industry members, consumer advocates, and government representatives, are requested to submit written comments on any issue of fact, law, or policy addressed at the workshop.

DATES: Written comments may be submitted between June 8, 1999 and July 1, 1999.

COMMENT SUBMISSION PROCEDURE: Written comments should be submitted to: Secretary, Federal Trade Commission, Room H–159, 600 Pennsylvania Ave., N.W., Washington, D.C., 20580. The Commission requests that commenters submit the original plus five copies, if feasible. To enable prompt review and accessibility to the public, responses also should be submitted, if possible, in electronic form, on either one 5¼ or one 3½ inch computer disk, with a disk label stating the name of the submitter and the name and version of the word processing program used to create the document. (Programs based on DOS or Windows are preferred. Files from other operating systems should be submitted in ASCII tex format.) Alternatively, the Commission will accept responses submitted to the following e-mail address <EMarketplace@ftc.gov>. All submissions should be captioned: ''U.S. Perspectives on Consumer Protection in the Global electronic Marketplace—Comment, P994312.''

FOR FURTHER INFORMATION: A complete and current agenda, including the list of participants, and all public comments submitted in connection with the workshop can be found at the Federal Trade Commission Web site at <http://www.ftc.gov/bcp/icpw>. For further questions about the workshop, contact either: Lisa Rosenthal, Legal Advisor for International Consumer Protection, Division of Planning and Information, Bureau of Consumer Protection, Federal Trade Commission, 600 Pennsylvania Avenue, N.W., Washington, D.C. 20580, telephone 202–326–2249, e-mail<lrosenthal@ftc.gov>; or Jonathan Smollen, Attorney, Division of Financial Practices, Bureau of Consumer Protection, Federal Trade Commission, 600 Pennsylvania Avenue, N.W., Washington, D.C. 20580, telephone 202–326–3457, e-mail <jsmollen@ftc.gov>.

Authority: 15 U.S.C. 41 *et seq.*

By direction of the Commission.

Donald S. Clark,

Secretary.

[FR Doc. 99–14550 Filed 6–4–99; 1:53 pm]

BILLING CODE 6750–01–M

GENERAL SERVICES ADMINISTRATION

[GSA Bulletin FPMR D–242, Supplement 1]

Placement of Commercial Antennas on Federal Property

AGENCY: Office of Governmentwide Policy, GSA.

ACTION: Notice.

SUMMARY: This supplement extends the expiration date of GSA Bulletin D–242, Placement of Commercial Antennas on Federal Property, published in the **Federal Register** on June 16, 1997 (62 FR 32611). The expiration date of the bulletin, June 30, 1999, has been extended indefinitely. The bulletin contains information of a continuing nature and will remain in effect until specifically canceled.

FOR FURTHER INFORMATION CONTACT: Stanley C. Langfeld, Director, Real Property Policy Division, at 202–501–1737.

Dated: May 28, 1999.

David L. Bibb,

Acting Associate Administrator, Office of Governmentwide Policy.

[FR Doc. 99–14335 Filed 6–7–99; 8:45 am]

BILLING CODE 6820–34–P

GENERAL SERVICES ADMINISTRATION

[GSA Bulletin FPMR D–246, Supplement 1]

Assessment of Fees and Recovery of Costs for Antennas of Federal Agencies and Public Service Organizations

AGENCY: Office of Governmentwide Policy, GSA.

ACTION: Notice.

SUMMARY: This supplement extends the expiration date of GSA Bulletin D–246, Assessment of Fees and Recovery of Costs for Antennas of Federal Agencies and Public Service Organizations, published in the **Federal Register** on March 4, 1998 (63 FR 10631). The expiration date of the bulletin, June 30, 1999, has been extended indefinitely. The bulletin contains information of a continuing nature and will remain in effect until specifically canceled.

FOR FURTHER INFORMATION CONTACT: Stanley C. Langfeld, Director, Real Property Policy Division, at 202–501–1737.

Dated: May 28, 1999.

David L. Bibb,

Acting Associate Administrator, Office of Governmentwide Policy.

[FR Doc. 99–14336 Filed 6–7–99; 8:45 am]

BILLING CODE 6820–34–P

DEPARTMENT OF HEALTH AND HUMAN SERVICES

Administration for Children and Families

Reallotment of FY 1998 Funds for Low Income Home Energy Assistance Program (LIHEAP)

AGENCY: Office of Community Services, ACF, DHHS.

ACTION: Notice of determination concerning funds available for reallotment.

