

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