publication. Under section 307(b)(2) of the Act, the requirements which are the subject of today's notice may not be challenged later in judicial proceedings, if any, brought by EPA to enforce these requirements.

This action is not a rule as defined by Executive Order 12291. Therefore, it is exempt from review by the Office of Management and Budget as required for rules and regulations by Executive Order 12291. Additionally a regulatory Impact Analysis is not being prepared under Executive Order 12291, for this waiver determination since it is not a rule.

This action also is not a "rule"as defined in the Regulatory Flexibility Act, 5 U.S.C. 601(2). Therefore EPA has not prepared a supporting regulatory flexibility analysis addressing the impact of this action on small business entities.

Dated: August 26, 1986.

Don R. Clay,

Acting Assistant Administrator for Air and Radiation.

[FR Doc. 86-19712 Filed 8-29-86; 8:45 am]

BILLING CODE 6560-50-M

[OPTS-59228; FRL-3073-2]

Lignosulfinic Acid, Triethanolamine Salt

AGENCY: Environmental Protection Agency (EPA).

ACTION: Notice.

SUMMARY: EPA may upon application exempt any person from the premanufacturing notification requirements of section 5(a) or (b) of the Toxic Substances Control Act (TSCA) to permit the person to manufacture or process a chemical for test marketing purposes under section 5(h)(1) of TSCA. Requirements for test marketing exemption (TME) applications, which must either be approved or denied within 45 days of receipt, are discussed in EPA's final rule published in the **Federal Register** of May 13, 1983 (48 FR 21722). This notice, issued under section 5(h)(6) of TSCA, announces receipt of one application for exemption, provides a summary, and requests comments on the appropriateness of granting the exemption.

DATE: Written comments by: September 17, 1986.

ADDRESS: Written comments, identified by the document control number "[OPTS-59228]" and the specific TME number should be sent to: Document Control Officer (TS-790), Confidential Data Branch, Information Management Division, Office of Toxic Substances, Environmental Protection Agency, Rm. E-201, 401 M Street, SW., Washington, DC 20460, (202) 382-3532.

FOR FURTHER INFORMATION CONTACT: Wendy Cleland-Hamnett, Premanufacture Notice Management Branch, Chemical Control Division, (TS-794), Office of Toxic Substances, Environmental Protection Agency, Rm. E-611, 401 M Street, SW., Washington, DC 20460, (202) 382-3725.

SUPPLEMENTARY INFORMATION: The following notice contains information extracted from the non-confidential version of the submission provided by the manufacturer on the TMEs received by EPA. The complete non-confidential document is available in the Public Reading Room NE-G004 at the above address between 8:00 a.m. and 4:00 p.m., Monday through Friday, excluding legal holidays.

T 86-57

Close of Review Period. October 1, 1986.

Manufacturer. Westvaco Corporation.

Chemical. (S) Lignosulfonic acid, triethanolamine salt.

Use/Production. (G) Dispersant for agricultural products, dyestuffs and colorants. Prod. range: Confidential.

Toxicity Data. No data submitted.

Exposure. Confidential.

Environmental Release/Disposal. Confidential.

Dated: August 22, 1986.

V. Paul Fuschini,

Acting Division Director, Information Management Division.

[FR Doc. 86-19714 Filed 8-29-86; 8:45 am]

BILLING CODE 6560-50-M

[SAB-FRL-3072-5]

Science Advisory Board; Radiation Advisory Committee; Drinking Water Subcommittee; Open Meeting—September 25-26, 1986

Under Pub. L. 92-463, notice is hereby given that a meeting of the Drinking Water Subcommittee of the Science Advisory Board's Radiation Advisory Committee will be held on September 25-26, 1986 at the U.S. Environmental Protection Agency, South Conference Area Room #6 on Thursday, and #12 on Friday. The Conference Area is located on the Ground Floor, near the EPA Washington Information Center, Waterside Mall, 401 M Street, SW, Washington, DC. The meeting will begin at 8:30 a.m. on Thursday and adjourn no later than 5:00 p.m. Friday. The Subcommittee will review the Office of Drinking Water's *Radionuclides in Drinking Water* and four supporting documents. Copies of the documents being reviewed may be obtained by calling or writing Dr. Joseph Cotruvo (202) 382-7575 at the Office of Drinking Water, WH-550D, U.S. Environmental Protection Agency, 401 M Street, SW., Washington, DC 20460.

