

South LaSalle Street, Chicago, Illinois 60690:

1. *Michigan National Corporation*, Farmington Hills, Michigan; to acquire Independence One Asset Management Corporation, Farmington Hills, Michigan, and thereby engage in providing asset management, servicing, and collection activities for unaffiliated third parties. *NCNB Corporation*, 77 Federal Reserve Bulletin 124 (1991).

Board of Governors of the Federal Reserve System, August 19, 1991.

Jennifer J. Johnson,
Associate Secretary of the Board.

[FR Doc. 91-20208 Filed 8-22-91; 8:45 am]

BILLING CODE 6210-01-F

Gad Zeevi; Change in Bank Control Notice; Acquisition of Shares of Banks or Bank Holding Companies

The notificant listed below has applied under the Change in Bank Control Act (12 U.S.C. 1817(j)) and § 225.41 of the Board's Regulation Y (12 CFR 225.41) to acquire a bank or bank holding company. The factors that are considered in acting on notices are set forth in paragraph 7 of the Act (12 U.S.C. 1817(j)(7)).

The notice is available for immediate inspection at the Federal Reserve Bank indicated. Once the notice has been accepted for processing, it will also be available for inspection at the offices of the Board of Governors. Interested persons may express their views in writing to the Reserve Bank indicated for the notice or to the offices of the Board of Governors. Comments must be received not later than September 12, 1991.

A. Federal Reserve Bank of Dallas (W. Arthur Tribble, Vice President) 400 South Akard Street, Dallas, Texas 75222:

1. *Gad Zeevi*, London, England; to acquire 94.9 percent of the voting shares of Ameritex Bancshares Corporation, Fort Worth, Texas, and thereby indirectly acquire Riverbend Bank, N.A., Fort Worth, Texas; American Bank, Grapevine, Texas; and American Bank of Haltom City, Haltom City, Texas.

Board of Governors of the Federal Reserve System, August 19, 1991.

Jennifer J. Johnson,
Associate Secretary of the Board.

[FR Doc. 91-20209 Filed 8-22-91; 8:45 am]

BILLING CODE 6210-01-F

FEDERAL TRADE COMMISSION

Granting of Request for Early Termination of the Waiting Period Under the Premerger Notification Rules

Section 7A of the Clayton Act, 15 U.S.C. 18a, as added by Title II of the Hart-Scott-Rodino Antitrust Improvements Act of 1976, requires persons contemplating certain mergers or acquisitions to give the Federal Trade Commission and the Assistant Attorney General advance notice and to wait designated periods before consummation of such plans. Section 7A(b)(2) of the Act permits the agencies, in individual cases, to terminate this waiting period prior to its expiration and requires that notice of this action be published in the *Federal Register*.

The following transactions were granted early termination of the waiting period provided by law and the premerger notification rules. The grants were made by the Federal Trade Commission and the Assistant Attorney General for the Antitrust Division of the Department of Justice. Neither agency intends to take any action with respect to these acquisitions during the applicable waiting period.

