membership on the National Vaccine Advisory Committee (NVAC). This committee studies and recommends ways to encourage the availability of an adequate supply of safe and effective vaccination products in the States; recommends research priorities and other measures the Director of the National Vaccine Program should take to enhance the safety and efficacy of vaccines; advises the Director of the Program in the implementation of sections 2102, 2103, and 2104, of the PHS Act; and identifies annually for the Director of the Program the most important areas of government and non-government cooperation that should be considered in implementing sections 2102, 2103, and 2104, of the PHS Act.

Nominations are being sought for individuals engaged in vaccine research or the manufacture of vaccines or who are physicians, members of parent organizations concerned with immunizations, or representatives of State or local health agencies, or public health organizations. Federal employees will not be considered for membership. Members may be invited to serve a four-year term.

Close attention will be given to minority and female representation; therefore nominations from these groups are encouraged.

The following information is requested: name, affiliation, address, telephone number, and a current curriculum vitae. Nominations should be sent, in writing, and postmarked by March 15, 1996, to: Gloria A. Kovach, Committee Management Specialist, NVAC, National Vaccine Program Office, Centers for Disease Control and Prevention, 1600 Clifton Road, NE, M/S D28, Atlanta, Georgia 30333. Telephone or facsimile submission cannot be accepted.

Dated: February 15, 1996.

Carolyn J. Russell,

Director, Management Services and Analysis Office, Centers for Disease Control and Prevention.

[FR Doc. 96–4217 Filed 2–23–96; 8:45 am]

BILLING CODE 4163–18–M

Food and Drug Administration

[Docket No. 96F–0053]

Eastman Chemical Co.; Filing of Food Additive Petition

AGENCY: Food and Drug Administration, HHS.

ACTION: Notice.

SUMMARY: The Food and Drug Administration (FDA) is announcing that Eastman Chemical Co. has filed a petition proposing that the food additive regulations be amended to provide for the safe use of dimethyl 1,4-cyclohexanedicarboxylate as a monomer in polyester resins employed in adhesives as components of articles intended for use in contact with food.

DATES: Written comments on the petitioner's environmental assessment by March 27, 1996.

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

FOR FURTHER INFORMATION CONTACT: Vir D. Anand, Center for Food Safety and Applied Nutrition (HFS–216), 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 5B4481) has been filed by Eastman Chemical Co., P.O. Box 1994, Kingsport, TN 37662. The petition proposes to amend the food additive regulations in § 175.105 Adhesives (21 CFR 175.105) to provide for the safe use of dimethyl 1,4-cyclohexanedicarboxylate as a monomer in polyester resins employed in adhesives as components of articles intended for 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 display at the Dockets Management Branch (address above) for public review and comment. Interested persons may, on or before March 27, 1996, 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 8, 1996.

Alan M. Rulis,

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

[FR Doc. 96–4286 Filed 2–23–96; 8:45 am]

BILLING CODE 4160–01–F

[Docket No. 96F–0052]

Milliken & Co.; Filing of Food Additive Petition

AGENCY: Food and Drug Administration, HHS.

ACTION: Notice.

SUMMARY: The Food and Drug Administration (FDA) is announcing that Milliken & Co. has filed a petition proposing that the food additive regulations be amended to provide for the additional safe use of dimethyldibenzylidene sorbitol as a clarifying agent for propylene homopolymers and high-propylene copolymers articles intended for use in contact with food.

DATES: Written comments on the petitioner's environmental assessment by March 27, 1996.

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

FOR FURTHER INFORMATION CONTACT: Vir D. Anand, Center for Food Safety and Applied Nutrition (HFS–216), Food and Drug Administration, 200 C Street 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 6B4495) has been filed by Milliken & Co., 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 § 178.3295 *Clarifying agents for polymers* (21 CFR 178.3295) to provide for the additional safe use of dimethyldibenzylidene sorbitol as a clarifying agent for olefin polymers complying with § 177.1520 (21 CFR 177.1520), items 1.1, 3.1, and 3.2, for contact with food under condition of use A, described in Table 2 of § 176.170(c) of this chapter.

The potential environmental impact of this action is being reviewed. To