summarizing the evidence that would be presented at a hearing.

Comments regarding this application must be received not later than January 23, 1989.

A. **Federal Reserve Bank of Atlanta** (Robert E. Heck, Vice President) 104 Marietta Street, NW., Atlanta, Georgia 30303:

1. *Parish National Corporation*, Bogalusa, Louisiana; to acquire 100 percent of the voting shares of Parish National Bank of St. Tammany Parish, Covington, Louisiana.

Board of Governors of the Federal Reserve System, December 30, 1988.

William W. Wiles,

Secretary of the Board.

[FR Doc. 89-178 Filed 1-5-89; 8:45 am]

BILLING CODE 6210-01-M

Change in Bank Control; Acquisition of Shares of Banks or Bank Holding Companies; Security Bank Holding Co.

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 notices are available for immediate inspection at the Federal Reserve Bank indicated. Once the notices have been accepted for processing, they 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 that notice or to the offices of the Board of Governors. Comments must be received not later than January 27, 1989.

A. **Federal Reserve Bank of San Francisco** (Harry W. Green, Vice President) 101 Market Street, San Francisco, California 94105:

1. *Security Bank Holding Company Employee Stock Ownership Trust*, Coos Bay, Oregon; to acquire up to 24.9 percent of the voting shares of Security Bank Holding Company, Coos Bay, Oregon, and thereby indirectly acquire Security Bank, Coos Bay, Oregon.

Board of Governors of the Federal Reserve System, December 30, 1988.

William W. Wiles,

Secretary of the Board.

[FR Doc. 89-179 Filed 1-5-89; 8:45 am]

BILLING CODE 6210-01-M

DEPARTMENT OF HEALTH AND HUMAN SERVICES

Office of the Secretary

Agency Forms Submitted to the Office of Management and Budget for Clearance

Each Friday the Department of Health and Human Services (HHS) publishes a list of information collection packages it has submitted to the Office of Management and Budget (OMB) for clearance in compliance with the Paperwork Reduction Act (44 U.S.C. Chapter 35). The following are those packages submitted to OMB since the last list was published on December 30, 1988.

Public Health Service

(Call Reports Clearance Office on 202-245-2100 for copies of package)

1. Cardiovascular Health Study (CHS)—NEW—A random sample of 5,336 men and women aged 65 and older will be selected from four communities. They will provide medical, social and demographic information and will participate in two clinical examinations to study risk factors for the onset and progression of clinical coronary heart disease and stroke, and related preclinical conditions. Individuals or households, Businesses or other for-profit, Small businesses or organizations.

	1st information collection	2d information collection
	Title—Cardiovascular Health	Physician Questionnaire
Number of Resondents	3,018	184
Number of Responses: Per Respondent	5.1	1
Average Burden: Per Response	.98	10

Total Burden Hours: 15,388

2. Common or Usual Name: Labeling of Peanut Spreads (21 CFR Part 102)—0910-0222—This regulation sets forth the nutritional requirements to be met if a non-standardized peanut spread is not to be labeled as "imitation" peanut butter. Respondents: Businesses or other for-profit: Number of Respondents: 40; Number of Responses per Respondent: 5; Average burden per Response: 107.4; Estimated Annual Burden: 2,148. This Burden is included in Nutritional Labeling, OMB NO. 0910-0177.

3. Application to Participate in the Public Health Capitation Program—0915-0089—The information collected from schools of public health is needed in order for the Public Health Service to determine the amount of the grants based on the number students reported and to audit grantee expenditures.

Respondents: Non-profit Institutions.

	1st information collection	2d information collection
	Title—Student Counts and Assurance on Non-Federal Funding—57.3504	Audits—57.3504(a)
Number of Resondents	26	26
Number of Responses: Per Respondent	1	1
Average Burden: Per Response	6 hours	1 hour

Total Burden Hours: 182

4. Identity Labeling of Food in Package Form—21 CFR 101.3—0910-0223—These regulations define what constitutes an imitation food product. Firms that manufacture substitute foods use the information to determine whether the term "imitation" is required on product labels. In addition these regulations specify when "imitation" cannot be used on the label.

Respondents: Businesses or other for-profit.

	1st Information Collection	2nd Information Collection
	Title: 21 CFR 101.3(e)(1)—Disclosure—Labeling (Approval of Language)*	21 CFR 101.3(e)(2) Disclosure Labeling (Approval of Language)*
Number of Respondents	0	0
Number of Responses Per Respondent	0	0
Average Burden Per Response	0	0

Total Burden hours: 1

* This burden is included in Nutritional Labeling, OMB No. 0910-01777

OMB Desk Officer: Shannah Koss-McCallum

Social Security Administration

(Call Reports Clearance Officer on 301-965-4149 for copies of package)

1. Certificate of Support—0960-0001—This form collects information which is used by the Social Security Administration to determine if the responder meets the one-half support requirement which is a requirement for entitlement to parents benefits, and is

required to meet the exception to Government Pension Offset (applicable in claims for spouse's benefits). Respondents: Individuals or households; Number of Respondents: 18,000; Frequency of Response: 15 minutes; Estimated Annual Burden: 4,500 hours.