TRANSACTIONS GRANTED EARLY TERMINATION BETWEEN: 080691 AND 081691

Name of acquiring person, name of acquired person, name of acquired entity	PMN No.	Date terminated
CPI Corp., Carl D. Newton, III, FPI Holding Corporation.....	91-1212	08/06/91
Chevron Corporation, Great Western Resources Inc., Great Western Offshore Inc.	91-1217	08/06/91
The Kassar Family Trust, B.T. Trustees (Jersey) Ltd., LIVE Entertainment, Inc.....	91-1240	08/06/91
Mosvold Shipping AS, Grosvenor Holding Limited, Hogarth Shipping Corporation	91-1251	08/06/91
Damon Group Inc., Ballantrae Partners, L.P., Damon Corporation	91-1223	08/07/91
Saratoga Partners II, L.P., National Record Mart, Inc., National Record Mart, Inc.....	91-1255	08/07/91
American Southwest Mortgage Investments Corp., Residential Mortgage Investments, Inc., RMA Financial Corporation/Residen. Mortg. Accept., Inc.....	91-1219	08/08/91
S.I. Newhouse, Jr., R.E. Turner, Turner Broadcasting System, Inc.....	91-1237	08/08/91
Mitsubishi Motors Corporation, Chrysler Corporation, Diamond-Star Motors Corporation	91-1192	08/09/91
Telephone and Data Systems, Inc. Voting Trust, Tri-State Cellular Partnership, Tri-State Cellular Partnership	91-1252	08/09/91
Donald E. Newhouse, R.E. Turner, Turner Broadcasting System, Inc.....	91-1260	08/09/91
Amoco Corporation, General Electric Company, General Electric Capital Corporation	91-1268	08/09/91
Information Partners Capital Fund, L.P., Richard C. Hawk, Hemar Corporation	91-1284	08/09/91
JWP Inc., Brandt Engineering Co., Inc., Brandt Engineering Co., Inc.....	91-1198	08/13/91
Aetna Life and Casualty Company, Bay Pacific Health Corporation, Bay Pacific Health Corporation	91-1208	08/13/91
Huntington Bancshares Inc., The Cumberland Federal Bancorporation, Inc., The Cumberland Federal Savings Bank	91-1229	08/13/91
Corning Incorporated, Jack L. Custer, Wadsworth/ALERT Laboratories, Inc.	91-1247	08/13/91
George Soros, Pansophic Systems, Incorporated, Pansophic Systems, Incorporated	91-1258	08/13/91
Anuhco, Inc., Estate of Paul E. Crouse, Deceased, CC Investment Corporation	91-1287	08/13/91
Glenn R. Jones, FKC Partners, Empire Cable Television, Inc.....	91-1181	08/14/91
Glenn R. Jones, Shapell Industries, Inc., Empire Cable Television, Inc.....	91-1199	08/14/91
Medco Containment Services, Inc., Rix Dunnington Inc., Rix Dunnington Inc	91-1241	08/14/91
Rix Dunnington Inc., Medco Containment Services, Inc., Synetic, Inc	91-1243	08/14/91
Kobe Portopia Hotel Co., Ltd., EIE-international corporation, EIE Saipan Corporation	91-1258	08/14/91
Perstorp AB, Arnold E. Ditrí, Lafayette Automotive Industrial Corp.	91-1274	08/14/91
JWP Inc., Telenova, Inc., Telenova, Inc.....	91-1288	08/14/91
CBS Inc., Midwest Communications, Inc., Midwest Communications, Inc.	91-1289	08/14/91
Continental Materials Corporation, Whitman Corporation, Krack Corporation	91-0989	08/15/91
General Electric Company, The Chase Manhattan Corporation, Chase Manhattan Leasing Company—Technology Equipment.....	91-1270	08/15/91
Western Mining Corporation Holdings Limited, Australian Consolidated Minerals Ltd., Australian Consolidated Minerals Ltd.	91-1282	08/16/91
Trustees of the Estate of Bernice Pauahi Bishop, Highness Kosan Co., Ltd., WM Associates.....	91-1293	08/16/91
Heidelberger Zement AG, J. Thomas Holton, Sherman International Corp.....	91-1295	08/16/91
Capital Holding Corporation, Durham Corporation, Durham Corporation.....	91-1301	08/16/91
Sumitomo Metal Industries, Ltd., Read-Rite Corporation, Read-Rite Corporation	91-1302	08/16/91

FOR FURTHER INFORMATION CONTACT:

Sandra M. Peay, or Renee A. Horton,
Contact Representative, Federal
Trade Commission, Premerger
Notification Office, Bureau of
Competition, room 303, Washington,
DC 20580, (202) 326-3100.

By Direction of the Commission,

Donald S. Clark,

Secretary.

