systems Plan. Allocates resources and monitors projects for all OSO activities, directs the preparation of detailed plans on the project or operational activity level and authorizes the use of resources by OSO components in support of these plans.

j. Monitors progress and use of manpower and equipment resources by OSO components against their approved plans.

k. Assists OSO components in the use of standard methods for project management.

Delete:

3. The Division of Operational Resource Management (SBL3) in its entirety.

Add:

3. The Division of Operational Resource Management (SBL3):

a. Directs OSO's participation in the Information Technology Systems (ITS) procurement process.

b. Performs technical and cost reviews of all OSO/ITS procurements.

c. Provides support for ITS Technical Evaluation Committees.

d. Support contract administration for all OSO ITS contracts.

e. Provides technical support to Project Officers in the development, modification and administration of ITS contracts.

f. Directs the renewal process for existing lease and maintenance contracts for ITS and telecommunications equipment and services.

g. Manages the fiscal administration of all implemented ITS contracts, collecting, analyzing and reporting performance data to support required fiscal and other contractual proceedings.

h. Manages a centralized inventory of all SSA ITS and telecommunications equipment and manages the ITS excess equipment process.

i. Provides for the centralized certification and authorization for the lease and maintenance of SSA's ITS and telecommunications equipment.

j. Provides necessary staff support to the users within OSO for the development of procurement documents and documentation.

k. Develops and maintains the OSO macroprocurement plan which relates to planned major acquisitions of ITS equipment, software, system design and system support services. Coordinates

the procurement plans and related budget items with appropriate Systems management and staffs.

l. Provides technical support to OSO and other SSA components during major procurement activities.

Section SU.10 *The Office of System Integration*—(Organization):

G. The Office of Programmatic Systems (SUF).

Delete:

7. The Division of SSI Systems (SUF7).

Add:

4. The Division of SSI Systems (SUF7).

Delete:

H. The Office of System Engineering (SUH) in its entirety.

Section SU.20 *The Office of System Integration*—(Functions):

G. The Office of Programmatic Systems (SUF).

Delete:

7. The Division of SSI Systems (SUF7) in its entirety.

Add:

4. The Division of SSI Systems (SUF7).
a. Provides the systems analysis, design, programming and testing necessary to develop and maintain current, new and redesigned systems in response to approved user system requirements for SSI claims and postentitlement transaction processing as well as a variety of supporting applications. These systems: edit incoming new records and transactions; maintain and revise the SSI master file to reflect changes; compute both Federal SSI benefit and State supplementary payments and produce payment information for the Treasury Department; account for disbursement of Federal and State funds; prepare recipient notices of claims decisions and changes in status and payment; identify and control overpayment activity; select and control cases requiring redetermination; exchange data with Government record systems to verify recipient income; generate data for State use in determining supplementation amounts and Medicaid eligibility; provide record query and response capability; control folder location and movement; produce statistical, management and actuarial data as needed and control exception processing and diary control mechanisms.

b. Translates user requirements, as approved by the Office of System Requirements (OSR), into detailed design, development and testing activities and activities and system documentation for current, new or redesigned systems.

c. Conducts liaison with other SSA components and Federal and State agencies to determine the feasibility,

and to plan the development, of SSI claims, transaction and support systems.

d. Provides OSR, the Associate Commissioner for System Integration and other SSA offices, as appropriate, with a technical assessment of the effect of legislative, administrative and systems modernization proposals on existing SSI claims, transaction and support system applications.

Delete:

H. The Office of Systems Engineering (SUH) in its entirety.

Dated: April 2, 1987.

Jaime L. Manzano,

Deputy Commissioner for Systems.

[FR Doc. 87-8256 Filed 4-13-87; 8:45 am]

BILLING CODE 4190-11-M

Privacy Act of 1974; Notification of New Routine Uses and Minor Revision

AGENCY: Social Security Administration (SSA), Department of Health and Human Services (HHA).

ACTION: New Routine Uses and Minor Revision.

