Ocean Freight Forwarder—Ocean Transportation Intermediary Applicants

Access Freight Forwarders, LLC, 184 East Bay Street, Suite 202, Charleston, SC 29401, Officers: David Holst, Jr., President (Qualifying Individual), John H. Chapman, Treasurer.

Gruen International, Inc., 6310 N. Port Washington Road, Milwaukee, WI 53217, Officers: Michael J. Karman, Asst. Vice President (Qualifying Individual), Steven Gruen, President.

Dated: October 10, 2003.

Bryant L. VanBrakle,

Secretary.

[FR Doc. 03–26237 Filed 10–16–03; 8:45 am]

BILLING CODE 6730–01–P

FEDERAL RESERVE SYSTEM

Formations of, Acquisitions by, and Mergers of Bank Holding Companies

The companies listed in this notice have applied to the Board for approval, pursuant to the Bank Holding Company Act of 1956 (12 U.S.C. 1841 *et seq.*) (BHC Act), Regulation Y (12 CFR Part 225), and all other applicable statutes and regulations to become a bank holding company and/or to acquire the assets or the ownership of, control of, or the power to vote shares of a bank or bank holding company and all of the banks and nonbanking companies owned by the bank holding company, including the companies listed below.

The applications listed below, as well as other related filings required by the Board, are available for immediate inspection at the Federal Reserve Bank indicated. The application also will be available for inspection at the offices of the Board of Governors. Interested persons may express their views in writing on the standards enumerated in the BHC Act (12 U.S.C. 1842(c)). If the proposal also involves the acquisition of a nonbanking company, the review also includes whether the acquisition of the nonbanking company complies with the standards in section 4 of the BHC Act (12 U.S.C. 1843). Unless otherwise noted, nonbanking activities will be conducted throughout the United States. Additional information on all bank holding companies may be obtained from the National Information Center website at *www.ffiec.gov/nic/*.

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 November 10, 2003.

A. Federal Reserve Bank of Chicago (Phillip Jackson, Applications Officer) 230 South LaSalle Street, Chicago, Illinois 60690-1414:

1. Northern States Financial Corporation, Waukegan, Illinois; to merge with Round Lake Bankcorp, Inc., Round Lake, Illinois, and thereby indirectly acquire First State Bank of Round Lake, Round Lake Beach, Illinois.

Board of Governors of the Federal Reserve System, October 10, 2003.

Robert deV. Frierson,

Deputy Secretary of the Board.

[FR Doc. 03–26233 Filed 10–16–03; 8:45 am]

BILLING CODE 6210–01–S

FEDERAL RESERVE SYSTEM

Sunshine Act Meeting

AGENCY HOLDING THE MEETING: Board of Governors of the Federal Reserve System.

TIME AND DATE: 11:15 a.m., Wednesday, October 15, 2003. The business of the Board requires that this meeting be held with less than one week's advance notice to the public, and no earlier announcement of the meeting was practicable.

PLACE: Marriner S. Eccles Federal Reserve Board Building, 20th and C Streets, NW., Washington, DC 20551.

STATUS: Closed.

MATTERS TO BE CONSIDERED:

1. Personnel actions (appointments, promotions, assignments, reassignments, and salary actions) involving individual Federal Reserve System employees.

2. Any items carried forward from a previously announced meeting.

FOR FURTHER INFORMATION CONTACT: Michelle A. Smith, Director, Office of Board Members; 202–452–2955.

SUPPLEMENTARY INFORMATION: You may call 202–452–3206 beginning at approximately 5 p.m. two business days before the meeting for a recorded announcement of bank and bank holding company applications scheduled for the meeting; or you may contact the Board's Web site at *http://www.federalreserve.gov* for an electronic announcement that not only lists applications, but also indicates procedural and other information about the meeting.

Dated: October 14, 2003.

Robert deV. Frierson,

Deputy Secretary of the Board.

[FR Doc. 03–26355 Filed 10–15–03; 8:49 am]

BILLING CODE 6210–01–P

DEPARTMENT OF HEALTH AND HUMAN SERVICES

Centers for Disease Control and Prevention

National Center for Health Statistics; Meeting

National Center for Health Statistics (NCHS), Classifications and Public Health Data Standards Staff, announces the following meeting.