[FR Doc. 91-20251 Filed 8-22-91; 8:45 am]

BILLING CODE 6750-01-M

[Docket No. C-3339]

The Perrier Group of America, Inc., et al.; Prohibited Trade Practices, and Affirmative Corrective Actions

AGENCY: Federal Trade Commission.

ACTION: Consent order.

SUMMARY: In settlement of alleged violations of federal law prohibiting unfair acts and practices and unfair methods of competition, this consent order prohibits, among other things, a Connecticut-based company and its subsidiary from making false claims that any mineral water it sells is unprocessed or unfiltered, or regarding the manner by which the water is carbonated.

DATES: Complaint and Order issued August 5, 1991.¹

FOR FURTHER INFORMATION CONTACT: Robert Cheek, FTC/S-4002, Washington, DC 20580. (202) 326-3045.

SUPPLEMENTARY INFORMATION: On Tuesday, March 26, 1991, there was published in the *Federal Register*, 56 FR 12541, a proposed consent agreement with analysis in the Matter of The Perrier Group of America, Inc., et al., for the purpose of soliciting public comment. Interested parties were given sixty (60) days in which to submit comments, suggestions or objections regarding the proposed form of the order.

Comments were filed and considered by the Commission. The Commission has ordered the issuance of the complaint in the form contemplated by the agreement, made its jurisdictional findings and entered an order to cease and desist, as set forth in the proposed consent agreement, in disposition of this proceeding.

¹ Copies of the Complaint and the Decision and Order are available from the Commission's Public Reference Branch, H-130, 6th Street & Pennsylvania Avenue, NW., Washington, DC 20580.

(Sec. 6, 38 Stat. 721; 15 U.S.C. 46. Interprets or applies sec. 5, 38 Stat. 719, as amended; 15 U.S.C. 45, 52)

Benjamin I. Berman,

Acting Secretary.

[FR Doc. 91-20250 Filed 8-22-91; 8:45 am]

BILLING CODE 9750-01-M

DEPARTMENT OF HEALTH AND HUMAN SERVICES

Food and Drug Administration

[Docket No. 91F-0289]

Rohm and Haas Co.; Filing of Food Additive Petition

AGENCY: Food and Drug Administration, HHS.

ACTION: Notice.

SUMMARY: The Food and Drug Administration (FDA) is announcing that the Rohm and Haas Co. has filed a petition proposing that the food additive regulations be amended to provide for the safe use of methyl methacrylate/butyl acrylate-grafted polypropylene as a component of propylene homopolymer and copolymer food-contact materials.

FOR FURTHER INFORMATION CONTACT: Hortense S. Macon, 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) (21 U.S.C. 348(b)(5))), notice is given that a petition (FAP 1B4272) has been filed on behalf of the Rohm and Haas Co., c/o 1150 17th St. NW., Washington, DC 20036, proposing that the food additive regulations in § 177.1520 *Olefin polymers* (21 CFR 177.1520) be amended to provide for the safe use of methyl methacrylate/butyl acrylate-grafted polypropylene for use as a component of propylene homopolymer and copolymer food-contact materials.

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 15, 1991.

Fred R. Shank,

Director, Center for Food Safety and Applied Nutrition.

[FR Doc. 91-20218 Filed 8-22-91; 8:45 am]

BILLING CODE 4160-01-M

[Docket No. 90P-0193]

Cottage Cheese Deviating From Identity Standard; Amendment of Temporary Permit of Market Testing

AGENCY: Food and Drug Administration, HHS.

ACTION: Notice.

SUMMARY: The Food and Drug Administration (FDA) is announcing that it is amending a temporary permit issued to The Kroger Co. to market test a product designated as "nonfat cottage cheese" that deviates from the U.S. standards of identity for cottage cheese (21 CFR 133.128), dry curd cottage cheese (21 CFR 133.129), and lowfat cottage cheese (21 CFR 133.131) to include 340-gram (g) (12-ounce (oz)) and 454-g (16-oz) container sizes not specified in the original temporary permit for market testing. In addition, the milkfat content allowed in the nonfat cottage cheese test product is changed from "0.1 percent" to "less than 0.4 percent."