SUMMARY: In accordance with the Privacy Act (5 U.S.C. 552a(e)(11)), we are issuing public notice of our intent to:

(1) Establish a new routine use of information maintained in the system of records entitled "Kentucky Birth Records System, HHS/SSA/DO(KY), 09-60-0220." This proposed routine use would permit SSA to disclose information to the General Services Administration (GSA) and the National Archives and Records Administration (NARA) for the purposes of conducting record management studies.

(2) Revise a routine use already applicable to the majority of SSA's systems of records which currently provides for disclosure to the Department of Justice (DOJ) to defend SSA components and employees in litigation matters affecting the systems to include additional disclosures involving litigation.

We are proposing to make a minor technical revision to the Federal Register applicable to the Kentucky Birth Records System to make it accurate and complete. We invite public comments on these proposals.

DATE: These proposals will become effective as proposed without further notice on May 14, 1987, unless we receive comments on or before that date which would result in a contrary determination.

ADDRESS: Interested individuals may comment on this proposal by writing to Mr. Bernard A. Oehlers, Acting SSA

Privacy Officer, Social Security Administration, Room L1140 West Low Rise Building, 6401 Security Boulevard, Baltimore, Maryland 21235. All comments received will be available for public inspection at that address.

FOR FURTHER INFORMATION CONTACT:

Mr. Willie J. Polk, Social Insurance Specialist, Social Security Administration, Office of Regulations, Privacy Branch, Room L1140 West Low Rise Building, 6401 Security Boulevard, Baltimore Maryland 21235, telephone (Area Code 301) 594-5599.

SUPPLEMENTARY INFORMATION:

I. Discussion of the Proposed Routine Uses

A. Disclosure to GSA and NARA for the Purpose of Conducting Records Management Studies

We are proposing to revise the routine use section of the Kentucky Birth Records System by adding a new routine use which permits disclosure to GSA and NARA for the purpose of conducting records management studies. The Administrator, GSA and the Archivist, NARA, are charged by 44 U.S.C. 2904, as amended by the National Archives and Records Administration Act of 1984, with promulgating standards, procedures and guidelines with respect to records management and the conduct of records management studies.

Section 2906 of this law, also amended by the NARA Act of 1984, provides that GSA and NARA shall have access to Federal agencies' records and that agencies shall cooperate with GSA and NARA. To facilitate GSA and NARA records management studies which may involve the Kentucky Birth Records System, we are adding the routine use to comply with the Privacy Act. The routine use is as follows:

Nontax return information which is not restricted from disclosure by Federal law may be disclosed to the General Services Administration and the National Archives and Records Administration for the purpose of conducting records management studies with respect to their duties and responsibilities under 44 U.S.C. 2904 and 2906, as amended by the National Archives and Records Administration Act of 1984.

We publish this routine use in the Federal Register for other SSA systems of records on March 10, 1986 (see 51 FR 8243, March 10, 1986). The notice of the Kentucky Birth Records System below contains the above new routine use.

B. Disclosure to DOJ, a Court or Other Tribunal, or Another Party Before Such Tribunal.

The majority of SSA's notices of systems of records currently contain a routine use which permits disclosure of DOJ for the purpose of defending SSA components and employees (or other Federal agencies where litigation is likely to affect SSA components) in litigation matters involving the systems of records. We are expanding this routine use to permit disclosure, in addition to DOJ, to a court or other tribunal or another party before such tribunal and to permit disclosure to these parties when SSA or another party brings suit or in cases in which SSA has an interest in the litigation. The revised routine use is as follows:

Disclosure may be to DOJ, to a court or other tribunal, or another party before such tribunal, when:

(a) SSA, or any component thereof; or
(b) Any SSA employee in his/her official capacity; or

(c) Any SSA employee in his/her individual capacity where DOJ (or SSA where it is authorized to do so) has agreed to represent the employee; or
(d) The United States or any agency thereof where SSA determines that the

litigation is likely to affect the operations of SSA or any of its components,

is a party to litigation or has an interest in such litigation, and SSA determines that the use of such records by DOJ, the tribunal, or the other party before such tribunal is relevant and necessary to the litigation, provided, however, that in each case, SSA determines that such disclosure is compatible with the purpose for which the records were disclosed.