The meeting is open to the public; however, seating is limited. Any member of the public wishing to attend or obtain information should contact Mrs. Kathleen Conway, Executive Secretary, or Mrs. Dorothy Clark, Staff Secretary, (A101-F) Radiation Advisory Committee, Science Advisory Board, by the close of business on September 22, 1986. The telephone number is (202) 382-2552.

Dated: August 21, 1986.

Terry F. Yosie,

Director, Science Advisory Board.

[FR Doc. 86-19713 Filed 8-29-86; 8:45 am

BILLING CODE 6560-50-M

FEDERAL COMMUNICATIONS COMMISSION

New TV Stations; Applications for Consolidated Hearing; Jacksonville Broadcasting Co. and James Capers, Jr.

1. The Commission has before it the following mutually exclusive applications for a new TV station:

Applicant and city/State	File No.	MM Docket No.
A. Sidney Popkin d/b/a Jacksonville Broadcasting Company, Jacksonville, NC.	BPCT-860304KE.	86-344
B. James Capers, Jr., Jacksonville, NC.	BPCT-860422KG.	.....

2. Pursuant to section 309(e) of the Communications Act of 1934, as amended, the above applications have been designated for hearing in a consolidated proceeding upon the issues whose headings are set forth below. The text of each of these issues has been standardized and is set forth in its entirety under the corresponding headings at 51 FR 19347, May 29, 1986. The letter shown before each applicant's name, above, is used below to signify whether the issue in question applies to that particular applicant.

Issue heading	Applicant(s)
Air hazard	B
Comparative	A, B
Ultimate	A, B

3. If there is any non-standardized issue(s) in this proceeding, the full text

of the issue and the applicant(s) to which it applies are set forth in an Appendix to this Notice. A copy of the complete HDO in this proceeding is available for inspection and copying during normal business hours in the FCC Dockets Branch (Room 230), 1919 M Street, NW., Washington, DC. The complete text may also be purchased from the Commission's duplicating contractor, International Transcription Services, Inc., 2100 M Street, NW., Washington, DC 20037 (Telephone No. (202) 857–3800).

Stephen F. Sewell,

Assistant Chief, Video Services Division, Mass Media Bureau.

[FR Doc. 86–19505 Filed 8–29–86; 8:45 am]

BILLING CODE 6712-01-M

DEPARTMENT OF HEALTH AND HUMAN SERVICES

Alcohol, Drug Abuse, and Mental Health Administration

Extramural Research Support Programs

AGENCY: National Institute of Mental Health.

ACTION: Issuance of notice of the revision of the program announcement on Extramural Research Support Programs.

SUMMARY: The National Institute of Mental Health announces the availability of a revised Extramural Research Programs announcement, MH–86–18. This announcement reflects changes in programs resulting from the recent reorganization of the Institute. It also contains the new schedule for the review of applications and an updated list of contact persons for the ongoing extramural research programs.

FOR FURTHER INFORMATION CONTACT: Anne W. Cooley, Extramural Policy Branch, Division of Extramural Activities, National Institute of Mental Health, Parklawn Building, Room 9–95, 5600 Fishers Lane, Rockville, Maryland 20857, Telephone: (301) 443–4673.

Donald Ian Macdonald,

Administrator, Alcohol, Drug Abuse, and Mental Health Administration.

[FR Doc. 86–19667 Filed 8–29–86; 8:45 am]

BILLING CODE 4160-20-M

Food and Drug Administration

[Docket No. 86F–0341]

Dow Chemical Co.; Filing of Food Additive Petition

AGENCY: Food and Drug Administration.

ACTION: Notice.

SUMMARY: The Food and Drug Administration (FDA) is announcing that The Dow Chemical Co. has filed a petition proposing that the food additive regulations be amended to provide for the safe use of ethylene-carbon monoxide copolymers as components of food-packaging materials.

FOR FURTHER INFORMATION CONTACT: Julius Smith, Center for Food Safety and Applied Nutrition (HFF–335), 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 6B3948) has been filed by The Dow Chemical Co., 1803 Building, Door 7, Midland, MI 48674, proposing that Part 177 (21 CFR Part 177) of the food additive regulations be amended to provide for the safe use of ethylene-carbon monoxide copolymers as components of food-packaging material.