FOR FURTHER INFORMATION CONTACT: Michelle A. Smith, Center for Food Safety and Applied Nutrition (HFF-414), Food and Drug Administration, 200 C Street SW., Washington, DC 20204, 202-485-0106.

SUPPLEMENTARY INFORMATION: In the *Federal Register* of August 9, 1990 (55 FR 32473), FDA issued a temporary permit under the provisions of 21 CFR 130.17 to The Kroger Co., 1014 Vine Street, Cincinnati, Ohio 45202, to market test in interstate commerce "nonfat cottage cheese." The agency issued the permit to facilitate interstate market testing of a nonfat cottage cheese, formulated from dry curd cottage cheese and a dressing, such that the finished product contained 0.1 percent milkfat. The food deviates from the U.S. standards of identity for cottage cheese (21 CFR 133.128) and lowfat cottage cheese (21 CFR 133.131) in that the test product contains 0.1 percent milkfat compared to not less than 4.0 percent milkfat in cottage cheese and 0.5 to 2.0 percent milkfat in lowfat cottage cheese. The test product also deviates from the U.S. standard of identity for dry curd cottage cheese (21 CFR 133.129) because of the added dressing. The test product meets all requirements of the standards with the

exception of these deviations. The purpose of the variation is to offer the consumer a product that is nutritionally equivalent to cottage cheese but contains less fat.

The Kroger Co. has requested that FDA amend its temporary permit to include container sizes not previously specified. The company states that these changes in package size are necessary, based on preliminary acceptance and consumer requests that the product be offered in container sizes beyond the current 680-g (24-oz) container size.

The Kroger Co. has also requested that FDA change the level of milkfat allowed in the test product from "0.1 percent" to "less than 0.4 percent." The Company maintains that this amendment will not alter the substance of the temporary permit (55 FR 32473), but will reflect a more realistic statement of the nutritional values of the product based on experience gained from mass production. Milkfat content remains less than 0.5 g per serving.

Therefore, under the provision of 21 CFR 130.17(f), FDA is amending the temporary permit to include 340-g (12-oz) and 454-g (16-oz) container sizes not previously specified. In addition, FDA is changing the level of milkfat allowed in the test product from "0.1 percent" to "less than 0.4 percent." All other terms and conditions of this permit remain unchanged.

Dated: August 15, 1991.

Fred R. Shank,

Director, Center for Food Safety and Applied Nutrition.

[FR Doc. 91-20291 Filed 8-22-91; 8:45 am]

BILLING CODE 4160-01-M

[Docket No. 91P-0246]

Cottage Cheese Deviating From Identity Standard; Temporary Permit for Market Testing

AGENCY: Food and Drug Administration, HHS.

ACTION: Notice.

SUMMARY: The Food and Drug Administration (FDA) is announcing that a temporary permit has been issued to Marigold Foods, Inc., to market test a product designated as "nonfat cottage cheese" that deviates from the U.S. standards of identity for cottage cheese (21 CFR 133.128), dry curd cottage cheese (21 CFR 133.129), and lowfat cottage cheese (21 CFR 133.131). The purpose of the temporary permit is to allow the applicant to measure consumer acceptance of the product, identify mass production problems, and assess commercial feasibility.

DATES: This permit is effective for 15 months, beginning on the date the food is introduced or caused to be introduced into interstate commerce, but not later than November 21, 1991.

FOR FURTHER INFORMATION CONTACT: Shellee A. Davis, Center for Food Safety and Applied Nutrition (HFF-414), Food and Drug Administration, 200 C Street SW., Washington, DC 20204, 202-485-0112.

