DEPARTMENT OF HEALTH AND HUMAN SERVICES

Administration for Children and Families

[Program Announcement No. ACYF–HS–93600–97–03–1]

Administration on Children, Youth and Families; Early Head Start Program Grant Availability

AGENCY: Administration on Children, Youth and Families, ACF, DHHS.

ACTION: Correction notice.

SUMMARY: This notice corrects the announcement of the availability of financial assistance and request for applications for the Early Head Start program, published in the **Federal Register** on April 17, 1997. Five of the geographic areas noted in Appendix D of the announcement under funding Category One are being changed.

FOR FURTHER INFORMATION CONTACT: Mireille Kanda (202) 205–8308.

SUPPLEMENTARY INFORMATION: On April 17, 1997, the Administration for Children and Families published in the **Federal Register** a notice which announced the availability of funds for competing applications for Early Head Start (62 FR 18966–19005). The purpose of this program is to provide early, continuous, intensive, and comprehensive child development and family support services on a year-round basis to low-income families with children under age three and pregnant women.

Geographic Areas

Georgia

The original citation of the ''Counties of Dekalb, Scottsdale and Decatur'' should be changed to ''Within Dekalb County the communities of Scottsdale, Decatur and that section of Atlanta in Dekalb County.''

Michigan

The original citation of ''Menominee, Delta and Schoolcraft Counties'' should be changed to ''Delta County.''

Texas

The original citation of ''Northeast Dallas'' should be changed to ''Southeast Dallas.''

Utah

Originally we listed Box Elder and Cache Counties in Utah and Franklin County in Idaho as served areas. This should be corrected. These counties are deleted from the list of served areas and are open now for competition under Category I.

Wisconsin

Originally the Counties of Barron, Chippewa, Dunn, Pepin, Pierce, Polk, and St. Croix were listed as served areas. This should be corrected. These counties are deleted from the list of served areas and are open now for competition under Category I.

Dated: May 15, 1997.

James A. Harrell,

Acting Commissioner, Administration on Children, Youth and Families.

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DEPARTMENT OF HEALTH AND HUMAN SERVICES

Food and Drug Administration

[Docket No. 97N–0201]

Agency Information Collection Activities: Proposed Collection; Comment Request

AGENCY: Food and Drug Administration, HHS.

ACTION: Notice.

SUMMARY: The Food and Drug Administration (FDA) is announcing an opportunity for public comment on the proposed collection of certain information by the agency pursuant to the Paperwork Reduction Act of 1995 (the PRA). This notice solicits comments on a data collection effort consisting of four consumer surveys regarding preferences for, and comprehension of information contained in different formats and methods for communication in over-the-counter (OTC) drug labels. For two of these studies (studies A and B), the agency has requested emergency processing of the proposed collection by the Office of Management and Budget (OMB).

DATES: Submit written comments on the collection of information for studies A and B by June 2, 1997. Submit written comments on the collection of information for studies C and D by July 22, 1997.

ADDRESSES: Submit written comments on the collection of information for studies A and B to the Office of Information and Regulatory Affairs, OMB, New Executive Office Bldg., 725 17th St. NW., rm. 10235, Washington, DC 20503, Attn: Desk Officer for FDA. Submit written comments on the collection of information for studies C and D to the Dockets Management Branch (HFA–305), ATTN: OTC Drug Labeling Data Collection, Food and Drug Administration, 12420 Parklawn Dr., rm. 1–23, Rockville, MD 20857. All comments should be identified with the docket number found in brackets in the heading of this document.

FOR FURTHER INFORMATION CONTACT: Denver Presley, Office of Information Resources Management (HFA–250), Food and Drug Administration, 5600 Fishers Lane, rm. 16B–19, Rockville, MD 20857, 301–827–1472.

SUPPLEMENTARY INFORMATION: Under the PRA (44 U.S.C. 3501–3520), Federal agencies must obtain approval from 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 requests 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 before submitting the collections to OMB for approval. Section 3507(j) of the P.A. and 5 CFR 1320.12 provides for emergency processing of proposed collection of information.

FDA intends to conduct four separate studies related to the labeling of OTC drug products. For studies A and B, the agency is requesting emergency processing because the information is necessary for the agency's deliberations on a proposed rule related to providing easier to read and easier to understand labeling on OTC drug products. (See 62 FR 9024.) The agency has determined that there is a public health need for revised OTC labeling, which is essential to the agency's mission, and if normal clearance procedures were followed, it would take longer to conclude the related OTC labeling rulemaking.

To comply with the PRA requirements, FDA is publishing notice of the proposed collections of information listed below.

With respect to the following collections of information, FDA invites comments on: (1) Whether the proposed collections of information are necessary for the proper performance of FDA's functions, including whether the information will have practical utility; (2) the accuracy of FDA's estimates of the burdens of the proposed collections 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 burdens of the collections of information on respondents, including through the use of automated collection