all-IPV schedule; revision of hepatitis B recommendations; consolidated Vaccines for Children resolution for hepatitis B vaccine; recommendation for use of pneumococcal conjugate vaccine; revaccination with pneumococcal polysaccharide vaccine; pneumococcal polysaccharide vaccine in adults with HIV infection; update on influenza; American Academy of Family Physicians recommendation for universal influenza vaccination starting at age 50 years; status of immunization of bone marrow transplant (BMT) recipients publication; discussion on vaccines related to bioterrorism; teaching immunization for medical education (TIME) project; recommendations for nursing home immunization: a HCFA/CDC collaboration; cost-effectiveness and economic analysis of immunization compared to other health interventions; electronic updating of ACIP recommendations; and Institute of Medicine report on priorities for vaccines development. Other matters of relevance among the committee's objectives may be discussed.

Agenda items are subject to change as priorities dictate.

CONTACT PERSON FOR MORE INFORMATION: John R. Livengood, M.D., Director, Division of Epidemiology and Surveillance, National Immunization Program, CDC, 1600 Clifton Road, NE, M/S E–61, Atlanta, Georgia 30333, telephone 404/639–8254.

The Director, Management Analysis and Services office has been delegated the authority to sign **Federal Register** notices pertaining to announcements of meetings and other committee management activities for both the Centers for Disease Control and Prevention and the Agency for Toxic Substances and Disease Registry.

Dated: May 20, 1999.

Carolyn J. Russell

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

[FR Doc. 99–13333 Filed 5–25–99; 8:45 am]

BILLING CODE 4163–18–P

DEPARTMENT OF HEALTH AND HUMAN SERVICES

Administration of Children and Families

[Program Announcement No. ACF/ACYF 99–06]

Fiscal Year 1999 Discretionary Announcement for Head Start Family Literacy Projects

AGENCY: Administration on Children, Youth and Families (ACYF), Administration for Children and Families (ACF), DHHS.

ACTION: Notice of announcement of the availability of funds and request for applications from organizations with experience in family literacy efforts to develop training and technical assistance programs in family literacy for Head Start and Early Head Start grantees.

SUMMARY: The Administration on Children, Youth and Families is making available $3,000,000 annually for each of the next five years to support one or more family literacy projects (FLPs). The project(s) funded under this effort will work cooperatively with the Head Start Bureau in designing and implementing training and technical assistance programs to support and strengthen the family literacy activities carried out by Head Start/Early Head Start grantees.

The overall goal of the family literacy project is to improve the quality, intensity and outcomes of the family literacy services provided by Head Start and Early Head Start grantees in order to increase lifelong learning for Head Start and Early Head Start children and their parents and to assist families in achieving self sufficiency. The cooperative agreement(s) will be awarded competitively to eligible applicant(s).

DATES: The closing date and time for receipt of applications is 5:00 p.m. (Eastern Time Zone).

FOR FURTHER INFORMATION CONTACT: A copy of the program announcement and Necessary application forms can be obtained by contacting: Family Literacy Projects, ACYF Operation Center, 1815 North Fort Myer Drive, Suite 300, Arlington, Virginia 22209. The telephone number is: 1–800–351–2293.

Copies of the program announcement can be downloaded from the Head Start web site at: *www.acf.dhhs.gov/programs/hsb*

Eligible Applicants

Applicants must be public or private nonprofit or for-profit organizations with the capability to implement a family literacy effort of national scope.

Project Duration

Awards, on a competitive basis, will be for a one-year budget period; project periods will be for five years.

Federal Share of Projects

Although there are no matching requirements, applicants are encouraged to provide non-Federal contributions to the project.

Statutory Authority: The Head Start Act, as amended, 42 U.S.C. 9831 *et seq.*

(Catalog of Federal Domestic Assistance. Number 93.600, Head Start)

Dated: May 20, 1999.

Patricia Montoya,

Commissioner, Administration on Children, Youth and Families.

[FR Doc. 99–13425 Filed 5–25–99; 8:45 am]

BILLING CODE 4184–01–M

DEPARTMENT OF HEALTH AND HUMAN SERVICES

Food And Drug Administration

[Docket No. 99F–1420]

Goodyear Tire and Rubber Co.; Filing of Food Additive Petition

AGENCY: Food and Drug Administration, HHS.

ACTION: Notice.

SUMMARY: The Food and Drug Administration (FDA) is announcing that Goodyear Tire and Rubber Co. has filed a petition proposing that the food additive regulations be amended to provide for the safe use of butylated reaction product of *p*-cresol and dicyclopentadiene as an antioxidant in pressure sensitive adhesives 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 9B4663) has been filed by Goodyear Tire and Rubber Co., c/o Keller and Heckman LLP, 1001 G St. NW., suite 500 West, Washington, DC 20001. The petition proposes to amend the food additive regulations in § 175.125 *Pressure-sensitive adhesives* (21 CFR 175.125) to provide for the safe use of butylated reaction product of *p*-cresol and dicyclopentadiene as an antioxidant in pressure sensitive

adhesives 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: May 5, 1999.