SUMMARY: Notice is hereby given that a preliminary determination has been made that fiscal year (FY) 1998 Low Income Home Energy Assistance Program (LIHEAP) funds are available for reallotment *to States, territories, and Tribes and tribal organizations receiving FY 1999 direct LIHEAP funding.* No subgrantees or other entities may apply for the funds. Section 2607(b)(1) of the Low Income Home Energy Assistance Act (the Act), Title XXVI of the Omnibus Budget Reconciliation Act of 1981 (42 U.S.C. 8621 et seq.), as amended, requires that if the Secretary of the Department of Health and Human Services determines that, as of September 1 of any fiscal, an amount in excess of certain levels allotted to a grantee for any fiscal year will not be used by the grantee during the fiscal year, the Secretary must notify the grantee and publish a notice in the **Federal Register** that such funds may be reallotted to LIHEAP grantees during the following fiscal year. If reallotted, the LIHEAP block grant allocation formula will be used to distribute the funds. (No funds may be allotted to entities that are not direct LIHEAP grantees during FY 1999.) It has been determined that $2,381,450.52 may be available for reallotment during FY 1999. This determination is based on revised reports from the State of North Carolina and the Delaware Tribe of Oklahoma, which were submitted to the Office of Community Services as required by 45 CFR 96.82.

The statute allows grantees who have funds unobligated at the end of the fiscal year for which they are awarded to request that they be allowed to carry over up to 10 percent of their allotments to the next fiscal year. Funds in excess

of this amount must be returned to DHHS and are subject to reallotment under section 2607(b)(1) of the Act. The amount described in this notice was reported as unobligated FY 1998 funds in excess of the amount that the State of North Carolina and the Delaware Tribe of Oklahoma could carry over to FY 1999.

The State of North Carolina was notified by certified mail that $2,375,000 of its FY 1998 funds may be reallotted. Additionally, the Delaware Tribe of Oklahoma was notified by certified mail that $6,450.52 of its FY 1998 funds may be reallotted. In accordance with section 2607(b)(3), the Chief Executive Officers of the State of North Carolina and of the Delaware Tribe of Oklahoma have 30 days from the date of the letter to submit comments to: Donald Sykes, Director, Office of Community Services, 3701 L'Enfant Promenade, SW, Washington, DC 20047. The comment period expires July 8, 1999.

After considering any comments submitted, the Chief Executive Officers will be notified of the decision, and the decision also will be published in the **Federal Register.** If funds are reallotted, they will be allocated in accordance with section 2604 of the Act and must be treated by LIHEAP grantees receiving them as an amount appropriated for FY 1999. As FY 1999 funds, they will be subject to all requirements of the Act, including section 2607(b)(2), which requires that a grantee obligate at least 90% of its total block grant allocation for a fiscal year by the end of the fiscal year for which the funds are appropriated, that is, by September 30, 1999.

FOR FURTHER INFORMATION CONTACT: Janet Fox, Director, Division of Energy Assistance, Office of Community Services, 370 L'Enfant Promenade, SW, Washington, DC 20447; telephone (202) 401–9351.

Dated: May 28, 1999.

Donald Sykes,

Director, Office of Community Services.

[FR Doc. 99–14480 Filed 6–7–99; 8:45 am]

BILLING CODE 99–4184–01–M

DEPARTMENT OF HEALTH AND HUMAN SERVICES

Food and Drug Administration

[Docket No. 99N–1522]

Agency Information Collection Activities: Proposed Collection; Comment Request; Temporary Marketing Permit Applications

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. 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 reporting requirements contained in existing FDA regulations governing temporary marketing permit applications.

DATES: Submit written comments on the collection of information by August 9, 1999.

ADDRESSES: Submit written comments on the collection of information to the Dockets Management Branch (HFA–305), Food and Drug Administration, 5630 Fishers Lane, rm. 1061, Rockville, MD 20852. All comments should be identified with the docket number found in brackets in the heading of this document.

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

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 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, including each proposed extension of an existing collection of information, before submitting the collection to OMB for approval. To comply with this requirement, FDA is publishing notice of the proposed collection of information set forth below.

With respect to the following collection of information, FDA invites comments on: (1) Whether the proposed collection of information is necessary for the proper performance of FDA's functions, including whether the information will have practical utility; (2) the accuracy of FDA'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.

Temporary Marketing Permit Applications—21 CFR 130.17(c) and (i) (OMB Control Number 0910–0133—Extension)

Section 401 of the Federal Food, Drug, and Cosmetic Act (the act) (21 U.S.C. 341) directs FDA to issue regulations establishing definitions and standards of identity for food ''whenever * * * such action will promote honesty and fair dealing in the interest of consumers.'' Under section 403(g) of the act (21 U.S.C. 343(g)), a food that is subject to a definition and standard of identity prescribed by regulation is misbranded if it does not conform to such definition and standard of identity. Section 130.17 (21 CFR 130.17) provides for the issuance by FDA of temporary marketing permits that enable the food industry to test consumer acceptance and measure the technological and commercial feasibility in interstate commerce of experimental packs of food that deviate from applicable definitions and standards of identity. Section 130.17(c) specifies the information that a firm must submit to FDA to obtain a temporary marketing permit. The information required in a temporary marketing permit application under § 130.17(c) enables the agency to monitor the manufacture, labeling, and distribution of experimental packs of food that deviate from applicable definitions or standards of identity. The information so obtained can be used in support of a petition to establish or amend the applicable definition or standard of identity to provide for the variations. Section 130.17(i) specifies the information that a firm must submit