FOR FURTHER INFORMATION CONTACT: Sandra M. Peay or Renee A. Horton, Contact Representatives, Federal Trade Commission, Premerger Notification Office, Bureau of Competition, Room 303, Washington, DC 20580, (202) 326–3100.

By direction of the Commission.

Donald S. Clark,

Secretary.

[FR Doc. 95–1960 Filed 1–25–95; 8:45 am]

BILLING CODE 6750–01–M

DEPARTMENT OF HEALTH AND HUMAN SERVICES

Administration for Children and Families

Agency Information Collection Under OMB Review

Under the provisions of the Paperwork Reduction Act (44 U.S.C. Chapter 35), the Administration for Children and Families (ACF) has submitted to the Office of Management and Budget (OMB) a request to extend the prior approval of the paperwork burden associated with the application required for the Head Start Program Information Report.

This request is being made to extend the OMB authorization of the Program Information Report to June 30, 1996. There is no change in the previously approved paperwork burden.

ADDRESSES: Copies of the request for approval may be obtained from Bob Sargis of the Office of Information Systems Management, ACF, by calling (202) 690–7275.

Consideration will be given to comments and suggestions received March 1, 1995. Written comments and recommendations for the proposed information should be sent directly to the following: Wendy Taylor, OMB Desk Officer for ACF, New Executive Office Building, Room 10235, 725 17th Street, N.W., Washington, D.C. 20503 (202) 395–7316

Information on Document

Title: Head Start Program Information Report

OMB No.: 0980–0017

Description: This Program Information Report is used to collect data necessary to evaluate the services delivered to participating children and families.

Respondents: States and Territories

Annual Number of Respondents: 2006 sites

Total annual responses: 2006 sites

Hours per response: 3.5

Total Burden Hours: 7,021

Dated: January 18, 1995.

Larry Guerrero,

Deputy Director, Office of Information Systems Management.

[FR Doc. 95–1912 Filed 1–25–95; 8:45 am]

BILLING CODE 4184–01–M

Food and Drug Administration

[Docket No. 91F–0324]

Goodyear Tire & Rubber Co.; Filing of Food Additive Petition; Amendment

AGENCY: Food and Drug Administration, HHS.

ACTION: Notice.

SUMMARY: The Food and Drug Administration (FDA) is amending the filing notice for a food additive petition filed by Goodyear Tire & Rubber Co. to indicate that the petitioned additive, alkylthiophendics, acid-catalyzed condensation reaction products of *p*-nonylphenol, formaldehyde, and 1-dodecanethiol, is also intended for use in pressure-sensitive adhesives. The previous filing notice indicated that the additive was intended for use only as an antioxidant for adhesives and repeat-use rubber articles.

DATES: Written comments on the petitioner's environmental assessment by February 27, 1995.

ADDRESSES: Submit written comments to the Dockets Management Branch (HFA–305), Food and Drug Administration, rm. 1–23, 12420 Parklawn Dr., Rockville, MD 20857.

FOR FURTHER INFORMATION CONTACT: Andrew J. Zajac, Center for Food Safety and Applied Nutrition (HFS–216), Food and Drug Administration, 200 C St. SW., Washington, DC 20204, 202–418–3095.

SUPPLEMENTARY INFORMATION: In a notice published in the **Federal Register** of September 12, 1991 (56 FR 46439), FDA announced that a food additive petition (FAP 1B4259) had been filed by Goodyear Tire & Rubber Co., Akron, OH 44316–0001 (currently 1001 G St. N.W., Suite 500 West, Washington, DC 20001), proposing that § 178.2010 *Antioxidants and/or stabilizers for polymers* (21 CFR 178.2010) be amended to provide for the safe use of the acid-catalyzed condensation reaction product of *p*-nonylphenol, formalin, and 1-dodecanethiol as an antioxidant for adhesives, listed under 21 CFR 175.105, and rubber articles, listed under 21 CFR 177.2600, intended for repeated use in food packaging.

Upon further review of the petition, the agency notes that the additive is specifically intended for use in pressure-sensitive adhesives rather than adhesives. However, the petitioner has subsequently amended the petition to also include the use of the additive in adhesives. In addition, the agency is also modifying the nomenclature for clarification. Therefore, FDA is amending the filing notice of September 12, 1991, to state that the petitioner requests that § 178.2010 *Antioxidants and/or stabilizers for polymers* be amended to provide for the safe use of alkylthiophendics formed by the acid-catalyzed condensation reaction of *p*-nonylphenol, formaldehyde, and 1-dodecanethiol as an antioxidant for adhesives, listed under 21 CFR 175.105, pressure-sensitive adhesives, listed under 21 CFR 175.125, and repeat-use rubber articles, listed under 21 CFR 177.2600.

The potential environmental impact of this action is being reviewed. To encourage public participation consistent with regulations promulgated under the National Environmental Policy Act (40 CFR 1501.4(b)), the agency is placing the environmental assessment submitted with the petition that is the subject of this notice on public display at the Dockets Management Branch (address above) for public review and comment. Interested persons may, on or before February 27, 1995, submit to the Dockets Management Branch (address above) written comments. Two copies of any comments are to be submitted, except that individuals may submit one copy. Comments are to be identified with the docket number found in brackets in the heading of this document. Received comments may be seen in the office above between 9 a.m. and 4 p.m., Monday through Friday. FDA will also place on public display any amendments to, or comments on, the petitioner's environmental assessment without further announcement in the **Federal Register**. If, based on its review, the agency finds that an environmental impact statement is not required and this petition results in a regulation, the notice of availability of the agency's finding of no significant impact and the evidence supporting that finding will be published with the final regulation in the **Federal Register** in accordance with 21 CFR 25.40(c).

Dated: January 18, 1995.

Alan M. Rulis,

Acting Director, Office of Premarket Approval, Center for Food Safety and Applied Nutrition.

[FR Doc. 95–2007 Filed 1–25–95; 8:45 am]

BILLING CODE 4160–01–F