Name: ICD–9–CM Coordination and Maintenance Committee meeting.

Time and Date: 9 a.m.–4 p.m., December 4–5, 2003.

Place: Centers for Medicare and Medicaid Services (CMS) Auditorium, 7500 Security Boulevard, Baltimore, Maryland.

Status: Open to the public.

Purpose: The ICD–9–CM Coordination and Maintenance (C&M) Committee will hold its final meeting of the 2003 calendar year cycle on Thursday and Friday December 4–5, 2003. The C&M meeting is a public forum for the presentation of proposed modifications to the International Classification of Diseases, Ninth-Revision, Clinical Modification.

Matters To Be Discussed: Agenda items include:

ICD–10–CM update
West Nile virus with and without encephalitis
Alpha-1 antitrypsin deficiency and other metabolic conditions
Long-term (current) use of aspirin
Multiple sclerosis
Sleep disorders
Genital prolapse
Bethesda system
Chondritis of ear
Worn out joint prosthesis
Awaiting heart transplant status
Dental expansions
Decubitus ulcers
Automatic implantable cardioverter/defibrillator (AICD) check
Spinal procedures—nucleus replacement device
Laparoscopic/Thorascopic approaches
Insertion/replacement of neurostimulator components
Axial flow left ventricular assist device
Prevention of vein graft failure
Intravascular ultrasound (IVUS)
ICD–10—Procedure classification system (ICD–10–PCS) update
Addenda

For Further Information Contact: Amy Blum, Medical Classification Specialist, Classifications and Public Health Data Standards Staff, NCHS, 3311 Toledo Road, rm. 2402, Hyattsville, Maryland 20782, telephone 301/458–4106 (diagnosis), Amy Gruber, Health Insurance Specialist, Division of Acute Care, CMS, 7500 Security Blvd., Room C4–07–07, Baltimore, Maryland 21244 telephone (410) 786–1542 (procedures).

Notice: In the interest of security, (CMS) has instituted stringent procedures for entrance into the building by non-government employees. Persons without a government I.D. will need to show a photo

I.D. and sign-in at the security desk upon entering the building.

Because of increased security requirements, those who wish to attend a specific ICD–9–CM C&M meeting in the CMS auditorium must submit their name and organization for addition to the meeting visitor list. Those wishing to attend the December 4–5, 2003 meeting must submit their name and organization by November 28, 2003 for inclusion on the visitor list.

This visitor list will be maintained at the front desk of the CMS building and used by the guards to admit visitors to the meeting. Those who attended previous ICD–9–CM C&M meetings will no longer be automatically added to the visitor list. You must request inclusion of your name prior to each meeting you attend.

Send your name and organization to one of the following by November 28, 2003 in order to attend the December 4–5, 2003 meeting: Pat Brooks *pbrooks1@cms.hhs.gov* (410) 786–5318. Ann Fagan *afagan@cms.hhs.gov* (410) 786–5662. Amy Gruber *agruber@cms.hhs.gov* (410) 786–1542.

Notice: This is a public meeting. However, because of fire code requirements, should the number of attendants meet the capacity of the room, the meeting will be closed.

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 CDC and the Agency for Toxic Substances and Disease Registry.

Dated: October 10, 2003.

Diane Allen,

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

[FR Doc. 03–26272 Filed 10–16–03; 8:45 am]

BILLING CODE 4160–18–P

DEPARTMENT OF HEALTH AND HUMAN SERVICES

Administration for Children and Families

President's Committee for People With Intellectual Disabilities (PCPID); Notice of Meeting

AGENCY: President's Committee for People with Intellectual Disabilities (PCPID); Department of Health and Human Services.

ACTION: Corrected notice of meeting.

