

Towanda and the townships of Wysox, North Towanda, and Monroe; (2) the Mount Carmel, Pennsylvania area, which includes the Borough of Mount Carmel and the Township of Mount Carmel; and (3) the Pittston, Pennsylvania area, which includes the city of Pittston, the townships of Pittston and Jenkins, and the boroughs of Dupont, Avoca, Hughestown, Duryea, Yatesville, and Laflin, Pennsylvania. According to the draft complaint, these markets are highly concentrated and entry is difficult or unlikely. Penn Traffic's acquisition of Acme may reduce competition in these markets by eliminating the direct competition between Penn Traffic and Acme, by increasing the likelihood that Penn Traffic will become a dominant firm, and by increasing the likelihood of collusive behavior among the few remaining competitors.

The agreement containing consent order attempts to remedy the Commission's competitive concerns about the acquisition. Under the terms of the proposed order, Penn Traffic must divest three supermarkets within twelve-months, to a purchaser approved by the Commission. The three stores to be divested include the "Acme" supermarket located in Towanda, Pennsylvania, the "Acme" supermarket located in Pittston, Pennsylvania, and either the "Acme" or the Penn Traffic store located in Mount Carmel, Pennsylvania.

For a period of ten years from the date the order becomes final, the order also prohibits Penn Traffic from acquiring, without prior Commission approval, stock, or any other interest in any supermarket, or entity that owns or operates a supermarket, located in the areas of Towanda, Pittston, or Mount Carmel, Pennsylvania. This prohibition will not apply to the construction of new facilities or the leasing of facilities not operated as supermarkets within six months of the offer to lease.

The purpose of this analysis is to invite public comment concerning the consent order and any other aspect of this matter. This analysis is not intended to constitute an official interpretation of the agreement and order or to modify its terms in any way.

Donald S. Clark,

Secretary.

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DEPARTMENT OF HEALTH AND HUMAN SERVICES

Food and Drug Administration

[Docket No. 95F-0016]

Johnson Matthey Chemicals; Filing of Food Additive Petition

AGENCY: Food and Drug Administration, HHS.

ACTION: Notice.

SUMMARY: The Food and Drug Administration (FDA) is announcing that Johnson Matthey Chemicals has filed a petition proposing that the food additive regulations be amended to provide for the safe use of silver chloride coated titanium dioxide.

DATES: Written comments on the petitioner's environmental assessment by March 15, 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: Diane E. Robertson, Center for Food Safety and Applied Nutrition (HFS-216), Food and Drug Administration, 200 C St. SW., Washington, DC 20204, 202-418-3089.

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 5B4442) has been filed by Johnson Matthey Chemicals, c/o 1000 Potomac St. NW., Washington, DC 20007. The petition proposes to amend the food additive regulations in § 176.170 *Components of paper and paperboard in contact with aqueous and fatty foods* (21 CFR 176.170) to provide for the safe use of silver chloride coated titanium dioxide.

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 March 15, 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 regulation in the **Federal Register** in accordance with 21 CFR 25.40(c).

Dated: February 3, 1995.

Alan M. Rulis,

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

[FR Doc. 95-3557 Filed 2-10-95; 8:45 am]

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[Docket No. 95F-0017]

Robinson Brothers Ltd.; Filing of Food Additive Petition

AGENCY: Food and Drug Administration, HHS.

ACTION: Notice.

SUMMARY: The Food and Drug Administration (FDA) is announcing that Robinson Brothers Ltd., has filed a petition proposing that the food additive regulations be amended to provide for the safe use of diisopropyl xanthogen polysulfide as a component of rubber articles intended for repeated use in contact with food.

DATES: Written comments on the petitioner's environmental assessment by March 15, 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: Diane E. Robertson, Center for Food Safety and Applied Nutrition (HFS-216), Food and Drug Administration, 200 C St. SW., Washington, DC 20204, 202-418-3089.

