number. Any written comments on the minutes should be directed to Ms. Samara Weinstein, Executive Director of the Working Group, as shown above.

Signed at Washington, DC, this 20th day of April, 1999.

David Gray Ross,

Commissioner, Office of Child Support Enforcement.

[FR Doc. 99–10487 Filed 4–26–99; 8:45 am]

BILLING CODE 4184–01–M

DEPARTMENT OF HEALTH AND HUMAN SERVICES

Food And Drug Administration

[Docket No. 99F–0994]

Ciba Specialty Chemicals Corp.; Filing of Food Additive Petition

AGENCY: Food and Drug Administration, HHS.

ACTION: Notice.

SUMMARY: The Food and Drug Administration (FDA) is announcing that Ciba Specialty Chemicals Corp. has filed a petition proposing that the food additive regulations be amended to provide for the safe use of phosphorothioic acid, *O,O,O*-triphenyl ester, tert-butyl derivatives, as extreme pressure-antiwear adjuvants for lubricants intended for incidental 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 9B4657) has been filed by Ciba Specialty Chemicals Corp., 540 White Plains Rd., P.O. Box 2005, Tarrytown, NY 10591–9005. The petition proposes to amend the food additive regulations in § 178.3570 *Lubricants with incidental food contact* (21 CFR 178.3570) to provide for the safe use of phosphorothioic acid, *O,O,O*-triphenyl ester, *tert*-butyl derivatives, as extreme pressure-antiwear adjuvants for lubricants intended for incidental 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: April 2, 1999.

Laura M. Tarantino,

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

[FR Doc. 99–10447 Filed 4–26–99; 8:45 am]

BILLING CODE 4160–01–F

DEPARTMENT OF HEALTH AND HUMAN SERVICES

Food and Drug Administration

[Docket No. 98F–0635]

General Electric Co.; Withdrawal of Food Additive Petition

AGENCY: Food and Drug Administration, HHS.

ACTION: Notice.

SUMMARY: The Food and Drug Administration (FDA) is announcing the withdrawal, without prejudice to a future filing, of a food additive petition (FAP 0B4615) proposing that the food additive regulations be amended to provide for the expanded safe use of phosphorous acid, cyclic butylethyl propanediol, 2,4,6-tri-*tert*-butylphenyl ester, which may contain up to 1 percent by weight of triisopropanolamine, as an antioxidant and/or stabilizer for polypropylene 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: In a notice published in the **Federal Register** of August 5, 1998 (63 FR 41855), FDA announced that a food additive petition (FAP 8B4615) had been filed by General Electric Co., One Lexan Lane, Mt. Vernon IN 47620–9364. The petition proposed to amend the food additive regulations in § 178.2010 *Antioxidants and/or stabilizers for polymers* (21 CFR 178.2010) to provide for the expanded safe use of phosphorous acid, cyclic butylethyl propanediol, 2,4,6-tri-*tert*-butylphenyl ester, which may contain up to 1 percent by weight of triisopropanolamine, as an antioxidant and/or stabilizer for polypropylene complying with § 177.1520(c), items 1.1, 1.2, or 1.3, intended for use in contact with food. General Electric Co. has now withdrawn the petition without prejudice to a future filing (21 CFR 171.7).

Dated: April 15, 1999.

Alan M. Rulis,

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

[FR Doc. 99–10445 Filed 4–26–99; 8:45 am]

BILLING CODE 4160–01–F

DEPARTMENT OF HEALTH AND HUMAN SERVICES

Food and Drug Administration

Dental Products Panel of the Medical Devices Advisory Committee; Notice of Meeting

AGENCY: Food and Drug Administration, HHS.

ACTION: Notice.

This notice announces a forthcoming meeting of a public advisory committee of the Food and Drug Administration (FDA). At least one portion of the meeting will be closed to the public.

Name of Committee: Dental Products Panel of the Medical Devices Advisory Committee.

General Function of the Committee: To provide advice and recommendations to the agency on FDA's regulatory issues.

Date and Time: The meeting will be held on May 10, 1999, 10:30 a.m. to 6:30 p.m., and May 11, 1999, 8 a.m. to 3 p.m.

Location: Holiday Inn, Walker-Whetstone Rooms, Two Montgomery Village Ave., Gaithersburg, MD.

Contact Person: Pamela D. Scott, Center for Devices and Radiological Health (HFZ–480), Food and Drug Administration, 9200 Corporate Blvd., Rockville, MD 20850, 301–827–5283, or FDA Advisory Committee Information Line, 1–800–741–8138 (301–443–0572 in the Washington, DC area), code 12518. Please call the Information Line for up-to-date information on this meeting.

Agenda: On May 10, 1999, the committee will discuss, make recommendations, and vote on a premarket approval application (PMA) for a total temporomandibular joint (TMJ) prosthesis, which consists of the glenoid fossa prosthesis and the mandibular condyle prosthesis, for reconstruction of the TMJ. On May 11, 1999, the committee will discuss, make recommendations, and vote on a PMA that includes both a total TMJ prosthesis and a glenoid fossa prosthesis that can be used alone without the mandibular condyle prosthesis to reconstruct the TMJ. These PMA's were received in response to the final rule issued in the **Federal Register** of December 30, 1998 (63 FR 71743), requiring the filing of a PMA or a notice of completion of a