standards in section 4 of the BHC Act. Unless otherwise noted, nonbanking activities will be conducted throughout the United States.

Unless otherwise noted, comments regarding each of these applications must be received at the Reserve Bank indicated or the offices of the Board of Governors not later than October 10, 1997.

A. Federal Reserve Bank of Cleveland (Jeffery Hirsch, Banking Supervisor) 1455 East Sixth Street, Cleveland, Ohio 44101-2566:

1. F.N.B. Corporation, Hermitage, Pennsylvania, and Southwest Banks, Inc., Naples, Florida; to acquire 100 percent of the voting shares of Mercantile Bank of Southwest Florida, Naples, Florida.

Board of Governors of the Federal Reserve System, September 11, 1997.

Jennifer J. Johnson,

Deputy Secretary of the Board.

[FR Doc. 97–24580 Filed 9–15–97; 8:45 am]

BILLING CODE 6210–01–F

GENERAL SERVICES ADMINISTRATION

Federal Acquisition Policy Division, FAR Secretariat; Cancellation of an Optional Form

AGENCY: General Services Administration.

ACTION: Notice.

SUMMARY: The Federal Acquisition Regulations eliminated the need for Optional Form 333, Procurement Integrity Certification For Procurement Officials removing the regulations that required its use. Therefore, OF 333 is cancelled. This deleted requirement was effective January 1, 1997.

FOR FURTHER INFORMATION CONTACT: Mr. Ralph DeStefano, (202) 501–1758.

DATES: Effective upon publication in the **Federal Register** September 16, 1997.

Dated: September 8, 1997.

Barbara M. Williams,

Deputy Standard and Optional Forms Management Officer.

[FR Doc. 97–24513 Filed 9–15–97; 8:45 am]

BILLING CODE 6820–34–M

GOVERNMENT PRINTING OFFICE

Depository Library Council to the Public Printer; Meeting

The Depository Library Council to the Public Printer (DLC) will hold its Fall 1997 meeting on Monday, October 20, 1997, through Thursday, October 23, 1997, in Clearwater Beach, Florida. The meeting sessions will take place from 9 a.m. until 5 p.m. on Monday, Tuesday, Wednesday, and from 9 a.m. until 12 noon on Thursday. The sessions will be held at the Adam's Mark Caribbean Gulf Hotel, 430 South Gulfview Boulevard, Clearwater Beach, Florida. The purpose of this meeting is to discuss the Federal Depository Library Program. The meeting is open to the public.

A limited number of hotel rooms have been reserved at the Adam's Mark Caribbean Gulf Hotel for anyone needing hotel accommodations. Telephone: 800–444–ADAM, 813–443–5714; FAX: 813–442–8389. Please specify the Depository Library Council when you contact the hotel. Room cost per night is $72.

Michael F. DiMario,

Public Printer.

[FR Doc. 97–24508 Filed 9–15–97; 8:45 am]

BILLING CODE 1520–01–P

DEPARTMENT OF HEALTH AND HUMAN SERVICES

Office of the Secretary

Agency Information Collection Activities: Submission for OMB Review; Comment Request

The Department of Health and Human Services, Office of the Secretary publishes a list of information collections it has submitted to the Office of Management and Budget (OMB) for clearance in compliance with the Paperwork Reduction Act of 1995 (44 U.S.C. Chapter 35) and 5 CFR 1320.5. The following are those information collections recently submitted to OMB.

National Study of Assisted Living Facilities for the Frail Elderly—New—The goal of this study is to determine where assisted living fits in the continuum of long term care and to examine its potential for addressing the needs of elderly persons with disabilities. The study will address such topics as trends in supply and demand; barriers to development; the effect of key assisted living features on resident satisfaction and other outcomes. Surveys of operators, staff and elderly residents will be conducted.

