

the safe use of chromium antimony titanium buff rutile (C.I. Pigment Brown 24) as a colorant for polymers intended for use in contact with food.

**FOR FURTHER INFORMATION CONTACT:** Vir D. Anand, Center for Food Safety and Applied Nutrition (HFS-215), Food and Drug Administration, 200 C St. SW., Washington, DC 20204, 202-418-3081.

**SUPPLEMENTARY INFORMATION:** Under the Federal Food, Drug, and Cosmetic Act (sec. 409(b)(5) (21 U.S.C. 348(b)(5))), notice is given that a food additive petition (FAP 8B4608) has been filed by BASF Corp., 3000 Continental Dr. North, Mt. Olive, NJ 07828-1234. The petition proposes to amend the food additive regulations to provide for the safe use of chromium antimony titanium buff rutile (C.I. Pigment Brown 24) as a colorant for polymers intended for use in contact with food.

The agency has determined under 21 CFR 25.32(i) that this action is of the type that does not individually or cumulatively have a significant effect on the human environment. Therefore, neither an environmental assessment nor an environmental impact statement is required.

Dated: July 6, 1998.

**Laura M. Tarantino,**

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

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## DEPARTMENT OF HEALTH AND HUMAN SERVICES

### Food and Drug Administration

[Docket No. 98F-0569]

#### Ticona; Filing of Food Additive Petition

**AGENCY:** Food and Drug Administration, HHS.

**ACTION:** Notice.

**SUMMARY:** The Food and Drug Administration (FDA) is announcing that Ticona has filed a petition proposing that the food additive regulations be amended to provide for the safe use of ethylene-norbornene copolymers as articles or components of articles in contact with dry food.

**DATES:** Written comments on the petitioner's environmental assessment by August 24, 1998.

**ADDRESSES:** Submit written comments to the Dockets Management Branch (HFA-305), Food and Drug Administration, 5630 Fishers Lane, rm. 1061, Rockville, MD 20852.

**FOR FURTHER INFORMATION CONTACT:** Julius Smith, Center for Food Safety and Applied Nutrition (HFS-215), Food and Drug Administration, 200 C St. SW., Washington, DC 20204, 202-418-3091.

**SUPPLEMENTARY INFORMATION:** Under the Federal Food, Drug, and Cosmetic Act (sec. 409(b)(5) (21 U.S.C. 348(b)(5))), notice is given that a food additive petition (FAP 8B4597) has been filed by Ticona, c/o Keller and Heckman, 1001 G St. NW., suite 500 West, Washington, DC 20001. The petition proposes to amend the food additive regulations in § 177.1520 *Olefin polymers* (21 CFR 177.1520) to provide for the safe use of ethylene-norbornene copolymers as articles or components of articles in contact with dry foods.

The potential environmental impact of this action is being reviewed. To encourage public participation consistent with regulations issued 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 August 24, 1998, 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 regulation in the **Federal Register** in accordance with 21 CFR 25.40(c).

Dated: July 6, 1998.

**Laura M. Tarantino,**

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

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## DEPARTMENT OF HEALTH AND HUMAN SERVICES

### Health Care Financing Administration

[HCFA-64, 64.21, 64.21U, 64.21P, 64.21UP, 64EC, 64.21E, 64.9P, 64.10P, 64.11A, 64.9d]

#### Agency Information Collection Activities: Proposed Collection; Comment Request

In compliance with the requirement of section 3506(c)(2)(A) of the Paperwork Reduction Act of 1995, the Health Care Financing Administration (HCFA), Department of Health and Human Services, is publishing the following summary of proposed collections for public comment. Interested persons are invited to send comments regarding the burden estimate or any other aspect of this collection of information, including any of the following subjects: (1) the necessity and utility of the proposed information collection for the proper performance of the agency's functions; (2) the accuracy of the estimated burden; (3) ways to enhance the quality, utility, and clarity of the information to be collected; and (4) the use of automated collection techniques or other forms of information technology to minimize the information collection burden.

#### Type of Information Collection

**Request:** Revision of a currently approved collection;

#### Title of Information Collection:

Quarterly Medicaid Statement of Expenditures for the Medical Assistance Program.

**Form Nos.:** HCFA-64, 64.21, 64.21U, 64.21P, 64.21UP, 64EC, 64.21E, 64.9, 64.10, 64.10P, 64.11a, 64.9d;

**Use:** These new forms are revisions of the currently approved collection report Form HCFA-64. These forms will be used by State Medicaid agencies to report their actual CHIP-related Medicaid expenditures and the numbers of CHIP-related children, and other children being served in the Medicaid program, to the Health Care Financing Administration (HCFA). The forms will be used by the HCFA to ensure that the appropriate level of Federal payments for the State's CHIP-related Medicaid program expenditures are made in accordance with the CHIP and related Medicaid provisions of the BBA of 1997, and to track, monitor, and evaluate the numbers of CHIP-related children and other individuals being served by the Medicaid program.

**Note:** at this time Forms HCFA-64.21E and HCFA-64EC of this package are for States to report the numbers of CHIP-related children and other