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).

Dated: August 5, 1986.

Richard J. Ronk,

Acting Director, Center for Food Safety and Applied Nutrition.

[FR Doc. 86–19680 Filed 8–29–86; 8:45 am]

BILLING CODE 4160-01-M

[Docket No. 86G–0321]

Donath-Kelterei; Filing of Petition for Affirmation of GRAS Status

AGENCY: Food and Drug Administration.

ACTION: Notice.

SUMMARY: The Food and Drug Administration (FDA) is announcing that a petition (GRASP 6G0314) has been filed by Donath-Kelterei, Gutenbergstrasse 4, 8043 Unterfohring, West Germany, proposing to affirm that the puree and juice from the sea buckthorn berry *Hippophae rhamnoides* L. is generally recognized as safe (GRAS) for use as a food ingredient.

DATE: Comments by November 3, 1986.

ADDRESS: Written comments to the Dockets Management Branch (HFA–305), Food and Drug Administration, Rm. 4–62, 5600 Fishers Lane, Rockville, MD 20857.

FOR FURTHER INFORMATION CONTACT: Geraldine E. Harris, Center for Food Safety and Applied Nutrition (HFF–334), Food and Drug Administration, 200 C St. SW., Washington, DC 20204, 202–426–9463.

SUPPLEMENTARY INFORMATION: Under the Federal Food, Drug, and Cosmetic Act (sec. 409(b)(5), 72 Stat. 1786 (21 U.S.C. 348(b)(5))) and the regulations for affirmation of GRAS status in § 170.35 (21 CFR 170.35), notice is given that a petition (GRASP 6G0314) has been filed by Donath-Kelterei, Gutenbergstrasse 4, 8043 Unterföhring, West Germany, proposing to affirm that the puree and juice from the sea buckthorn berry *Hippophae rhamnoides* L. is GRAS for use as a food ingredient.

The petition has been placed on display at the Dockets Management Branch (address above).

Any petition that meets the format requirements outlined in § 170.35 is filed by the agency. There is no prefiling review of the adequacy of data to support a GRAS conclusion. Thus, the filing of a petition for GRAS affirmation should not be interpreted as a preliminary indication of suitability for GRAS affirmation.

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).

Interested persons may, on or before November 3, 1986, review the petition and/or file comments (two copies, identified with the docket number found in brackets in the heading of this document) with the Dockets Management Branch (address above). Comments should include any available information that would be helpful in determining whether the substance is, or is not, GRAS. A copy of the petition and received comments may be seen in the Dockets Management Branch between 9 a.m. and 4 p.m., Monday through Friday.

Dated: August 25, 1986.

Richard J. Ronk,

Acting Director, Center for Food Safety and Applied Nutrition.

[FR Doc. 86–19679 Filed 8–29–86; 8:45 am]

BILLING CODE 4160-01-M

[Docket No. 86G-0202]

The Hereld Organization; Filing of Petition for Affirmation of GRAS Status; Correction

AGENCY: Food and Drug Administration.

ACTION: Notice; correction.

SUMMARY: The Food and Drug Administration (FDA) is correcting the GRASP number in the notice that announced the filing of a petition for affirmation of GRAS status on behalf of the Hereld Organization.

FOR FURTHER INFORMATION CONTACT: Lola Batson, Regulations Editorial Staff (HFC-222), 5600 Fishers Lane, Food and Drug Administration, Rockville, MD 20857, 301-443-2994.

SUPPLEMENTARY INFORMATION: In FR Doc. 86-12097, appearing on page 19612 in the **Federal Register** of Friday, May 30, 1986, in the first column, third line and in the second column under "Supplementary Information," seventh line, the GRASP number is changed to read "(GRASP 5G0305)".

Dated: August 25, 1986.

Richard J. Ronk,

Acting Director, Center for Food Safety and Applied Nutrition.