SUPPLEMENTARY INFORMATION: In accordance with 21 CFR 130.17 concerning temporary permits to facilitate market testing of foods deviating from the requirements of the standards of identity promulgated under section 401 of the Federal Food, Drug, and Cosmetic Act (21 U.S.C. 341), FDA is giving notice that a temporary permit has been issued to Marigold Foods, Inc., 2929 University Ave. SE., Minneapolis, MN 55414.

The permit covers limited interstate marketing tests of a nonfat cottage cheese, formulated from dry curd cottage cheese and a dressing, such that the finished product contains less than 0.5 gram of milkfat per serving. The food deviates from the U.S. standards of identity for cottage cheese (21 CFR 133.128) and lowfat cottage cheese (21 CFR 133.131) in that the test product contains 0.5 percent milkfat compared to not less than 4.0 percent milkfat in cottage cheese, and 0.5 to 2.0 percent milkfat in lowfat cottage cheese. The test product also deviates from the U.S. standard of identity for dry curd cottage cheese (21 CFR 133.129) because of the added dressing. The test product meets all requirements of the standards with the exception of these deviations. The purpose of the variation is to offer the consumer a product that is nutritionally equivalent to cottage cheese products with dressing but contains less fat.

For the purpose of this permit, the name of the product is "nonfat cottage cheese." The information panel of the label will bear nutrition labeling in accordance with 21 CFR 101.9.

This permit provides for the temporary marketing of 680,400 kilograms (1,500,000 pounds) of the test product. The product will be manufactured at Marigold Foods, Inc., 15 Fourth St., Farmington, MN 55024, and distributed in Illinois, Indiana, Iowa, Michigan, Minnesota, and Wisconsin.

Each of the ingredients used in the food must be declared on the label as required by the applicable sections of 21 CFR Part 101. This permit is effect for 15 months, beginning on the date the food is introduced or caused to be introduced into interstate commerce, but not later than November 21, 1991.

Dated: August 15, 1991.

Fred R. Shank,

Director, Center for Food Safety and Applied Nutrition.

[FR Doc. 91-20292 Filed 8-22-91; 8:45 am]

BILLING CODE 4160-01-M

[Docket No. 91P-0270]

Cottage Cheese Deviating From Identity Standard; Temporary Permit for Market Testing

AGENCY: Food and Drug Administration, HHS.

ACTION: Notice.

SUMMARY: The Food and Drug Administration (FDA) is announcing that a temporary permit has been issued to Oak Farms Dairy to market test a product designated as "nonfat cottage cheese" that deviates from the U.S. standards of identity for cottage cheese (21 CFR 133.128), dry curd cottage cheese (21 CFR 133.129), and lowfat cottage cheese (21 CFR 133.131). The purpose of the temporary permit is to allow the applicant to measure consumer acceptance of the product, identify mass production problems, and assess commercial feasibility.

DATES: This permit is effective for 15 months, beginning on the date the food is introduced or caused to be introduced into interstate commerce, but not later than November 21, 1991.

FOR FURTHER INFORMATION CONTACT: Michelle A. Smith, Center for Food Safety and Applied Nutrition (HFF-414), Food and Drug Administration, 200 C Street SW, Washington DC 20204, 202-485-0106.

SUPPLEMENTARY INFORMATION: In accordance with 21 CFR 130.17 concerning temporary permits to facilitate market testing of foods deviating from the requirements of the standards of identity promulgated under section 401 of the Federal Food, Drug, and Cosmetic Act (21 U.S.C. 341), FDA is giving notice that a temporary permit has been issued to Oak Farms Dairy, 1114 North Lancaster Avenue, Dallas, TX 75203.

The permit covers limited interstate marketing tests of a nonfat cottage cheese, formulated from dry curd cottage cheese and a dressing, such that the finished product contains less than 0.5 percent milkfat and less than 0.5 gram fat per serving. The food deviates from the U.S. standards of identity for cottage cheese (21 CFR 133.128) and lowfat cottage cheese (21 CFR 133.131) in that the milkfat content of cottage cheese is not less than 4.0 percent, and