Respondents: Assisted Living Facilities operators, staff and residents—Burden Information on Operator Screen—*Number of Responses:* 1958; *Burden per Response:* 10 minutes; *Total Screen Burden:* 326 hours—Burden Information for Operator Telephone Interview—*Number of Responses:* 230; *Burden per Response:* 20 minutes; *Total Burden:* 77 hours—Burden Information for Operator In-Person Interview and Supplement—*Number of Responses:* 690; *Burden per Response:* 45 minutes; *Total Burden:* 518 hours—Burden Information for Staff Interview—*Number of Responses:* 1380; *Burden per Response:* 20 minutes; *Total Burden:* 460 hours—Burden Information for Resident Interview—*Number of Responses:* 2496; *Burden per Response:* 30 minutes; *Total Burden:* 1248 hours—Burden Information for Resident Proxy Interview—*Number of Responses:* 1230; *Burden per Response:* 15 minutes; *Total Burden:* 308 hours—Burden Information for Family Member Interview—*Number of Responses:* 897; *Burden per Response:* 20 minutes; *Total Burden:* 299 hours—Burden Information for Discharged Resident Interview—*Number of Responses:* 117; *Burden per Response:* 10 minutes; *Total Burden:* 20 hours—Burden Information for Discharged Resident Proxy Interview—*Number of Responses:* 470; *Burden per Response:* 12 minutes; *Total Burden:* 94 hours—*Total Burden for the Survey:* 3,350 hours.

OMB Desk Officer: Allison Eydt.

Copies of the information collection packages listed above can be obtained by calling the OS Reports Clearance Officer on (202) 690–6207. Written comments and recommendations for the proposed information collection should be sent directly to the OMB desk officer designated above at the following address: Human Resources and Housing Branch, Office of Management and Budget, New Executive Office Building, Room 10235, 725 17th Street NW., Washington, DC 20503.

Comments may also be sent to Cynthia Agens Bauer, OS Reports Clearance Officer, Room 503H, Humphrey Building, 200 Independence Avenue SW., Washington, DC, 20201. Written comments should be received within 30 days of this notice.

Dated: September 5, 1997.

Dennis P. Williams,

Deputy Assistant Secretary, Budget.

[FR Doc. 97–24434 Filed 9–15–97; 8:45 am]

BILLING CODE 4150–04–M

DEPARTMENT OF HEALTH AND HUMAN SERVICES

Centers for Disease Control and Prevention

[30DAY–23–97]

Agency Forms Undergoing Paperwork Reduction Act Review

The Centers for Disease Control and Prevention (CDC) publishes a list of

information collection requests under review by the Office of Management and Budget (OMB) in compliance with the Paperwork Reduction Act (44 U.S.C. Chapter 35). To request a copy of these requests, call the CDC Reports Clearance Office on (404) 639–7090. Send written comments to CDC, Desk Officer; Human Resources and Housing Branch, New Executive Office Building, Room 10235; Washington, DC 20503. Written comments should be received within 30 days of this notice.

Proposed Projects

1. Multi-Center Cohort Study to Assess the Risk and Consequences of Hepatitis C Virus Transmission from Mother to Infant (0920–0344)—Reinstatement—The purpose of the study is to determine the incidence of vertical hepatitis C virus (HCV) transmission, to assess risk factors for vertical HCV transmission, to assess the clinical course of disease among infants with HCV infection, and to assess diagnostic methods for detecting HCV infection in infants. Respondents for the study will be anti-HCV positive mothers.

There is no cost to the respondents. They will be remunerated for travel costs; provided well-child visits and free vaccinations for infants enrolled in the study; and, provided anti-HCV testing to all family members free of charge. The total annual burden hours are 277.