[FR Doc. 86-19677 Filed 8-29-86; 8:45 am]

BILLING CODE 4160-01-M

[Docket No. 86F-0294]

EMS-CHEMIE AG; Filing of Food Additive Petition

Correction

In FR Doc. 86-17182, appearing on page 27461, in the issue of Thursday, July 31, 1986, first column, in the third line of the "SUMMARY", "EMS-CHEMIC" should read "EMS-CHEMIE", and in the seventh line, "contact" was misspelled.

BILLING CODE 1505-01-M

[Docket No. 86G-0289]

The National Fish Meal and Oil Association; Filing of Petition For Affirmation of GRAS Status

Correction

In FR Doc. 86-17183, beginning on page 27461, in the issue of Thursday, July 31, 1986, make the following correction:

On page 27461, third column, in the **"FOR FURTHER INFORMATION CONTACT"** caption, third line, "NW." should read "SW."

BILLING CODE 1505-01-M

[Docket No. 86F-0339]

Rohm & Haas Co.; Filing of Food Additive Petition

AGENCY: Food and Drug Administration.

ACTION: Notice.

SUMMARY: The Food and Drug Administration (FDA) is announcing that Rohm & Haas Co., has filed a petition proposing that the food additive regulations be amended to provide for the safe use of 5-chloro-2-methyl-4-isothiazolin-3-one and 2-methyl-4-isothiazolin-3-one mixture with magnesium nitrate as an antimicrobial agent for fillers, binders, pigment slurries, sizing solutions, and coating formulations employed in the manufacture of paper and paperboard for use in contact with food.

FOR FURTHER INFORMATION CONTACT: Julius Smith, Center for Food Safety and Applied Nutrition (HFF-335), Food and Drug Administration, 200 C Street, 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 6B3947) has been filed by Rohm & Haas Co., Independence Mall West, Philadelphia, PA 19105, proposing that § 176.170 *Components of paper and paperboard in contact with aqueous and fatty foods* (21 CFR 176.170) be amended to provide for the safe use of 5-chloro-2-methyl-4-isothiazolin-3-one and 2-methyl-4-isothiazolin-3-one mixture with magnesium nitrate as an antimicrobial agent for fillers, binders, pigment slurries, sizing solutions, and coating formulations employed in the manufacture of paper and paperboard 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).

Dated: August 25, 1986.

Richard J. Ronk,

Acting Director, Center for Food Safety and Applied Nutrition.

[FR Doc. 86-19675 Filed 8-29-86; 8:45 am]

BILLING CODE 4160-01-M

Health Care Financing Administration

[OA-1-N]

Meeting of the Task Force on Long-Term Health Care Policies

AGENCY: Health Care Financing Administration (HCFA), HHS.

ACTION: Notice of public meeting.

SUMMARY: In accordance with section 10(a)(2) of the Federal Advisory Committee Act (Pub. L. 92-463), this notice announces a meeting of the Task Force on Long-Term Health Care Policies.

DATE: The meeting will be held on September 25, 1986 from 1:00 p.m. to 4:00 p.m., and on September 26, 1986 from 9:00 a.m. to 3:00 p.m., e.s.t. The meeting will be open to the public.

ADDRESS: The meeting will be held in the Marriott Hotel, 1999 Jefferson Davis Highway, Crystal City, Arlington, Virginia.

FOR FURTHER INFORMATION CONTACT: Dennis DeWitt, 202-245-0063.

SUPPLEMENTARY INFORMATION:

Purpose

The Task Force on Long-Term Health Care Policies, created by Section 9601 of the Consolidated Omnibus Budget Reconciliation Act of 1985, will evaluate current issues relating to private long-term care insurance. To ensure the evolution of sound private long-term care policies and to help foster consumer confidence in them, the Task Force will develop guidelines that can be used by State regulators, persons involved in the insurance industry, and consumers who may wish to purchase such policies.

The term "long-term health care policy" means an insurance policy, or similar health benefits plan, that is designed for or marketed as providing (or making payment for) health care services (such as nursing home care and home health care) or related services (which may include home and community-base services), or both, over an extended period of time.

The Task Force on Long-Term Health Care Policies will advise the Secretary of Health and Human Services and the Administrator of the Health Care Financing Administration concerning the development of insurance policies for long-term care that are privately marketed to individuals or groups. The Task Force will develop recommendations for long-term health care policies, including recommendations designed to: (1) Limit marketing and agent abuse for those