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 June 16, 2003.

A. Federal Reserve Bank of Dallas (W. Arthur Tribble, Vice President) 2200 North Pearl Street, Dallas, Texas 75201– 2272:

1. Inwood Bancshares, Inc., Dallas, Texas, and Inwood Delaware, Inc., Dover, Delaware; to acquire 100 percent of WB&T Bancshares, Inc., Duncanville, Texas, and thereby indirectly acquire WB&T Delaware, Inc., Duncanville, Texas, and Western Bank & Trust, Duncanville, Texas.

Board of Governors of the Federal Reserve System, May 15, 2003.

Robert deV. Frierson,

Deputy Secretary of the Board.
[FR Doc. 03–12655 Filed 5–19–03; 8:45 am]
BILLING CODE 6210–01–S

FEDERAL RESERVE SYSTEM

Notice of Proposals to Engage in Permissible Nonbanking Activities or to Acquire Companies that are Engaged in Permissible Nonbanking Activities

The companies listed in this notice have given notice under section 4 of the Bank Holding Company Act (12 U.S.C. 1843) (BHC Act) and Regulation Y (12 CFR Part 225) to engage *de novo*, or to acquire or control voting securities or assets of a company, including the companies listed below, that engages

either directly or through a subsidiary or other company, in a nonbanking activity that is listed in § 225.28 of Regulation Y (12 CFR 225.28) or that the Board has determined by Order to be closely related to banking and permissible for bank holding companies. Unless otherwise noted, these activities will be conducted throughout the United States.

Each notice is available for inspection at the Federal Reserve Bank indicated. The notice also will be available for inspection at the offices of the Board of Governors. Interested persons may express their views in writing on the question whether the proposal complies with the standards of section 4 of the BHC Act. 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 the applications must be received at the Reserve Bank indicated or the offices of the Board of Governors not later than June 3, 2003.

A. Federal Reserve Bank of New York (Betsy Buttrill White, Senior Vice President) 33 Liberty Street, New York, New York 10045-0001:

1. Commerzbank Aktiengesellschaft, Frankfurt, Germany; to engage de novo through its subsidiary, Commerzbank Capital Markets Corporation, New York, New York, in serving as the investment adviser to and the general partner or general member of, and holding and placing equity interest in, certain investment funds which invest only in securities and other assets which are permitted to be held directly under the Bank Holding Company Act (the "BHC Act"), including acting as a commodity pool operator for private investment funds organized as commodity pools that invest in assets which a bank holding company is permitted to hold directly under the BHC Act, as permitted under Board precedent, see First Security Corporation, 85 Fed. Res. Bull. 207 (1999), Dresdner Bank AG, 84 Fed. Res. Bull. 361 (1998).

Board of Governors of the Federal Reserve System, May 14, 2003.

Robert deV. Frierson,

Deputy Secretary of the Board. [FR Doc. 03–12501 Filed 5–19–03; 8:45 am] BILLING CODE 6210–01–S

DEPARTMENT OF HEALTH AND HUMAN SERVICES

Centers for Disease Control and Prevention

[60Day-03-67]

Proposed Data Collections Submitted for Public Comment and Recommendations

In compliance with the requirement of section 3506(c)(2)(A) of the Paperwork Reduction Act of 1995 for opportunity for public comment on proposed data collection projects, the Centers for Disease Control and Prevention (CDC) will publish periodic summaries of proposed projects. To request more information on the proposed projects or to obtain a copy of the data collection plans and instruments, call the CDC Reports Clearance Officer on (404) 498–1210.

Comments are invited on: (a) Whether the proposed collection of information is necessary for the proper performance of the functions of the agency, including whether the information shall have practical utility; (b) the accuracy of the agency's estimate of the burden of the proposed collection of information; (c) ways to enhance the quality, utility, and clarity of the information to be collected; and (d) ways to minimize the burden of the collection of information on respondents, including through the use of automated collection techniques or other forms of information technology. Send comments to Anne O'Connor, CDC Assistant Reports Clearance Officer, 1600 Clifton Road, MS-D24, Atlanta, GA 30333. Written comments should be received within 60 days of this notice.

Proposed Project: Evaluating the Effectiveness of the Asthma Intervention Program, Power Breathing—New—National Center for Environmental Health (NCEH), Centers for Disease Control and Prevention (CDC).