2. Request for Earnings and Benefit Estimate Statement—0960–7004—The information will be used to provide a statement of earnings, quarters of coverage and future benefit estimates to individuals in response to requests. Respondents: Individuals or households; Number of Respondents: 6,000,000; Frequency of Response: 1; Average Burden Per Response: 5 minutes; Estimated Annual Burden: 500,000 hours.

OMB Desk Officer: Justin Kopca

As mentioned above, copies of the information collection clearance packages can be obtained by calling the Reports Clearance Officer, on one of the following numbers:
PHS: (202) 245–2100
HCFA: (301) 966–2088
FSA: (202) 252–5605
SSA: (301) 965–4149
OS: (202) 245–6511
OHDS: (202) 472–4415

Written comments and recommendations for the proposed information collections should be sent directly to the appropriate OMB Desk Officer designated above at the following address: OMB Reports Management Branch, New Executive Office Building, Room 3208, Washington, DC 20503.

Date: January 3, 1989.

James E. Larson,

Acting Deputy Assistant Secretary for Information Business Management.

[FR Doc. 89–283 Filed 1–5–89; 8:45 am]

BILLING CODE 4150–04–M

Agency Forms Submitted to the Office of Management and Budget for Clearance

Each Friday the Department of Health and Human Services (HHS) publishes a list of information collection packages it has submitted to the Office of Management and Budget (OMB) for clearance in compliance with the Paperwork Reduction Act (44 U.S.C. Chapter 35). The following are those packages submitted to OMB since the last list was published on December 30, 1988.

Health Care Financing Administration

(Call Reports Clearance Officer on 301–966–2088 for copies of package)

1. Request for Termination of Premium Hospital and/or Supplementary Medical Insurance—0938–0025—The HCFA–1763 is the form an individual completes when he/she wishes to terminate Medicare coverage. This form is the vehicle by which the SSA Program Services Center is made aware of the beneficiary's desire to withdraw from Medicare. Respondents: Individuals or households; Number of Respondents: 30,000; Frequency of Response: 1; Average Burden Per Response: .166; Estimated Annual Burden: 5,000 hours.

2. Application for Health Insurance Benefits under Medicare for Individuals with Chronic Renal Disease—0938–0080—The law requires the filing of an application to establish Medicare entitlement based on end-stage renal disease. The HCFA–43 is the application form used to obtain information needed to determine Medicare eligibility. It guides district office personnel in securing the required development and becomes a permanent part of the claims. Respondent: Individuals or households; Number of Respondents: 13,500; Frequency of Response: 1; Average Burden Per Response: .43; Estimated Annual Burden: 5,850.

Public Health Service

(Call Reports Clearance Officer on 202–245–2100 for copies of package)

1. Registration of Cosmetic Product Establishment—0910–0027—The registration of cosmetic manufacturers and repackers supplies FDA with current locations for onsite inspection, addresses for information and regulatory mailings, business trading names supplying product distribution sources, and aids FDA in responding to Freedom of Information requests. Respondents: Small businesses; Number of Respondents: 50; Number of Responses Per Respondent: 1; Average Burden Per Response: 0.4; Estimated Annual Burden: 20 hours.

2. Notice of Discontinuance of Commercial Distribution or Cosmetic Product or Cosmetic Raw Material—0910–0029—The purpose of Form FDA 2514 is to notify the FDA of removal from commercial distribution of a cosmetic product or raw material previously filed with the FDA under 21 CFR 730 thereby allowing that data to be maintained in a current state. Respondents: Businesses or other for-profit, Small businesses or organizations; Number of Respondents: 850; Number of Responses Per Respondent: 3; Average Burden Per Response: 0.2; Estimated Annual Burden: 510 hours.

3. Cosmetic Product Ingredient Statement (21 CFR 720)—0910–0030—This information collection assists FDA in evaluating alleged injuries and adverse reactions from use of cosmetic products. It is also utilized in defining and planning analytical and toxicological studies. Data on ingredients and formulations is also available to other government agencies such as GAO, NCI, NIOSH, and the public and industry may access it through FOI. Respondents: Businesses or other for-profit, and small businesses or organizations.

	1st Information Collection	2nd Information Collection
	Title: Registration/ Amended product brand name/ ingredient change	Request for Confidentiality
Number of Respondents	280	12.5
Number of Responses Per Respondent	10	1
Average Burden Per Response	0.5	
Burden Hours	1,400	

As mentioned above, copies of the information collection clearance packages can be obtained by calling the Reports Clearance Officer, on one of the following numbers:
PHS: (202) 245–2100
HCFA: (301) 966–2088
FSA: (202) 252–5605
SSA: (301) 965–4149
OS: (202) 245–6511
OHDS: (202) 472–4415

Written comments and recommendations for the proposed information collections should be sent directly to the appropriate OMB Desk Officer designated above at the following address: OMB Reports Management Branch, New Executive Office Building, Room 3208, Washington, DC 20503.

Date: January 3, 1989.

James E. Larson,

Deputy Assistant Secretary for Information Resources Management.

