DATES: Submit written or electronic comments on the collection of information by May 30, 2002.

ADDRESSES: Submit electronic comments on the collection of information to: *www.mtolson@aoa.gov.* Submit written comments on the collection of information to Administration on Aging, 330 Independence Avenue, SW., Washington, DC 20201.

FOR FURTHER INFORMATION CONTACT: Margaret A. Tolson at (202) 401–0838.

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.

The Certification of Maintenance of Effort form will be used by the Administration on Aging to verify the amount of State expenditures for Title III of the Older Americans Act, and make comparisons with such expenditures for the three previous years to assure that the State Agency on Aging is in compliance with 45 CFR 1321.49.

This information will be used for federal oversight of the Title III program. AoA estimates the burden of this collection of information as follows: ½ hour per State Agency on Aging annually, for a total of 28 hours.

Dated: March 7, 2003.

Josefina G. Carbonell,

Assistant Secretary for Aging.

[FR Doc. 02–5856 Filed 3–11–02; 8:45 am]

BILLING CODE 4154–01–P

DEPARTMENT OF HEALTH AND HUMAN SERVICES

Food and Drug Administration

Advisory Committee; Renewals

AGENCY: Food and Drug Administration, HHS.

ACTION: Notice.

SUMMARY: The Food and Drug Administration (FDA) is announcing the renewal of certain FDA advisory committees by the Commissioner of Food and Drugs (the Commissioner). The Commissioner has determined that it is in the public interest to renew the charters of the committees listed below for an additional 2 years beyond charter expiration date. The new charters will be in effect until the dates of expiration listed below. This notice is issued under the Federal Advisory Committee Act of October 6, 1972 (Pub. L. 92–463 (5 U.S.C. app. 2)).

DATES: Authority for these committees will expire on the date indicated below unless the Commissioner formally determines that renewal is in the public interest.

Name of committee	Date of expiration
Antiviral Drugs Advisory Committee.	Feb. 15, 2003.
National Mammography Quality Assurance Advisory Committee,	July 6, 2003.
Nonprescription Drugs Advisory Committee.	Oct. 27, 2003.
Advisory Committee on Special Studies Relating to the Possible Long-Term Health Effects of Phenoxy Herbicides and Contaminants.	Dec. 2, 2003.
Food Advisory Committee	Dec. 18, 2003.
Vaccines and Related Biological Products Advisory. Committee	Dec. 31, 2003.
Advisory Committee for Pharmaceutical Science.	Jan. 22, 2004.
Pharmacy Compounding Advisory Committee.	Feb. 3, 2004.

FOR FURTHER INFORMATION CONTACT: Linda A. Sherman, Advisory Committee Oversight and Management Staff (HF–4), Food and Drug Administration, 5600 Fishers Lane, Rockville, MD 20857, 301–827–1220.

Dated: March 5, 2002.

Linda A. Suydam,

Senior Associate Commissioner for Communications and Constituent Relations.

[FR Doc. 02–5791 Filed 3–11–02; 8:45 am]

BILLING CODE 4160–01–P

DEPARTMENT OF HEALTH AND HUMAN SERVICES

Health Resources and Services Administration

National Vaccine Injury Compensation Program; List of Petitions Received

AGENCY: Health Resources and Services Administration, HHS.

ACTION: Notice.

SUMMARY: The Health Resources and Services Administration (HRSA) is publishing this notice of petitions received under the National Vaccine Injury Compensation Program (''the Program''), as required by section 2112(b)(2) of the Public Health Service (PHS) Act, as amended. While the Secretary of Health and Human Services is named as the respondent in all proceedings brought by the filing of petitions for compensation under the Program, the United States Court of Federal Claims is charged by statute with responsibility for considering and acting upon the petitions.

FOR FURTHER INFORMATION CONTACT: For information about requirements for filing petitions, and the Program in general, contact the Clerk, United States Court of Federal Claims, 717 Madison Place, NW., Washington, DC 20005, (202) 219–9657. For information on HRSA's role in the Program, contact the Director, National Vaccine Injury Compensation Program, 5600 Fishers Lane, Room 8A–46, Rockville, MD 20857; (301) 443–6593.

SUPPLEMENTARY INFORMATION: The Program provides a system of no-fault compensation for certain individuals who have been injured by specified childhood vaccines. Subtitle 2 of title XXI of the PHS Act, 42 U.S.C. 300aa–10 *et seq.*, provides that those seeking compensation are to file a petition with the U.S. Court of Federal Claims and to serve a copy of the petition on the Secretary of Health and Human Services, who is named as the respondent in each proceeding. The Secretary has delegated his responsibility under the Program to HRSA. The Court is directed by statute to appoint special masters who take