Wage and other information which are subject to the disclosure provisions of the Internal Revenue Code (IRC) (26 U.S.C. 6103) will not be disclosed under this routine use unless disclosure is expressly permitted by the IRC.

We will disclose information under this routine use only as necessary to defend SSA components or employees in litigation matters involving the systems of records which will contain the routine use or when SSA has an interest in litigation which might affect SSA operations.

Because of the number of systems of records to which the routine use will apply and the costs of republishing the individual notices of each one, we are republishing only the identification number and the name of each system and the volume, page number and date of the Federal Register issue in which a notice of the systems last was published

in the Federal Register. The systems of records are as follows:

- 09-60-0001—Commissioner's Correspondence File, HHS/SSA/OC (51 FR 8243, March 10, 1986);
- 09-60-0002—Automated Document Control and Retrieval System, HHS/SSA/OGA (51 FR 8243, March 10, 1986);
- 09-60-0003—Attorney Fee File, HHS/SSA/OHA (51 FR 8243, March 10, 1986);
- 09-60-0004—Working File of the Appeals Council, HHS/SSA/OHA (51 FR 8243, March 10, 1986);
- 09-60-0005—Hearing Office File, HHS/SSA/OHA (51 FR 8243, March 10, 1986);
- 09-60-0006—Storage of Hearings Records: Tape Cassettes and Audiograph Discs, HHS/SSA/OHA (51 FR 8243, March 10, 1986);
- 09-60-0008—Administrative Law Judge's File, HHS/SSA/OHA (51 FR 8243, March 10, 1986);
- 09-60-0009—Hearings and Appeals Case Control System, HHS/SSA/OHA (51 FR 8243, March 10, 1986);
- 09-60-0012—Listing and Alphabetical Name File (Folder) of Vocational Experts, Medical Advisors and Medical Consultants, HHS/SSA/OHA (51 FR 8243, March 10, 1986);
- 09-60-0013—Records of Usage of Medical Advisors, Medical Consultants and Vocational Experts, HHS/SSA/OHA (51 FR 8243, March 10, 1986);
- 09-60-0014—Curriculum Vitae and Professional Qualifications of Medical Advisors, Medical Consultants and Resume of Vocational Experts, HHS/SSA/OHA (51 FR 8243, March 10, 1986);
- 09-60-0017—Personnel Research and Merit Promotion Test Records, HHS/SSA/OHA (51 FR 8243, March 10, 1986);
- 09-60-0031—Employee Production and Accuracy Records, HHS/SSA/OMB (51 FR 8243, March 10, 1986);
- 09-60-0032—Employee Indebtedness System, HHS/SSA/OMB (51 FR 8243, March 10, 1986);
- 09-60-0033—Report on Paid Consultants, HHS/SSA/OMB (51 FR 8243, March 10, 1986);
- 09-60-0037—General Criminal Investigations Files, HHS/SSA/OMB (51 FR 8243, March 10, 1986);
- 09-60-0038—Employee Building Pass Files, HHS/SSA/OMB (51 FR 8243, March 10, 1986);
- 09-60-0040—Quality Review System, HHS/SSA/OA (51 FR 8243, March 10, 1986);
- 09-60-0042—Quality Review Case Files, HHS/SSA/OA (51 FR 8243, March 10, 1986);
- 09-60-0044—Disability Determination Service Processing File, HHS/SSA/OD (51 FR 8243, March 10, 1986);
- 09-60-0045—Black Lung Payment System, HHS/SSA/OSR (51 FR 8243, March 10, 1986);
- 09-60-0046—Consultative Physicians File, HHS/SSA/OD (51 FR 8243, March 10, 1986);
- 09-60-0050—Completed Determination Record—Continuing Disability Determinations, HHS/SSA/OP (51 FR 8243, March 10, 1986);

