

North Pearl Street, Dallas, Texas 75201–2272:

1. *Integrity Bancshares, Inc.*, Houston Texas; to become a bank holding company by acquiring 100 percent of Integrity Bank, SSB, Houston, Texas.

Board of Governors of the Federal Reserve System.

Dated: July 6, 2011.

Robert deV. Frierson,

Deputy Secretary of the Board.

[FR Doc. 2011–17283 Filed 7–8–11; 8:45 am]

BILLING CODE 6210–01–P

FEDERAL RESERVE SYSTEM

Government in the Sunshine; Meeting Notice

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

TIME AND DATE: 10 a.m. on July 7, 2011.

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 Matters.

FOR MORE INFORMATION PLEASE CONTACT: Michelle Smith, Director, or Dave Skidmore, Assistant to the Board, Office of Board Members at 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: July 7, 2011.

Robert deV. Frierson,

Deputy Secretary of the Board.

[FR Doc. 2011–17487 Filed 7–7–11; 4:15 pm]

BILLING CODE 6210–01–P

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.

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 July 26, 2011.

A. Federal Reserve Bank of Richmond (Adam M. Drimer, Assistant Vice President) 701 East Byrd Street, Richmond, Virginia 23261–4528:

1. ASB Bancorp, Inc., Asheville, North Carolina; to engage *de novo* in extending credit and servicing loans activities, pursuant to section 225.28(b)(1) of Regulation Y.

Board of Governors of the Federal Reserve System, July 6, 2011.

Robert deV. Frierson,

Deputy Secretary of the Board.

[FR Doc. 2011–17281 Filed 7–8–11; 8:45 am]

BILLING CODE 6210–01–P

FEDERAL TRADE COMMISSION

Agency Information Collection Activities; Proposed Collection; Comment Request

AGENCY: Federal Trade Commission.

ACTION: Notice.

SUMMARY: The information collection requirements described below will be submitted to the Office of Management and Budget (OMB) for review, as

required by the Paperwork Reduction Act (PRA). The Federal Trade Commission (FTC) is seeking public comments on its proposal to extend through October 31, 2014, the current PRA clearance for information collection requirements contained in its Trade Regulation Rule entitled Power Output Claims for Amplifiers Utilized in Home Entertainment Products (Amplifier Rule or Rule), 16 CFR Part 432 (OMB Control Number 3084–0105). That clearance expires on October 31, 2011.

DATES: Comments must be filed by September 9, 2011.

ADDRESSES: Interested parties may file a comment online or on paper, by following the instructions in the Request for Comment part of the **SUPPLEMENTARY INFORMATION** section below. Write “Amplifier Rule: FTC File No. P974222” on your comment, and file your comment online at <https://ftcpublic.commentworks.com/ftc/amplifierrulepra>, by following the instructions on the web-based form. If you prefer to file your comment on paper, mail or deliver your comment to the following address: Federal Trade Commission, Office of the Secretary, Room H–113 (Annex J), 600 Pennsylvania Avenue, NW., Washington, DC 20580.

FOR FURTHER INFORMATION CONTACT: Requests for additional information should be addressed to Jock K. Chung, Attorney, Division of Enforcement, Bureau of Consumer Protection, Federal Trade Commission, M–8133, 600 Pennsylvania Avenue, NW., Washington, DC 20580, (202) 326–2984.

SUPPLEMENTARY INFORMATION:

Proposed Information Collection Activities

Under the PRA, 44 U.S.C. 3501–3521, federal agencies must obtain approval from OMB for each collection of information they conduct or sponsor. “Collection of information” means agency requests or requirements that members of the public submit reports, keep records, or provide information to a third party. 44 U.S.C. 3502(3), 5 CFR 1320.3(c). Because the number of entities affected by the Commission's requests will exceed ten, the Commission plans to seek OMB clearance under the PRA. As required by § 3506(c)(2)(A) of the PRA, the Commission is providing this opportunity for public comment before requesting that OMB extend the existing paperwork clearance for the information collection requirements associated with the Commission's Amplifier Rule.

