

**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

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**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]

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**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]

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