- 09-60-0057—Quality Evaluation Data Records, HHS/SSA/OA (51 FR 8243, March 10, 1986);
- 09-60-0058—Master Files of Social Security Number Holders, HHS/SSA/OSR (51 FR 8243, March 10, 1986);
- 09-60-0059—Earnings Recording and Self-Employment Income System, HHS/SSA/OSR (51 FR 8243, March 10, 1986);
- 09-60-0063—Resource Accounting and Project Management System, HHS/SSA/OSPI (51 FR 8243, March 10, 1986);
- 09-60-0066—Claims Development Record, HHS/SSA/RO (51 FR 8243, March 10, 1986);
- 09-60-0075—Congressional Bills Tracking System, HHS/SSA/OLRP (51 FR 8243, March 10, 1986);
- 09-60-0077—Congressional Inquiry File, HHS/SSA/RO (51 FR 8243, March 10, 1986);
- 09-60-0078—Public Inquiry Correspondence File, HHS/SSA/RO (51 FR 8243, March 10, 1986);
- 09-60-0089—Claims Folders System, HHS/SSA/OP (51 FR 8243, March 10, 1986);
- 09-60-0090—Master Beneficiary Record, HHS/SSA/OSR (51 FR 8243, March 10, 1986);
- 09-60-0094—Recovery Accounting for Overpayments, HHS/SSA/OSR (51 FR 8243, March 10, 1986);
- 09-60-0095—Health Insurance Overpayment Ledger Cards, HHS/SSA/OSR (51 FR 8243, March 10, 1986);
- 09-60-0103—Supplemental Security Income Record, HHS/SSA/OSR (51 FR 8243, March 10, 1986);
- 09-60-0110—Supplemental Security Income File of Refunds, HHS/SSA/OSR (51 FR 8243, March 10, 1986);
- 09-60-0111—Debit Voucher File (Supplemental Security Income), HHS/SSA/OSR (51 FR 8243, March 10, 1986);
- 09-60-0118—Non-Contributory Military Service Reimbursement System, HHS/SSA/OACT (51 FR 8243, March 10, 1986);
- 09-60-0119—Special Age 72 Benefit Trust Fund Transfer Project, HHS/SSA/OACT (51 FR 8243, March 10, 1986);
- 09-60-0184—Hearing Office Master Calendar, HHS/SSA/OHA (51 FR 8243, March 10, 1986);
- 09-60-0186—SSA Litigation Tracking System, HHS/SSA/LS (51 FR 8243, March 10, 1986);
- 09-60-0210—Record of Individuals Authorized Entry to Secured ADP Areas, HHS/SSA/OS (51 FR 8243, March 10, 1986);
- 09-60-0212—Supplemental Security Income Quality Initial Claims Review Process System, HHS/SSA/OA (51 FR 8243, March 10, 1986);
- 09-60-0213—Quality Review of Hearing/Appellate Process, HHS/SSA/OHA (51 FR 8243, March 10, 1986);
- 09-60-0214—Personal Identification Number File, HHS/SSA/OA (48 FR 37526, August 18, 1983);
- 09-60-0219—Representative Disqualification/Suspension Information System, HHS/SSA/ORSI (51 FR 8243, March 10, 1986).

II. Compatibility of the Proposed Routine Use

The Privacy Act (5 U.S.C. 552a(a)(7) and 552a(b)(3)) and our disclosure

regulation (20 CFR Part 401) both permit us to disclose information under a routine use if the information will be used for a purpose which is compatible with the purpose for which we collected the information. Section 401.310 of the regulation permits us to disclose information under a routine use for administering our programs or for use in similar income-maintenance or health-maintenance programs of other agencies. Section 401.205 of the regulation requires us to disclose information when a law specifically requires it.

With regard to GSA and NARA, disclosure would assist SSA in its records management. Further 44 U.S.C. 2906, as amended by the NARA Act of 1984, requires us to disclose information to GSA and NARA for records management purposes.

Disclosure would be made to DOJ, a court or other tribunal, or another party before such tribunal, as necessary, in litigation matters involving SSA operations. We consider disclosure in this instance as an extension of program administration. Thus, both routine uses are appropriate and meet the criteria in the Privacy Act and the regulation.