Background

The Centers for Disease Control and Prevention (CDC) seeks to conduct an evaluation of the effectiveness of the asthma intervention program, Power Breathing, in two school districts in Kansas City, KS and Fredericksburg, VA. The overall aim of this program, developed by the Asthma and Allergy Foundation of America, is to provide adolescents with a basic understanding of asthma and its management in a peerfriendly environment and to empower them to take control of their asthma on a personal level. The proposed data collection for the evaluation will provide feedback to CDC about the

usefulness and cost-effectiveness of this teen asthma intervention program. Sample participants will come from students, parents, program facilitators, and school personnel (school nurses and teachers) in the selected two school districts. Self-administered questionnaires will be given to students at baseline (pre-intervention program), immediately post-program, and at 6-months post-program, while parents

receive baseline and 6-month postprogram surveys. The student survey will focus on: knowledge, attitudes, and behaviors regarding their asthma; perception of their health status and quality of life; assessment of the program; and impact of the program on their asthma management skills. Parents will be asked about their child's asthma condition, assessment of the program, and cost-related issues for their child's asthma. Individual, one-time interviews will be conducted with program facilitators and school personnel regarding their perceptions of the intervention program and its impact on the students. Two focus groups will be conducted with students post-program to obtain additional, in-depth information about their perceptions of the program.

| Respondents | Number of respondents | Number of responses/ respondent | Average burden/ response (in hrs.) | Total burden (in hrs.) |
|-----------------------|-----------------------|---------------------------------|---|------------------------------|
| Students: | | | | |
| Baseline | 524 | 1 | 30/60 | 262 |
| Post-program | 524 | 1 | 15/60 | 131 |
| 6-month follow-up | 524 | 1 | 30/60 | 262 |
| Focus group | 16 | 1 | 1 | 16 |
| Parents: | | | | |
| Baseline | 524 | 1 | 10/60 | 87 |
| 6-month follow-up | 524 | 1 | 15/60 | 131 |
| Program facilitators: | | | | |
| Interview | 6 | 1 | 40/60 | 4 |
| Program sessions | 6 | 12 | 30/60 | 36 |
| School nurses: | | | | |
| School profile | 6 | 1 | 10/60 | 1 |
| Record abstraction | 6 | 87 | 10/60 | 87 |
| Interview | 6 | 1 | 40/60 | 4 |
| Teachers Interview | 12 | 1 | 40/60 | 8 |
| Total | | | | 1029 |

Dated: May 13, 2003.

Thomas Bartenfeld,

Acting Associate Director for Policy, Planning and Evaluation, Centers for Disease Control and Prevention.

[FR Doc. 03–12535 Filed 5–19–03; 8:45 am] BILLING CODE 4163–18–P

DEPARTMENT OF HEALTH AND HUMAN SERVICES

Food and Drug Administration

Endocrinologic and Metabolic Drugs Advisory Committee; Amendment of Notice

AGENCY: Food and Drug Administration, HHS.

ACTION: Notice.

The Food and Drug Administration (FDA) is announcing an amendment to the notice of the meeting of the Endocrinologic and Metabolic Drugs Advisory Committee. This meeting was announced in the **Federal Register** of May 6, 2003 (68 FR 24003). The amendment is being made to reflect a change in the *Agenda* portion of the document. There are no other changes. **FOR FURTHER INFORMATION CONTACT:**

Dornette Spell-LeSane, Center for Drug Evaluation and Research (HFD–21), Food and Drug Administration, 5600 Fishers Lane (for express delivery, 5630 Fishers Lane, rm. 1093), Rockville, MD 20857, 301-827-7001, or FDA Advisory Committee Information Line, 1–800– 741–8138 (301–443–0572 in the Washington, DC area), code 12536. Please call the Information Line for upto-date information on this meeting.

SUPPLEMENTARY INFORMATION: In the **Federal Register** of May 6, 2003 (68 FR 24003), FDA announced that a meeting of the Endocrinologic and Metabolic Drugs Advisory Committee would be held on June 10, 2003. On page 24003, in the third column, the *Agenda* portion of the meeting is amended to read as follows:

Agenda: The committee will discuss supplemental new drug application (sNDA) 19–604/S–033 HUMATROPE (somatropin recombinant deoxyribonucleic acid (rDNA) origin) for injection), Eli Lilly and Co., for the proposed indication of treatment of nongrowth hormone deficiency short stature.

The notice of this meeting is given under the Federal Advisory Committee Act (5 U.S.C. app. 2) and 21 CFR part 14, relating to advisory committees. Dated: May 13, 2003.

Peter J. Pitts,

Associate Commissioner for External Belations.

[FR Doc. 03–12544 Filed 5–19–03; 8:45 am] **BILLING CODE 4160–01–S**

DEPARTMENT OF HEALTH AND HUMAN SERVICES

Food and Drug Administration

Obstetrics and Gynecology Devices Panel of the Medical Devices 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). At least one portion of the meeting will be closed to the public.

Name of the 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's regulatory issues.

Date and Time: The meeting will be held on June 9, 2003, from 12:30 p.m.