Alan M. Rulis,

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

[FR Doc. 99–13347 Filed 5–25–99; 8:45 am]

BILLING CODE 4160–01–F

DEPARTMENT OF HEALTH AND HUMAN SERVICES

Food And Drug Administration

[Docket No. 99F–1422]

Sumitomo Chemical Co. Ltd.; Filing of Food Additive Petition

AGENCY: Food and Drug Administration, HHS.

ACTION: Notice.

SUMMARY: The Food and Drug Administration (FDA) is announcing that Sumitomo Chemical Co., Ltd. has filed a petition proposing that the food additive regulations be amended to provide for the expanded safe use of 2,4-di-*tert*-pentyl-6-[1-(3,5-di-*tert*-pentyl-2-hydroxyphenyl)ethyl]phenyl acrylate as an antioxidant and/or stabilizer for polypropylene, polystyrene, rubber modified polystyrene, and styrene block copolymers 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 9B4661) has been filed by Sumitomo Chemical Co., Ltd., c/o Keller and Heckman LLP, 1001 G St. NW., suite 500 West, Washington, DC 20001. The petition proposes 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 2,4-di-*tert*-pentyl-6-[1-(3,5-di-*tert*-pentyl-2-hydroxyphenyl)ethyl]phenyl acrylate as an antioxidant and/or stabilizer for polypropylene, polystyrene, rubber modified polystyrene, and styrene block copolymers 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: May 5, 1999.

Alan M. Rulis,

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

[FR Doc. 99–13254 Filed 5–25–99; 8:45 am]

BILLING CODE 4160–01–F

DEPARTMENT OF HEALTH AND HUMAN SERVICES

Food and Drug Administration

Ear, Nose, and Throat Devices 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: Ear, Nose, and Throat Devices 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 June 18, 1999, 8:30 a.m. to 5:30 p.m.

Location: Corporate Bldg., conference room 020B, 9200 Corporate Blvd., Rockville, MD.

Contact Person: Harry R. Sauberman, Center for Devices and Radiological Health (HFZ–460), Food and Drug Administration, 9200 Corporate Blvd., Rockville, MD 20850, 301–594–2080, or FDA Advisory Committee Information Line, 1–800–741–8138 (301–443–0572 in the Washington, DC area), code 12522. Please call the Information Line for up-to-date information on this meeting.

Agenda: The committee will discuss generic issues relating to the safety and efficacy of middle ear amplification devices.

Procedure: Interested persons may present data, information, or views, orally or in writing, on issues pending before the committee. Written submissions may be made to the contact person by June 4, 1999. Oral presentations from the public will be scheduled between approximately 9 a.m. and 10 a.m., and for an additional 30 minutes near the end of the committee deliberations. Time allotted for each presentation may be limited. Those desiring to make formal oral presentations should notify the contact person before June 4, 1999, and submit a brief statement of the general nature of the evidence or arguments they wish to present, the names and addresses of proposed participants, and an indication of the approximate time requested to make their presentation.

Closed Committee Deliberations: On June 18, 1999, from 4:30 p.m. to 5:30 p.m., the meeting will be closed to the public to permit discussion and review of trade secret and/or confidential commercial information (5 U.S.C. 552b(c)(4)) relating to present and future agency issues.

Notice of this meeting is given under the Federal Advisory Committee Act (5 U.S.C. app. 2).

Dated: May 20, 1999.

Michael A. Friedman,

Deputy Commissioner for Operations.

[FR Doc. 99–13348 Filed 5–25–99; 8:45 am]

BILLING CODE 4160–01–F

DEPARTMENT OF HEALTH AND HUMAN SERVICES

Health Resources and Services Administration

In accordance with section 10(a)(2) of the Federal Advisory Committee Act (Public Law 92–463), announcement is made of the following National Advisory body scheduled to meet during the month of June 1999:

Name: Maternal and Child Health Research Grants Review Committee

Date: June 16–18, 1999 (Wednesday, Thursday and Friday)

Time: 8:00 a.m. to 5:00 p.m.

Place: Holiday Inn Chevy Chase, 5520 Wisconsin Avenue, Chevy Chase, Maryland 20815.

The meeting is open on Wednesday, June 16 from 9:00–10:00 a.m., and closed for the remainder of the meeting.

Agenda

The open portion of the meeting will cover opening remarks by the Acting Director, Division of Research, Training and Education, who will report on program issues, congressional activities, and other topics of interest to the field of maternal and child health. The meeting will be closed to the public on Wednesday, June 16, 1999 from 10:00 a.m., to the remainder of the meeting, for the review of grant applications. The closing is in accordance with the provisions