III. Effect of the Proposed Routine Uses on Individual Rights

Information will be disclosed to GSA and NARA for the purpose of their reviewing SSA's records management practices as mandated by Federal law and the only for this purpose. Information will be disclosed to DOJ, a court or other tribunal, or another party before such tribunal in litigation, as necessary, to defend SSA components or employees or if the litigation would affect SSA operations. The disclosures will be made in complete compliance with the Privacy Act. Thus, we do not anticipate that they would have an unwarranted affect on the rights of individuals.

IV. Minor Revision

We initially published a notice of the Kentucky Birth Records System in the *Federal Register* on June 3, 1986 (51 FR 19899). In that publication, we inadvertently did not publish the "Authority for Maintenance of the System" section of the notice. That section of the notice should have identified section 205 of the Social Security Act as the authority for the system. We have revised the notice to reflect this information.

Dated: April 6, 1987.

Dorcas R. Hardy,
Commissioner of Social Security.

09-60-0220

SYSTEM NAME:

Kentucky Birth Records System, HHS/SSA/DO(Ky).

SECURITY CLASSIFICATION:

None.

SYSTEM LOCATION:

Social Security district and branch offices located in the State of Kentucky. Individuals should consult Kentucky telephone directories for address and telephone information.

CATEGORIES OF INDIVIDUALS COVERED BY THE SYSTEM:

Members of the general public whose birth records have been registered in the State of Kentucky.

CATEGORIES OF RECORDS IN THE SYSTEM:

The system consists of an index of Kentucky birth records. Included on the index are the individual's name, mother's maiden name, date and place of birth, certificate number and volume number of the index.

AUTHORITY FOR MAINTENANCE OF THE SYSTEM:

Section 205 of the Social Security Act.

PURPOSE:

Information in the system will be used by Social Security Administration (SSA) offices in the State of Kentucky to provide evidentiary proof of age and other facts about individuals applying for various Social Security benefits, Supplemental Security Income payments and Social Security numbers.

ROUTINE USES OF RECORDS MAINTAINED IN THE SYSTEM, INCLUDING CATEGORIES OF USERS AND THE PURPOSES OF SUCH USES:

Disclosure may be made as routine uses as indicated below:

1. To a Congressional office in response to an inquiry from that office made at the request of the subject of a record.

2. To the Department of Justice (DOJ), to a court or other tribunal, or to another party before such tribunal, when:

(a) The Department of Health and Human Services (HHS)/SSA, or any component thereof; or

(b) Any HHS/SSA employee in his/her official capacity; or

(c) Any HHS/SSA employee in his/her individual capacity where DOJ (or HHS/SSA where it is authorized to do so) has agreed to represent the employee; or

(d) The United States or any agency thereof where HHS/SSA determines that the litigation is likely to affect the operations of HHS/SSA or any of its components.

Is a party to litigation or has an interest in such litigation, and HHS/SSA determines that the use of such records by DOJ, the tribunal, or the other party before such tribunal is relevant and necessary to the litigation, provided, however, that in each case, HHS/SSA determines that such disclosure is compatible with the purpose for which the records were collected.

3. *Nontax return information which is not restricted from disclosure by Federal law may be disclosed to the General Services Administration and the National Archives and Records Administration for the purpose of conducting records management studies with respect to their duties and responsibilities under 44 U.S.C. 2904 and 2906, as amended by the National Archives and Records Administration Act of 1984.*

POLICIES AND PRACTICES FOR STORING, RETRIEVING, ACCESSING, RETAINING, AND DISPOSING OF RECORDS IN THE SYSTEM:

STORAGE:

Records will be stored on microfilm.

RETRIEVABILITY:

Records will be retrieved by the individuals's name and other identifying information (e.g., mother's name and date of birth).

SAFEGUARDS:

Access to records in the system will be restricted to personnel who need them in the performance of their official duties. Also, the information will be maintained in the secured facilities and kept from access by unauthorized individuals (e.g., stored in locked filing cabinets) when not in use.

RETENTION AND DISPOSAL:

Records in the system will be updated biennially. Out-of-date microfilm records will be disposed of by the application of heat.