Respondents	Form name	Number of respondents	Number of responses/ respondent	Avg. burden/ response (in hrs.)
Individual Mothers	Form A	300	1	0.25
Mothers	Form B	1200	1	0.25
Mothers	Form C	300	1	0.10
Mothers	Form D	300	1	0.25
Family members	Form E	700	1	0.25
Mothers	Form F	300	1	0.25
Mothers	Form G	300	8	0.10

*The annualized response burden is estimated to be 970 hours/3.5 years= 277 hours. (Target enrollment in the study is 300; the target population will be drawn from those who complete Form B. Family members will complete Form E.)

2. Information Collection Procedures for Evaluating Toxicological Profiles—New—The Agency for Toxic Substances and Disease Registry (ATSDR) prepares toxicological profiles in accordance with guidelines developed with guidelines developed by ATSDR and EPA and each profile is revised and republished as necessary, but no less often than every three years. The principal audiences for the toxicological profiles are health professionals at the federal, state, and local levels, interested private sector organizations and groups, and members of the public.

This is a request for approval to collect information in the profiles from users on: (a) Affiliation of users of the profiles, (b) clarity of discussion in the profiles, (c) consistency of information in the profiles, (d) completeness of information in the profile, and (e) utility of information in the profile.

The information will be used in an effort to maintain customer satisfaction concerning use of the profiles by these multi-disciplinary users. This will also ensure that we continue to provide a client-oriented product. This effort will be accomplished through enhancement of the built-in system used for updating existing toxicological profiles and improving the utility of newly developed profiles by use of these user surveys.

The only cost to respondents will be the time to complete the form, which we estimate at less than 15 minutes per respondent. We expect respondents of the toxicological profile survey to come from a wide range of occupational and professional backgrounds and have an average hourly wage of $15. The total annual burden hours are 750.

Respondents	Number of respondents	Number of responses/respondent	Average burden/response
Questionnaire	6000	1	0.25

3. NIOSH Training Grants, 42 CFR part 86, Application And Regulations—(0920–02610)—Reinstatement—Public Law 91–596 authorizes CDC/NIOSH to support ''education programs that provide an adequate supply of qualified personnel * * * by grants or contracts'' to assure a safe and healthful work environment. NIOSH awards grants for both short-term and long-term training to academic institutions and other organizations interested in providing training for professionals. Grants are also provided to Educational Resource Centers (ERCs) which provide multi disciplinary graduate training for industrial hygienists, occupational physicians, occupational health nurses, safety professionals and other occupational health-related disciplines in addition to continuing education for practicing professionals and outreach in the Region. 42 CFR Part 86, ''Grants for Education Programs in Occupational Safety and Health, Subpart B-Occupational Safety and Health Training, provides guidelines for implementing Public Law 91–596. The training grant application form is used by the National Institute of Occupational Safety and Health to collect information from potential applicants. The information is used to determine the eligibility of applicants for review, to calculate the amount of each award and to judge the merit of each application. CDC Form 2.145A is used for new and competing continuation grants; CDC Form 2.145B is used for non-competing awards. If this information is not collected, grants cannot be reviewed and awarded. The total annual burden hours are 4,635.

Respondents	Number of respondents	Number of responses/respondent	Average burden/response (in hrs.)
Training Grant Application			
ERC	3	1	348
Training Grant	12	1	63
Continuation Grant Application			
ERC	11	1	189
Training Grant	28	1	27

Dated: September 10, 1997.

Wilma G. Johnson,

Acting Associate Director for Policy Planning And Evaluation, Centers for Disease Control and Prevention (CDC).

[FR Doc. 97–24482 Filed 9–15–97; 8:45 am]

BILLING CODE 4163–18–P

DEPARTMENT OF HEALTH AND HUMAN SERVICES

Food and Drug Administration

[Docket No. 97F–0375]

General Electric Co.; Filing of Food Additive Petition

AGENCY: Food and Drug Administration, HHS.

ACTION: Notice.

SUMMARY: The Food and Drug Administration (FDA) is announcing that General Electric Co. has filed a petition proposing that the food additive regulations be amended to provide for the expanded safe use of phosphorous acid, cyclic butylethyl propanediol, 2,4,6-tri-*tert*-butylphenyl ester, which may contain up to 1 percent by weight of triisopropanolamine, as an antioxidant and/or stabilizer in olefin copolymers intended for use in contact with food.

