b. Printing and distributing reports, studies, guidelines, and other health related technical material.

c. Developing new/redesigning existing data systems to meet specific program needs.

d. Acquisition, development, and implementation of ADP systems.

4. The awarding of funds through grants and cooperative agreements for training, research, investigations, and technical assistance. Typical actions include:

a. Occupational safety and health reserach and training.

b. Childhood immunization.

c. Veneral disease control, research, demonstration, and public information and education.

d. Health programs for refugees under the Immunization and Nationality Act.

e. State-based diabetes control or other programs.

f. Preventive health and health services block grants.

g. Investigations and technical assistance.

5. Actions associated with the conduct of liaison functions with other government and nongovermental entities. CDC is represented on a wide range of groups such as:

a. Intergovernmental task forces.

b. Ad hoc committees.

c. Work groups.

d. National code setting organizations.

e. International committees.

f. Interdepartmental groups.

6. Actions related to routine maintenance, repair, or replacement of equipment or structural components (doors, windows, roof, etc.) of CDC controlled facilities and improvements to those facilities. (Note: This exclusion does not apply to facilities listed or eligible for listing on National Register of Historic Places.)

7. Actions associated with data collection, storage, and dissemination. These actions typically involve:

a. Surveillance of health, population, and other indices and analysis for program management and budget justification purposes.

b. Identification and definition of preventable health problems including conducting research and demonstrations.

c. Surveillance of diseases through epidemiologic, laboratory, and field investigations and data collection, analysis, and distribution.

d. Planning, developing, and producing the Morbidity and Mortality Weekly Report and various other surveillance reports.

8. Technical assistance by CDC program personnel. These actions typically consist of:

a. Technical assistance to other Federal agencies, other HHS components, State and local governments, universities, nonprofit organizations, foreign governments, and international organizations.

b. The assignment of CDC personnel to Federal, State, and local governmental agencies, universities, nonprofit organizations, foreign governments, and international organizations for technical assistance.

9. Actions related to the adoption of regulations and guidelines pertaining to the above activities (except technical assistance and those resulting in population changes).

C. The following CDC program actions are excluded from environmental review requirements under provisions of HHS GAM Section 30–20–40 based on the determination that they will not normally: (a) significantly affect the human environment (as defined in NEPA), or (b) affect an asset (as defined in the related acts) regardless of location or magnitude:

1. Direct delivery of medical, laboratory, or other related health services by CDC staff or by contract providers.

2. Utilization of health professionals or paraprofessionals to supplement existing manpower resources in medically underserved areas or during health emergencies.

3. Application of pesticides which are not classified for restricted use under provisions of the Federal Insecticide, Fungicide, and Rodenticide Act when used for routine pest control purposes.

4. Relocation of employees into existing owned or office space currently leased within the same metropolitan area.

D. The following CDC program actions are partially excluded from environmental review requirements under provisions of HHS GAM Section 30–20–40 based on the determination that they may cause a significant environmental effect or impact an asset at some but not all locations and/or levels of magnitude or they may have an effect/impact associated with some but not all environmental and related acts:

1. Actions associated with the construction of 10,000 square feet or less of occupiable space are excluded except when such construction impacts properties: (a) listed or eligible for listing on the National Register of Historic Places; (b) with possible archeological, prehistoric, or scientific importance; and/or (c) located where natural asset review is mandated (see GAM Section 30–50).

2. Program actions with similar or related actions subject to previous environmental review if the effects of the action have been determined environmentally insignificant and the historic/natural asset implications are the same as for the action(s) previously reviewed.

Dated: February 23, 1983.

William H. Foege,

Director, Centers for Disease Control.

[FR Doc. 83-5648 Filed 3-3-83; 8:45]

BILLING CODE 4160-18-M

Food and Drug Administration

[Docket No. 83F-0037]

EMS-CHEMIE AG; Filing of Food Additive Petition

AGENCY: Food and Drug Administration.

ACTION: Notice.

SUMMARY: The Food and Drug Administration (FDA) is announcing that the EMS-CHEMIE AG has filed a petition proposing that the food additive regulations be amended to provide for the safe use of Nylon 12T in food-contact articles.

FOR FURTHER INFORMATION CONTACT: Julia L. Ho, Bureau of Foods (HFF-334), Food and Drug Administration, 200 C St. SW., Washington, D.C. 20204, 202-472-5690.