SYSTEM MANAGER(S) AND ADDRESS:

Managers of Social Security district/branch offices in the State of Kentucky. Individuals seeking office addresses and telephone numbers should consult Kentucky telephone directories.

NOTIFICATION PROCEDURE:

An individual wishing to find out if this system of records contains information about him/her may do so by contacting any Social Security office and furnishing his/her name, date and

place of birth and mother's maiden name. These procedures are in accordance with HHS Regulations 45 CFR Part 5b.

RECORD ACCESS PROCEDURE:

Same as notification procedures above. Also, individuals requesting access to their records should reasonably describe the records they are seeking. These procedures are in accordance with HHS Regulations 45 CFR Part 5b.

CONTESTING RECORD PROCEDURES:

Same as notification procedures above. Also, individuals contesting the contents of records in the system should reasonably describe the records, specify the information they are contesting and state the corrective action sought with supporting justification showing how the records are untimely, incomplete, inaccurate or irrelevant. These procedures are in accordance with HHS Regulations 45 CFR Part 5b.

RECORD SOURCE CATEGORIES:

Records in the system will be obtained from the Kentucky Office of Vital Statistics.

SYSTEMS EXEMPTED FROM CERTAIN PROVISIONS OF THE ACT:

None.

[FR Doc. 87-8255 Filed 4-13-87; 8:45 am]

BILLING CODE 4190-11-M

DEPARTMENT OF HOUSING AND URBAN DEVELOPMENT

[Docket No. N-87-1692]

Agency Information Collection Activities Under OMB Review

AGENCY: Office of Administration, HUD.
ACTION: Notices.

SUMMARY: The proposed information collection requirements described below have been submitted to the Office of Management and Budget (OMB) for review, as required by the Paperwork Reduction Act. The Department is soliciting public comments on the subject proposals.

ACTION: Interested persons are invited to submit comments regarding these proposals. Comments should refer to the proposal by name and should be sent to: John F. Morrall, OMB Desk Officer, Office of Management and Budget, New Executive Office Building, Washington, DC 20503.

FOR FURTHER INFORMATION CONTACT: David S. Cristy, Reports Management Officer, Department of Housing and Urban Development, 451, 7th Street SW.,

Washington, DC 20410, telephone (202) 755-6050. This is not a toll-free number.

SUPPLEMENTARY INFORMATION: The Department has submitted the proposal described below for the collection of information to OMB for review, as required by the Paperwork Reduction Act (44 U.S.C. Chapter 35).

The Notice lists the following information: (1) The title of the information collection proposal; (2) the office of the agency to collect the information; (3) the description of the need for the information and its proposed use; (4) the agency form number, if applicable; (5) what members of the public will be affected by the proposal; (6) how frequently information submissions will be required; (7) an estimate of the total number of hours needed to prepare the information submission; (8) whether the proposal is new, an extension, reinstatement, or revision of an information collection requirement; and (9) the names and telephone numbers of an agency official familiar with the proposal and of the OMB Desk Officer for the Department.

Copies of the proposed forms and other available documents submitted to OMB may be obtained from David S. Cristy, Reports Management Officer for the Department. His address and telephone number are listed above. Comments regarding the proposal should be sent to the OMB Desk Officer at the address listed above.

The proposed information collection requirement is described as follows:

Notice of Submission of Proposed Information Collection to OMB

Proposal: Request for Exceptions to Income Limits for Public and Indian Housing Programs, 24 CFR 913.105(B)
Office: Public and Indian Housing
Description of the need for the information and its proposed use: PHAs and IHAs may request approval from HUD for admission of families with incomes greater than fifty percent of the area median to units which were available for occupancy under contracts executed under the U.S. Housing Act of 1937 on or after October 1, 1981. HUD will maintain the five percent statutory cap and approve requests which comply with the regulation. Without a procedure to grant exceptions, it would be impossible to comply with the five percent statutory cap
Form Number: None
Respondent: State or Local Governments and Non-Profit Institutions
Frequency of Response: On Occasion
Estimated Burden Hours: 280