[FR Doc. 89–284 Filed 1–5–89; 8:45 am]

BILLING CODE 4150–04–M

Food and Drug Administration

[Docket No. 88F–0381]

Betz Laboratories, Inc.; Filing of Food Additive Petition

AGENCY: Food and Drug Administration.

ACTION: Notice.

SUMMARY: The Food and Drug Administration (FDA) is announcing

that a food additive petition has been filed by Betz Laboratories, Inc., proposing that the food additive regulations be amended to provide for the safe use of poly(isopropenylphosphonic acid), sodium salt in the manufacture of paper and paperboard for food-contact use.

FOR FURTHER INFORMATION CONTACT: Marvin D. Mack, 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 9B4114) has been filed by Betz Laboratories, Inc., Somerton Rd., Trevose, PA 19047, 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 poly(isopropenylphosphonic acid), sodium salt in the manufacture of paper and paperboard for food-contact use.

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: December 22, 1988.

Richard J. Ronk,

Acting Director, Center for Food Safety and Applied Nutrition.

[FR Doc. 89–188 Filed 1–5–89; 8:45 am]

BILLING CODE 4160–01–M

[Docket No. 88F–0427]

Cryovac Division of W.R. Grace & Co.; Filing of Food Additive Petition

AGENCY: Food and Drug Administration.

ACTION: Notice.

SUMMARY: The Food and Drug Administration (FDA) is announcing that Cryovac Division, W.R. Grace & Co., has filed a petition proposing that the food additive regulations be amended by raising the limitation on the maximum absorbed dose of radiation that may be used to produce molecular crosslinking of ethylene-vinyl acetate copolymers.

FOR FURTHER INFORMATION CONTACT: Laura M. Tarantino, Center for Food Safety and Applied Nutrition (HFF–330), Food and Drug Administration, 200 C St. SW., Washington, DC 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 Cryovac Division of W.R. Grace & Co., P.O. Box 464, Duncan, SC 29334, has filed a petition (FAP 9M4117) proposing that § 177.1350 *Ethylene-vinyl acetate copolymers* (21 CFR 177.1350) of the food additive regulations be amended by raising the limitation, in paragraph (d)(1) and (d)(3), on the maximum absorbed dose of radiation that may be used to produce molecular crosslinking of ethylene-vinyl acetate copolymers.

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: December 27, 1988.

Richard J. Ronk,

Acting Director, Center for Food Safety and Applied Nutrition.

[FR Doc. 89–184 Filed 1–5–89; 8:45 am]

BILLING CODE 4160–01–M

[Docket No. 88F–0404]

Mitsui Petrochemical Industries, Ltd.; Filing of Food Additive Petition

AGENCY: Food and Drug Administration.

ACTION: Notice.

SUMMARY: The Food and Drug Administration (FDA) is announcing that Mitsui Petrochemical Industries, Ltd., has filed a petition proposing that the food additive regulations be amended to provide for the safe use of ethylene/1,3-phenyleneoxyethylene isophthalate/terephthalate copolymer as a nonfood contact layer of food packaging laminates intended for use in contact with food.

FOR FURTHER INFORMATION CONTACT: Gillian Robert-Baldo, 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 Mitsui Petrochemical Industries, Ltd., Kasumigaseki Bldg., P.O. Box 90, 2–5 Kasumigaseki 3-chrome, Chiyoda-Ku, Tokyo 100, Japan, has filed a petition (FAP 8B4107), proposing that § 177.1395 *Laminate structures for use at temperatures between 120 °F and 250 °F* (21 CFR 177.1395) be amended to provide for the safe use of ethylene/1,3-phenyleneoxyethylene isophthalate/terephthalate copolymer as a nonfood contact layer of food packaging laminates 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).

Dated: December 27, 1988.

Richard J. Ronk,

Acting Director, Center for Food Safety and Applied Nutrition.

[FR Doc. 89–185 Filed 1–5–89; 8:45 am]

BILLING CODE 4160–01–M

[Docket No. 82F–0309]

Monsanto Chemical Corp.; Withdrawal of Food Additive Petition

AGENCY: Food and Drug Administration.

ACTION: Notice.

SUMMARY: The Food and Drug Administration (FDA) is announcing the withdrawal, without prejudice to a future filing, of food additive petition 2B3655 proposing that the food additive regulations be amended to provide for the safe use of a mixture of partially hydrogenated terphenyl and quaterphenyl as components of adhesives for food-contact use.

FOR FURTHER INFORMATION CONTACT: Marvin D. Mack, 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: In the **Federal Register** of October 19, 1982 (47 FR 46576), FDA published a notice that it had filed a petition (FAP 2B3655) submitted by Monsanto Industrial Chemicals Co. (later named Monsanto Chemical Co.), 800 North Lindberg Blvd., St. Louis, MO 63167, proposing to amend § 175.105 *Adhesives* (21 CFR 175.105) of the food additive regulations to provide for the safe use of a mixture of partially hydrogenated terphenyl and quarterphenyl as components of adhesives for food-contact use. Monsanto Chemical Co. has now