SUPPLEMENTARY INFORMATION: Under the Federal Food, Drug, and Cosmetic Act (sec. 409(b)(5), 72 Stat. 1786 (21 U.S.C. 348(b)(5))), notice is given that a petition (FAP 2B3670) has been filed by EMS-CHEMIE AG, CH-7013 Domat/Ems Switzerland, proposing that Part 177 (21 CFR Part 177) of the food additive regulations be amended to provide for the safe use of Nylon 12T manufactured by polymerization of *omega*-laurolactam, isophthalic acid and bis(4-amino-3-methylcyclohexyl)methane in food-contact articles.

The agency has carefully considered the potential environmental effects of this action and has concluded that the action will not have a significant impact on the human environment and that an environmental impact statement is not required. The agency's finding of no significant impact and the evidence supporting that finding may be seen in the Dockets Management Branch (HFA-305), Food and Drug Administration, Rm. 4-62, 5600 Fishers Lane, Rockville, Md 20857, between 9 a.m. and 4 p.m., Monday through Friday.

Dated: February 24, 1983.

Sanford A. Miller,

Director, Bureau of Foods.

[FR Doc. 83-5479 Filed 3-3-83; 8:45 am]

BILLING CODE 4160-01-M

[Docket No. 83F-0006]

Monsanto Co.; Filing of Food Additive Petition

AGENCY: Food and Drug Administration.

ACTION: Notice.

SUMMARY: The Food and Drug Administration (FDA) is announcing that Monsanto Co. has filed a petition proposing that the food additive regulations be amended to provide for the safe use of beverage containers fabricated from acrylonitrile/styrene copolymer resins.

FOR FURTHER INFORMATION CONTACT: Terry C. Troxell, Bureau of Foods (HFF-334), Food and Drug Administration, 200 C St. SW., Washington, DC 20204, 202-472-5690.

SUPPLEMENTARY INFORMATION: Under the Federal Food, Drug, and Cosmetic Act (sec. 409(b)(5), 72 Stat. 1786 (21 U.S.C. 348(b)(5))), notice is given that a petition (FAP 3B3690) has been filed by Keller and Heckman, 1150 17th St. NW., Washington, DC 20036, on behalf of Monsanto Co., proposing that § 177.1040 (21 CFR 177.1040) of the food additive regulations be amended to provide for the safe use of beverage containers fabricated from certain acrylonitrile/styrene copolymer resins.

The potential environmental impact of this action is being reviewed. If 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) (proposed December 11, 1979; 44 FR 71742).

Dated February 24, 1983.

Sanford A. Miller,

Director, Bureau of Foods.

[FR Doc. 83-5481 Filed 3-3-83; 8:45 am]

BILLING CODE 4160-01-M

[Docket No. 83C-0041]

Precision-Cosmet Co., Inc.; Filing of Color Additive Petition

AGENCY: Food and Drug Administration.

ACTION: Notice.

SUMMARY: The Food and Drug Administration (FDA) is announcing that Precision-Cosmet Co., Inc., has filed a petition proposing that the color additive regulations be amended to provide for the safe use of 2-[[2,5-diethoxy-4-[(4-methylphenyl)thio]phenyl]azo]-1,3,5-benzenetriol (a diazonium compound) for coloring soft (hydrophilic) contact lenses.

FOR FURTHER INFORMATION CONTACT: George C. Murray, National Center for Devices and Radiological Health (HFK-460), Food and Drug Administration, 8757 Georgia Ave., Silver Spring, MD 20910, 301-427-7940.

SUPPLEMENTARY INFORMATION: Under the Federal Food, Drug, and Cosmetic Act (sec. 706(b)(1), 74 Stat. 399-402 as amended (21 U.S.C. 376(b)(1))), notice is given that a petition, CAP 3CO159, has been filed by Precision-Cosmet Co., Inc., Minnetonka, MN 55343, proposing that the color additive regulations be amended to provide for the safe use of 2-[[2,5-diethoxy-4-[(4-methylphenyl)thio]phenyl]azo]-1,3,5-benzenetriol (a diazonium compound) for placing an identification mark on and in soft (hydrophilic) contact lenses.

The potential environmental impact of this action is being reviewed. If 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) (proposed December 11, 1979; 44 FR 71742).

Dated: February 24, 1983.

Sanford A. Miller,

Director, Bureau of Foods.

[FR Doc. 83-5483 Filed 3-3-83; 8:45 am]

BILLING CODE 4160-01-M

[Docket No. 83F-0042]

Schenectady Chemicals, Inc.; Filing of Food Additive Petition

AGENCY: Food and Drug Administration.

ACTION: Notice.

SUMMARY: The Food and Drug Administration (FDA) is announcing that Schenectady Chemicals, Inc., has filed a petition proposing that the food additive regulations be admended to provide for the safe use of 2-2'-ethylidene bis(4,6-di-*tert*-butylphenol) as an antioxidant and/or stabilizer in acrylonitrile-butadiene-styrene copolymers and high-impact polystyrene and also for use in adhesive formulations intended for food-contact applications.

FOR FURTHER INFORMATION CONTACT: Vir Anand, Bureau of Foods (HFF-334),Food and Drug Administration, 200 C St. SW., Washington, DC 20204, 202-472-5690.

SUPPLEMENTARY INFORMATION: Under the Federal Food, Drug, and Cosmetic Act (sec. 409(b)(5), 72 Stat. 1786 (21 U.S.C. 348(b)(5))), notice is given that a petition (FAP 3B3691) has been filed by Schenectady Chemicals, Inc., c/o Jerome H. Heckman, Keller and Heckman, 1150 17th St. NW., Washington, DC 20036, proposing that the food additive regulations be amended to provide for the safe use of 2-2'-ethylidene bis(4,6-di-*tert*-butylphenol) as an antioxidant and/or stabilizer in acrylonitrile-butadiene-styrene copolymers and high-impact polystyrene and also for use in adhesive formulations intended for use in contact with food.

The potential environmental impact of this action is being reviewed. If 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) (proposed December 11, 1979; 44 FR 71742).

Dated: February 24, 1983.

Sanford A. Miller,

Director, Bureau of Foods.

[FR Doc. 83-5480 Filed 3-3-83; 8:45 am]

BILLING CODE 4160-01-M

[Docket No. 83F-0036]

Standard Oil Co. (Indiana); Filing of Food Additive Petition

AGENCY: Food and Drug Administration.

ACTION: Notice.

SUMMARY: The Food and Drug Administration (FDA) is announcing that Standard Oil Co. (Indiana) has filed a petition proposing that the food additive regulations be amended to provide for the safe use of 2,5-dimethyl-2,5-di(*tert*-butylperoxy)hexane in the production of polyolefins.

FOR FURTHER INFORMATION CONTACT: Mary W. Lipien, Bureau of Foods (HFF-334), Food and Drug Administration, 200 C St. SW., Washington, D.C. 20204, 202-472-5740.

SUPPLEMENTARY INFORMATION: Under the Federal Food, Drug, and Cosmetic Act (sec. 409(b)(5), 72 Stat. 1786 (21 U.S.C. 348(b)(5))), notice is given that a petition (FAP 2B3658) has been filed by Standard Oil Co. (Indiana), 200 E.

Randolph Dr., Chicago, IL 60601, proposing that the food additive regulations be amended to provide for the safe use of 2,5-dimethyl-2,5-di(*tert*-butylperoxy)hexane in the production of polyolefins.

The potential environmental impact of this action is being reviewed. If 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) (proposed December 11, 1979; 44 FR 71742).

Dated: February 24, 1983.

Sanford A. Miller,

Director, Bureau of Foods.

[FR Doc. 83-5482 Filed 3-3-83; 8:45 am]

BILLING CODE 4160-01-M

Consumer Participation; Open Meeting

AGENCY: Food and Drug Administration.

ACTION: Notice.

SUMMARY: The Food and Drug Administration (FDA) is announcing the following consumer exchange meeting: San Juan District Office, chaired by Lynn Campbell, District Director.

DATE: Thursday, March 17, 1983, 10:30 a.m.

ADDRESS: Ponce Regional College, University of Puerto Rico, Ponce, PR 00731.

FOR FURTHER INFORMATION CONTACT: William E. Martinez-Soto, Consumer Affairs Officer, Food and Drug Administration, P.O. Box S-4427, Old San Juan Station, San Juan, PR 00905; 809-753-4264.

SUPPLEMENTARY INFORMATION: The purpose of this meeting is to encourage dialogue between consumers and FDA officials, to identify and set priorities for current and future health concerns, to enhance relationships between local consumers and FDA's District Offices, and to contribute to the agency's policymaking decisions on vital issues. The topics to be discussed are orphan drugs, food labeling formats, and an update on sodium.