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 5B4437) has been filed by Robinson Brothers Ltd., Phoenix St., West Bromwich, West Midland, B70 OAH, England. The petition proposes to amend the food additive regulations in

§ 177.2600 *Rubber articles intended for repeated use* (21 CFR 177.2600) to provide for the safe use of diisopropyl xanthogen polysulfide as a component of rubber articles intended for repeated use in contact with food.

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 March 15, 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 regulation in the **Federal Register** in accordance with 21 CFR 25.40(c).

Dated: February 3, 1995.

Alan M. Rulis,

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

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HEALTH RESOURCES AND SERVICES ADMINISTRATION

Special Project Grants and Cooperative Agreements; Maternal and Child Health Services; Federal Set-Aside Program; Genetic Services and Maternal and Child Health Improvement Projects

AGENCY: Health Resources and Services Administration (HRSA), PHS.

ACTION: Notice of availability of funds.

SUMMARY: The Maternal and Child Health Bureau (MCHB), HRSA, announces that fiscal year (FY) 1995 funds are available for grants and cooperative agreements for the following activities: Maternal and Child Health (MCH) Special Projects of Regional and National Significance (SPRANS), including special MCH improvement projects (MCHIP) which contribute to the health of mothers, children, and children with special health care needs (CSHCN); and genetic disease testing, counseling and information services. All awards will be made under the program authority of section 502(a) of the Social Security Act, the MCH Federal Set-Aside Program. No new hemophilia SPRANS grants will be funded in FY 1995. Grants for MCH research and training are being announced in a separate notice.

Of the approximately \$44 million available for SPRANS activities in FY 1995 in categories covered by this announcement, about \$9.7 million will be available to support approximately 65 new and competing renewal projects at an average of \$150,000 per award for one year. The remaining funds will be used to support continuation of existing SPRANS activities. The actual amounts available for awards and their allocation may vary, depending on unanticipated program requirements and the volume and quality of applications. Awards are made for grant periods which may run from 1 to 5 years in duration. Funds for the MCH Federal Set-Aside Program are appropriated by Public Law 103-333. Revised regulations implementing the Federal Set-Aside Program (42 CFR part 51a) were published in the July 19, 1994, issue of the **Federal Register** at 59 FR 36703.

The Public Health Service (PHS) is committed to achieving the health promotion and disease prevention

objectives of Healthy People 2000, a PHS-led national activity for setting priority areas. The MCH Block Grant Federal Set-Aside Program addresses issues related to the Healthy People 2000 objectives of improving maternal, infant, child and adolescent health and developing service systems for children with special health care needs.

Potential applicants may obtain a copy of Healthy People 2000 (Full Report: Stock No. 017-001-00474-0) or Healthy People 2000 (Summary Report: Stock No. 017-001-00473-1) through the Superintendent of Documents, Government Printing Office, Washington, DC 20402-9325 (telephone: 202 783-3238).

The Public Health Service strongly encourages all grant recipients to provide a smoke-free workplace and promote the non-use of all tobacco products. This is consistent with the PHS mission to protect and advance the physical and mental health of the American people. In addition, Public Law 103-227, The Pro-Children Act Of 1994, prohibits smoking in certain facilities in which education, library, day care, regular and routine health care and early childhood development services are provided to children. Smoking must also be prohibited in indoor facilities that are constructed, operated or maintained with Federal funds.

ADDRESSES: Grant applications for the MCH SPRANS Federal Set-Aside Program must be obtained from and submitted to: Acting Chief, Grants Management Branch, Office of Program Support, Maternal and Child Health Bureau, Health Resources and Services Administration, Room 18-12, Parklawn Building, 5600 Fishers Lane, Rockville, Maryland 20857, (301) 443-1440. Applicants for all projects covered by this announcement will use application Form PHS 5161-1 with revised face page DHHS Form 424, approved by OMB under control number 0937-0189. Requests should specify the category or categories of activities for which an application is requested so that the appropriate forms, information and materials may be provided.

DATES: Deadlines for receipt of applications differ for the several categories of grants and cooperative agreements. These deadlines are as follows: