research program, mining industry health and safety statistics, and improving miner's health and safety.

Agenda items are subject to change as priorities dictate.

Contact Person for More Information: Lewis V. Wade, Ph.D., Executive Secretary, MSHRAC, NIOSH, CDC, 200 Independence Avenue, SW., Room 715–H, Hubert Humphrey Building, P12 Washington, DC 20201–0004, telephone 202/401–2192, fax 202/260–4464.

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: June 26, 2003.

John Burckhardt,

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

[FR Doc. 03–16677 Filed 7–1–03; 8:45 am]

BILLING CODE 4163–19–U

DEPARTMENT OF HEALTH AND HUMAN SERVICES

Food and Drug Administration

[Docket No. 2002N–0354]

Agency Information Collection Activities; Announcement of OMB Approval; The Evaluation of Long-Term Antibiotic Drug Therapy for Persons Involved in Anthrax Remediation Activities

AGENCY: Food and Drug Administration, HHS.

ACTION: Notice.

SUMMARY: The Food and Drug Administration (FDA) is announcing that a collection of information entitled ''The Evaluation of Long-Term Antibiotic Drug Therapy for Persons Involved in Anthrax Remediation Activities'' has been approved by the Office of Management and Budget (OMB) under the Paperwork Reduction Act of 1995 (the PRA).

FOR FURTHER INFORMATION CONTACT: Karen L. Nelson, Office of Management Programs (HFA–250), Food and Drug Administration, 5600 Fishers Lane, Rockville, MD 20857, 301–827–1482.

SUPPLEMENTARY INFORMATION: In the **Federal Register** of April 2, 2003 (68 FR 16059), the agency announced that the proposed information collection had been submitted to OMB for review and clearance under 44 U.S.C. 3507. An agency may not conduct or sponsor, and a person is not required to respond to, a collection of information unless it displays a currently valid OMB control number. OMB has now approved the information collection and has assigned OMB control number 0910–0494. The approval expires on April 30, 2004. A copy of the supporting statement for this information collection is available on the Internet at *http://www.fda.gov/ohrms/dockets*.

Dated: June 25, 2003.

Jeffrey Shuren,

Assistant Commissioner for Policy.

[FR Doc. 03–16618 Filed 7–1–03; 8:45 am]

BILLING CODE 4160–01–S

DEPARTMENT OF HEALTH AND HUMAN SERVICES

Food and Drug Administration

[Docket No. 2002N–0496]

Agency Information Collection Activities; Announcement of OMB Approval; Aluminum in Large and Small Volume Parenterals Used in Total Parenteral Nutrition

AGENCY: Food and Drug Administration, HHS.

ACTION: Notice.

SUMMARY: The Food and Drug Administration (FDA) is announcing that a collection of information entitled ''Aluminum In Large and Small Volume Parenterals Used in Total Parenteral Nutrition'' has been approved by the Office of Management and Budget (OMB) under the Paperwork Reduction Act of 1995 (the PRA).

FOR FURTHER INFORMATION CONTACT: Karen L. Nelson, Office of Management Programs (HFA–250), Food and Drug Administration, 5600 Fishers Lane, Rockville, MD 20857, 301–827–1482.

SUPPLEMENTARY INFORMATION: In the **Federal Register** of March 17, 2003 (68 FR 12701), the agency announced that the proposed information collection had been submitted to OMB for review and clearance under 44 U.S.C. 3507. An agency may not conduct or sponsor, and a person is not required to respond to, a collection of information unless it displays a currently valid OMB control number. OMB has now approved the information collection and has assigned OMB control number 0910–0439. The approval expires on June 30, 2006. A copy of the supporting statement for this information collection is available on the Internet at *http://www.fda.gov/ohrms/dockets*.

Dated: June 25, 2003.

Jeffrey Shuren,

Assistant Commissioner for Policy.