SUMMARY: This document clarifies and corrects the notice that was published in the **Federal Register** on October 9, 2003 (68 FR 58352). It corrects a statement that a portion of the meeting would be closed to the public. The full Committee meeting of the President's Committee for People with Intellectual Disabilities will be open to the public pursuant to section 10(a)(1) of the Federal Advisory Committee Act (5 U.S.C. App. 2). Subcommittees of the Committee will have breakout working sessions from 1:30 p.m. to 4 p.m. for the purpose of preliminary discussions on issues of the PCPID. This notice is filed less than 15 calendar days prior to the meeting date due to scheduling conflicts.

FOR FURTHER INFORMATION CONTACT: Sally Atwater, Executive Director, President's Committee for People with Intellectual Disabilities. Aerospace Center Building, Suite 701, 370 L'Enfant Promenade, SW., Washington, DC 20447, Telephone (202) 619–0634, Fax (202) 205–9519, E-mail: *satwater@acf.hhs.gov.*

Dated: October 9, 2003.

Sally Atwater,

Executive Director, President's Committee for People with Intellectual Disabilities.

[FR Doc. 03–26277 Filed 10–16–03; 8:45 am]

BILLING CODE 4184–01–M

DEPARTMENT OF HEALTH AND HUMAN SERVICES

Food and Drug Administration

[Docket No. 2003N–0463]

Agency Information Collection Activities; Proposed Collection; Comment Request; Infant Formula Requirements

AGENCY: Food and Drug Administration, HHS.

ACTION: Notice.

SUMMARY: The Food and Drug Administration (FDA) is announcing an opportunity for public comment on the proposed collection of certain information by the agency. Under the Paperwork Reduction Act of 1995 (the PRA), Federal agencies are required to publish notice in the **Federal Register** concerning each proposed collection of information including each proposed extension of an existing collection of information, and to allow 60 days for public comment in response to the notice. This notice solicits comments on information collection regarding the manufacture of infant formula, including infant formula labeling, quality control procedures, notification requirements, and recordkeeping.

DATES: Submit written or electronic comments on the collection of information by December 16, 2003.

ADDRESSES: Submit electronic comments on the collection of information to *http://www.fda.gov/dockets/ecomments.* Submit written comments on the collection of information to the Division of Dockets Management (HFA–305), Food and Drug Administration, 5630 Fishers Lane, rm. 1061, Rockville, MD 20852. All comments should be identified with the docket number found in brackets in the heading of this document.

FOR FURTHER INFORMATION CONTACT: Peggy Robbins, Office of Management Programs (HFA–250), Food and Drug Administration, 5600 Fishers Lane, Rockville, MD 20857, 301–827–1223.

SUPPLEMENTARY INFORMATION: Under the PRA (44 U.S.C. 3501–3520), Federal agencies must obtain approval from the Office of Management and Budget (OMB) for each collection of information they conduct or sponsor. ''Collection of information'' is defined in 44 U.S.C. 3502(3) and 5 CFR 1320.3(c) and includes agency requests or requirements that members of the public submit reports, keep records, or provide information to a third party. Section 3506(c)(2)(A) of the PRA (44 U.S.C. 3506(c)(2)(A)) requires Federal agencies to provide a 60-day notice in the **Federal Register** concerning each proposed collection of information, including each proposed extension of an existing collection of information, before submitting the collection to OMB for approval. To comply with this requirement, FDA is publishing notice of the proposed collection of information set forth in this document.

With respect to the following collection of information, FDA invites comments on these topics: (1) Whether the proposed collection of information is necessary for the proper performance of FDA's functions, including whether the information will have practical utility; (2) the accuracy of FDA's estimate of the burden of the proposed collection of information, including the validity of the methodology and assumptions used; (3) ways to enhance the quality, utility, and clarity of the information to be collected; and (4) ways to minimize the burden of the collection of information on respondents, including through the use of automated collection techniques, when appropriate, and other forms of information technology.

Infant Formula Requirements—21 CFR Parts 106 and 107 (OMB Control Number 0910–0256)—Extension

Statutory requirements for infant formula under the Federal Food, Drug, and Cosmetic Act (the act) are intended to protect the health of infants and include a number of reporting and recordkeeping requirements. Among other things, section 412 of the act (21 U.S.C. 350a) requires manufacturers of infant formula to establish and adhere to quality control procedures, notify FDA