The Amplifier Rule assists consumers by standardizing the measurement and disclosure of power output and other performance characteristics of amplifiers in stereos and other home entertainment equipment. The Rule also specifies the test conditions necessary to make the disclosures that the Rule requires.

Request for Comments

The FTC invites comments on: (1) Whether the proposed collection of information is necessary for the proper performance of the functions of the agency, including whether the information will have practical utility; (2) the accuracy of the agency'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 those who are to respond, including through the use of appropriate automated, electronic, mechanical, or other technological collection techniques or other forms of information technology, e.g., permitting electronic submission of responses. All comments should be filed as prescribed in the **ADDRESSES** section above, and must be received on or before September 9, 2011.

You can file a comment online or on paper. For the Commission to consider your comment, we must receive it on or before September 9, 2011. Write "Amplifier Rule: FTC File No. P974222" on your comment. Your comment—including your name and your state—will be placed on the public record of this proceeding, including, to the extent practicable, on the public Commission Web site, at <http://www.ftc.gov/ios/publiccomments.shtml>. As a matter of discretion, the Commission tries to remove individuals' home contact information from comments before placing them on the Commission Web site.

Because your comment will be made public, you are solely responsible for making sure that your comment doesn't include any sensitive personal information, like anyone's Social Security number, date of birth, driver's license number or other state identification number or foreign country equivalent, passport number, financial account number, or credit or debit card number. You are also solely responsible for making sure that your comment doesn't include any sensitive health information, like medical records or other individually identifiable health information. In addition, don't include

any "[t]rade secret or any commercial or financial information which is obtained from any person and which is privileged or confidential," as provided in Section 6(f) of the FTC Act, 15 U.S.C. 46(f), and FTC Rule 4.10(a)(2), 16 CFR 4.10(a)(2). In particular, don't include competitively sensitive information such as costs, sales statistics, inventories, formulas, patterns, devices, manufacturing processes, or customer names.

If you want the Commission to give your comment confidential treatment, you must file it in paper form, with a request for confidential treatment, and you have to follow the procedure explained in FTC Rule 4.9(c), 16 CFR 4.9(c).¹ Your comment will be kept confidential only if the FTC General Counsel, in his or her sole discretion, grants your request in accordance with the law and the public interest.

Postal mail addressed to the Commission is subject to delay due to heightened security screening. As a result, we encourage you to submit your comments online. To make sure that the Commission considers your online comment, you must file it at <https://ftcpublic.commentworks.com/ftc/amplifierrulepra>, by following the instructions on the web-based form. If this Notice appears at <http://www.regulations.gov>, you also may file a comment through that Web site.

If you file your comment on paper, write "Amplifier Rule: FTC File No. P974222" on your comment and on the envelope, and mail or deliver it to the following address: Federal Trade Commission, Office of the Secretary, Room H-113 (Annex J), 600 Pennsylvania Avenue, NW., Washington, DC 20580. If possible, submit your paper comment to the Commission by courier or overnight service.

Visit the Commission Web site at <http://www.ftc.gov> to read this Notice and the news release describing it. The FTC Act and other laws that the Commission administers permit the collection of public comments to consider and use in this proceeding as appropriate. The Commission will consider all timely and responsive public comments that it receives on or before September 9, 2011. You can find more information, including routine uses permitted by the Privacy Act, in the Commission's privacy policy, at <http://www.ftc.gov/ftc/privacy.htm>.

¹ In particular, the written request for confidential treatment that accompanies the comment must include the factual and legal basis for the request, and must identify the specific portions of the comment to be withheld from the public record. See FTC Rule 4.9(c), 16 CFR 4.9(c).

Amplifier Rule Burden Statement

Estimated annual hours burden: 450 hours (300 testing-related hours; 150 disclosure-related hours).