[FR Doc. 03–16619 Filed 7–1–03; 8:45 am]

BILLING CODE 4160–01–S

DEPARTMENT OF HEALTH AND HUMAN SERVICES

Food and Drug Administration

[Docket Nos. 97F–0284, 88F–0182, 98F–0706, 98F–0391, 97F–0170, 92F–0315, 99F–4694, 88F–0340, 95F–0021, 99F–0720, 94F–0290, and 00F–1366]

Withdrawal of Food Additive Petitions Subsequently Converted to Food Contact Notifications

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 12 food additive petitions (FAPs) proposing that the food additive regulations be amended to provide for the safe use of certain new food additives. The petitioners subsequently requested that their petitions be converted to food contact notifications for review under the agency's new food contact notification (FCN) program for food contact substances. The requested uses are now the subjects of effective notifications.

FOR FURTHER INFORMATION CONTACT: Sylvia Dodson, Center for Food Safety and Applied Nutrition (HFS–275), Food and Drug Administration, 5100 Paint Branch Pkwy., College Park, MD 20740, 202–418–3087.

SUPPLEMENTARY INFORMATION: In notices published in the **Federal Register**, on the dates indicated in table 1 of this document, FDA announced the filing of 12 FAPs. These petitions proposed to amend the food additive regulations in the sections listed in the table to provide for the safe use of the listed substances intended for use in food contact articles. Since publication of these filing notices, the petitioners have requested that their respective petitions be converted to FCNs for review under the agency's new FCN process for food contact substances and that their petitions be withdrawn when the corresponding notifications become effective. These petitions were converted to notifications and subsequently reviewed under the FCN process. The requested uses are now the subjects of effective notifications. The corresponding FAPs are now withdrawn

without prejudice to a future filing (21 CFR 171.7).

TABLE 1.—FOOD ADDITIVE PETITIONS SUBSEQUENTLY CONVERTED TO FOOD CONTACT NOTIFICATIONS

FAP No.[1] and Docket No.	FCN No.[2]	FR Citation and Date	Company	21 CFR Section/ Part	Additive	Use
7B4547, 97F–0284	87	62 FR 37266, July 11, 1997.	Eastman Chemical Co.	175.300	1,4-cyclohexanedimethanol as a polyhydric alcohol.	In polyester resins intended for coatings in contact with food.
8B4083, 88F–0182	106	53 FR 23455, June 22, 1988.	Dow Chemical Co.	176.170	Styrene-butadiene-acrylonitrile copolymers copolymerized with not more than 10 percent of one or more of the monomers of acrylic acid, fumaric acid, 2-hydroxyethyl acrylate, itaconic acid and menacrylic acid.	As components of paper and paperboard in contact with food.
8B4620, 98F–0706	115	63 FR 45820, Aug. 27, 1998.	BASF Corp.	178.3297	2,9-bis(3,5-dimethylphenyl)anthra(2,1,9-def:6,5,10-d′e′f′)diisoquinoline-1,3,8,10(2H, 9H)-tetrone. (C.I. Pigment Red 149).	As a colorant for all polymers intended for use in contact with food.
8B4595, 98F–0391	118	63 FR 32672, June 15, 1998.	BASF Corp.	178.3297	2,9-bis[4-(phenylazo)phenyl]anthra[2,1,9-def:6,5,10-d′e′f′]diisoquinoline-1,3,8,10(2H, 9H)-tetrone. (C.I. Pigment Red 178).	As a colorant for all polymers intended for use in contact with food.
7B4538, 97F–0170	123	62 FR 23467, Apr. 30, 1997.	Toyo-Morton, Ltd. c/o Keller and Heckman, LLP.	177.1390	Polyester-epoxy-urethane adhesive.	As a nonfood contact layer of laminated articles intended for use in contact with food.
2B4337, 92F–0315	124	57 FR 43740, Sept. 22, 1992.	Fina Oil and Chemical Co.	177.1640 178.2010	Rubber-modified polystyrene resin containing not less than 71 weight percent of polymer units derived from styrene monomer and octadecyl 3,5-di-*tert*-butyl-4-hydroxyhydrocinnamate.	In contact with food. As a stabilizer in the rubber-modified polystyrene.
0B4699, 99F–4694	131	64 FR 61132, Nov. 9, 1999.	Rohm and Haas Co.	175.105 and 176.170.	2-methyl-4- isothiazolin-3-one.	As an antimicrobial additive for adhesives, paper additives, and paper coatings that are intended to contact food.
8B4105, 88F–0340	146	53 FR 43272, Oct. 26, 1988.	Shell Oil Co.	177.1570	Poly-1-butene resins and butene/ethylene copolymers containing no more than 6-weight-percent ethylene.	As articles or components of articles intended for food-contact use.