Dated: February 25, 1983.

William F. Randolph,

Acting Associate Commissioner for Regulatory Affairs.

[FR Doc. 83-5332 Filed 3-3-83; 8:45 am]

BILLING CODE 4160-01-M

Consumer Participation; Open Meetings

Correction

In FR Doc. 82-1519, appearing on page 2836 in the issue of Friday, January 21, 1983, make the following correction.

On page 2836, in the **"FOR FURTHER INFORMATION CONTACT"** section, the Zip Code reading "21202" should read "21201".

BILLING CODE 1505-01-M

[Docket No 79N-0113; DESI 2847]

Certain Parenteral Multivitamin Products; Drug Efficacy Study Implementation; Withdrawal of Approval

Correction

In FR. Doc. 83-1419, beginning on page 2835 in the issue of Friday, January 21, 1983, make the following correction.

On page 2836, first column, fourth line of the last paragraph, "approval" should read "approved".

BILLING CODE 1505-01-M

National Institutes of Health

Biomedical Research Support Subcommittee of the General Research Support Review Committee; Meeting

Pursuant to Pub. L. 92-463, notice is hereby given of the meeting of the Biomedical Research Support Subcommittee of the General Research Support Review Committee, Division of Research Resources, National Institutes of Health, May 6, 1983, Building 31C, Conference Room 7, Bethesda, Maryland 20205, from 9:30 a.m. to adjournment.

The meeting will be open to the public on May 6 from 9:30 a.m. to adjournment to discuss program policies and planning for the Biomedical Research Support Grant Program and the Biomedical Research Support Shared Instrumentation Grant Program. Attendenance by the public will be limited to space available.

Mr. James Augustine, Information Officer, Division of Research Resources, Room 5B10, Building 31, National Institutes of Health, Bethesda, Maryland 20205, (301) 496-5545, will provide summaries of the meeting and rosters of the Committee members. Dr. Marjorie A. Tingle, Executive Secretary, Biomedical Research Support Subcommittee of the General Research Support Review Committee will furnish substantive program information and will receive any comments pertaining to this announcement.

(Catalogue of Federal Domestic Assistance Program No. 13.337, Biomedical Research Support, National Institutes of Health)

Dated: February 28, 1983.

Betty J. Beveridge,

Committee Management Officer, National Institutes of Health.

[FR Doc. 83-5569 Filed 3-3-83; 8:45 am]

BILLING CODE 4140-01-M

Environmental Health Sciences Review Committee; Meeting;

Pursuant to Pub. L. 92-463, notice is hereby given of the meeting of the Environmental Health Sciences Review Committee on March 31-April 1, 1983 in Building 101 Conference Room, Research Triangle Park, North Carolina. This meeting will be open to the public from 8:30 a.m. to approximately 10:30 a.m. on March 31, 1983, for general discussions. Attendance by the public is limited to space available.

In accordance with provisions set forth in sections 552b(c)(4) and 552(c)(6), Title 5, U.S. Code and section 10(d) of Pub. L. 92-463, the meeting will be closed to the public from 10:30 a.m., March 31, to adjournment on April 1, 1983, for the review, discussion and evaluation of individual grant applications and contract proposals. These applications and proposals and the discussions could reveal confidential trade secrets or commercial property such as patentable material, and personal information concerning individuals associated with the applications, the disclosure of which would constitute a clearly unwarranted invasion of personal privacy.

Dr. Carol Shreffler, Executive Secretary, Environmental Health Sciences Review Committee, National Institute of Environmental Health Sciences, National Institutes of Health, P.O. Box 12233, Research Triangle Park, North Carolina 27709, (telephone 919-541-7826), will provide summaries of meetings, rosters of committee members, and substantive program information.

(Catalog of Federal Domestic Assistance Program Nos. 13.892, Prediction, Detection and Assessment of Environmental Caused Diseases and Disorders; 13.893, Mechanisms of Environmental Diseases and Disorders; 13.894, Environmental Health Research and Manpower Development Resources, National Institutes of Health)

Dated: February 17, 1983.

Betty J. Beveridge,

Committee Management Officer, NIH.

[FR Doc. 83-5568 Filed 3-3-83; 8:45 am]

BILLING CODE 4140-01-M