FOR FURTHER INFORMATION CONTACT: Vir D. Anand, Center for Food Safety and Applied Nutrition (HFS–215), Food and Drug Administration, 200 C St. SW., Washington, DC 20204, 202–418–3081.

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 food additive petition (FAP 7B4553) has been filed by General Electric Co., One Lexan Lane, Mt. Vernon, IN 47620–9364. The petition proposes to amend the food additive regulations in § 178.2010 *Antioxidants and/or stabilizers for polymers* (21 CFR 178.2010) to provide for the expanded safe use of phosphorous acid, cyclic butylethyl propanediol, 2,4,6-tri-*tert*-butylphenyl ester, which may contain up to 1 percent by weight of triisopropanolamine, as an antioxidant and/or stabilizer for olefin copolymers complying with 21 CFR 177.1520(c), items 3.1 and 3.2, intended for use in contact with food.

The agency has determined under 21 CFR 25.32(i) that this action is of the type that does not individually or cumulatively have a significant effect on the human environment. Therefore, neither an environmental assessment nor an environmental impact statement is required.

Dated: August 29, 1997.

Alan M. Rulis,

Director, Office of Premarket Approval, Center for Food Safety and Applied Nutrition.

[FR Doc. 97–24424 Filed 9–15–97; 8:45 am]

BILLING CODE 4160–01–F

DEPARTMENT OF HEALTH AND HUMAN SERVICES

Food and Drug Administration

Advisory Committee; Notice of Meeting

AGENCY: Food and Drug Administration, HHS.

ACTION: Notice.

This notice announces a forthcoming meeting of a public advisory committee of the Food and Drug Administration (FDA). The meeting will be open to the public.

Name of Committee: Obstetrics and Gynecology Devices Panel of the Medical Devices Advisory Committee.

General Function of the Committee: To provide advice and recommendations to the agency on FDA regulatory issues.

Date and Time: The meeting will be held on October 6, 1997, 8:30 a.m. to 5 p.m., and October 7, 1997, 8:30 a.m. to 2 p.m.

Location: Corporate Bldg., conference room 020B, 9200 Corporate Blvd., Rockville, MD.

Contact Person: Elisa D. Harvey, Center for Devices and Radiological Health (HFZ–470), Food and Drug Administration, 9200 Corporate Blvd., Rockville, MD 20850, 301–594–1180, or FDA Advisory Committee Information Line, 1–800–741–8138 (301–443–0572 in the Washington, DC area), Obstetrics and Gynecology Devices Panel, code 12524. Please call the Information Line for up-to-date information on this meeting.

Agenda: On October 6, 1997, the committee will discuss and make recommendations on a premarket approval application for a thermal endometrial ablation device. On October 7, 1997, the committee will discuss and advise FDA on a petition for reclassification of home uterine activity monitors from class III (premarket approval) to class II (special controls).

Procedure: Interested persons may present data, information, or views, orally or in writing, on issues pending before the committee. Written submissions may be made to the contact person by September 29, 1997. Oral presentations from the public will be scheduled between approximately 8:30 a.m. and 9:30 a.m. on October 6 and 7, 1997. Time allotted for each presentation may be limited. Those desiring to make formal oral presentations should notify the contact person before September 29, 1997, and submit a brief statement of the general nature of the evidence or arguments they wish to present, the names and addresses of proposed participants, and an indication of the approximate time required to make their presentation.

Notice of this meeting is given under the Federal Advisory Committee Act (5 U.S.C. app. 2).

Dated: September 8, 1997

Michael A. Friedman,

Deputy Commissioner for Operations.

[FR Doc. 97–24509 Filed 9–15–97; 8:45 am]

BILLING CODE 4160–01–F