TABLE 1.—FOOD ADDITIVE PETITIONS SUBSEQUENTLY CONVERTED TO FOOD CONTACT NOTIFICATIONS—Continued

FAP No.[1] and Docket No.	FCN No.[2]	FR Citation and Date	Company	21 CFR Section/ Part	Additive	Use
5B4444, 95F–0021	151	60 FR 7974, Feb. 10, 1995.	M & G Ricerche S.p.A.	177.1630	Ethylene terephthalate-isophthalate copolymers prepared with pyromellitic dianhydride such that the finished copolymers contain at least 95 weight percent of polymer units derived from ethylene terephthalate.	In contact with food.
9B4653, 99F–0720	166	64 FR 16742, Apr. 6, 1999.	Arakawa Chemical Industries, Ltd. c/o Keller and Heckman, LLP.	178	Hydrogenated aromatic petroleum hydrocarbon resins.	In blends with polymers intended for contact with food.
4B4427, 94F–0290	179	59 FR 43847, Aug. 25, 1994.	Eastman Chemical Co.	177.1315	Ethylene-1,4-cyclohexylene dimethylene terephthalate copolymers that include 1 to 100 mole percent of repeat units derived from 1,4-cyclohexylene dimethylene terephthalate.	As components of articles for food contact use.
0B4713, 00F–1366	220	65 FR 41079, July 3, 2000.	Nippon Shokubai c/o Keller and Heckman, LLP.	177.1520	Methylmethacrylate-trimethylolpropane trimethacrylate copolymer as an antiblocking agent in linear low-density polyethylene.	Intended for use in contact with food.

[1] Food Additive Petition Number.
[2] Food Contact Notification Number.

Dated: June 17, 2003.

Laura M. Tarantino

Deputy Director, Office of Food Additive Safety, Center for Food Safety and Applied Nutrition.

[FR Doc. 03–16616 Filed 7–1–03; 8:45 am]

BILLING CODE 4160–01–S

DEPARTMENT OF HEALTH AND HUMAN SERVICES

Food and Drug Administration

[Docket No. 2003N–0201]

Minimizing Medication Errors—Methods for Evaluating Proprietary Names for Their Confusion Potential; Public Meeting; Correction

AGENCY: Food and Drug Administration; HHS.

ACTION: Notice of public meeting; correction.

SUMMARY: The Food and Drug Administration (FDA) is correcting a notice that appeared in the **Federal Register** of May 30, 2003 (68 FR 32529). The document announced a public meeting to explore current methods being used to evaluate proprietary drug names to reduce medication errors due to similarity in drug names. The document was published with the incorrect docket number. This document corrects that error.

FOR FURTHER INFORMATION CONTACT: Joyce A. Strong, Chief, Regulations Editorial Section (HF–27), Food and Drug Administration, 5600 Fishers Lane, Rockville, MD 20857, 301–827–7010.

SUPPLEMENTARY INFORMATION: In FR Doc. 03–13591, appearing on page 35679 in the **Federal Register** of May 30, 2003, the following correction is made:

1. On page 32529, in the third column, the Docket No. ''02N–0201'' should be corrected to read ''2003N–0201''.

Dated: June 25, 2003.

Jeffrey Shuren,

Assistant Commissioner for Policy.

[FR Doc. 03–16617 Filed 7–1–03; 8:45 am]

BILLING CODE 4160–01–S

DEPARTMENT OF HEALTH AND HUMAN SERVICES

Health Resources and Services Administration

Small Rural Hospital Improvement Grant Program

AGENCY: Health Resources and Services Administration (HRSA), HHS.

ACTION: Notice of availability of funds.

SUMMARY: The Health Resources and Services Administration (HRSA) announces that applications are being accepted for grants to help small rural hospitals do any or all of the following: (1) Pay for costs related to the implementation of prospective payment systems (PPS), (2) comply with