The Rule's provisions require affected entities to test the power output of amplifiers in accordance with a specified FTC protocol. The Commission staff estimates that approximately 300 new amplifiers and receivers come on the market each year. High fidelity manufacturers routinely conduct performance tests on these new products prior to sale. Because manufacturers conduct such tests, the Rule imposes no additional costs except to the extent that the FTC protocol is more time-consuming than alternative testing procedures. In this regard, a warm-up period that the Rule requires before measurements are taken may add approximately one hour to the time testing would otherwise entail. Thus, staff estimates that the Rule imposes approximately 300 hours (1 hour × 300 new products) of added testing burden annually.

In addition, the Rule requires disclosures if a manufacturer makes a power output claim for a covered product in an advertisement, specification sheet, or product brochure. This requirement does not impose any additional costs on manufacturers because, absent the Rule, media advertisements, as well as manufacturer specification sheets and product brochures, would contain a power specification obtained using an alternative to the Rule-required testing protocol. The Rule, however, also requires disclosure of harmonic distortion, power bandwidth, and impedance ratings in manufacturer specification sheets and product brochures that might not otherwise be included.

Staff assumes that manufacturers produce one specification sheet and one brochure each year for each new amplifier and receiver. The burden of disclosing the harmonic distortion, bandwidth, and impedance information on the specification sheets and brochures is limited to the time needed to draft and review the language pertaining to the aforementioned specifications. Staff estimates the time involved for this task to be a maximum of fifteen minutes for each new specification sheet and brochure for a total of 150 hours [(300 new products × 1 specification sheet) + (300 new products × 1 brochure)] × 15 minutes).

The total annual burden imposed by the Rule, therefore, is approximately 450 burden hours for testing and disclosures.

Estimated annual cost burden:
\$18,300.²

Generally, electronics engineers perform the testing of amplifiers and receivers. Staff estimates a labor cost of \$12,900 for such testing (300 hours for testing × \$43 mean hourly wages). Staff assumes advertising or promotions managers prepare the disclosures contained in product brochures and manufacturer specification sheet and estimates a labor cost of \$5,400 (150 hours for disclosures × \$36 mean hourly wages). Accordingly, staff estimates the total labor costs associated with the Rule to be approximately \$18,300 per year (\$12,900 for testing + \$5,400 for disclosures).

The Rule imposes no capital or other non-labor costs because its requirements are incidental to testing and advertising done in the ordinary course of business.

Willard K. Tom,

Willard K. Tom, General Counsel.

[FR Doc. 2011–17300 Filed 7–8–11; 8:45 am]

BILLING CODE 6750–01–M

DEPARTMENT OF HEALTH AND HUMAN SERVICES

Agency for Healthcare Research and Quality

Meeting of the National Advisory Council for Healthcare Research and Quality

AGENCY: Agency for Healthcare Research and Quality (AHRQ), HHS.

ACTION: Notice of public meeting.

SUMMARY: In accordance with section 10(a) of the Federal Advisory Committee Act, 5 U.S.C. App. 2, this notice announces a meeting of the National Advisory Council for Healthcare Research and Quality.

DATES: The meeting will be held on Friday, July 22 2011, from 8:30 a.m. to 3 p.m.

ADDRESSES: The meeting will be held at the Eisenberg Conference Center, Agency for Healthcare Research and Quality, 540 Gaither Road, Rockville, Maryland 20850.

FOR FURTHER INFORMATION CONTACT: Jaime Zimmerman, Coordinator of the Advisory Council, at the Agency for Healthcare Research and Quality, 540 Gaither Road, Rockville, Maryland, 20850, (301) 427–1456. For press-related information, please contact Karen Migdail at (301) 427–1855.

² Staff's labor cost estimates are based on recent data from the Bureau of Labor and Statistics found here: <http://www.bls.gov/news.release/ocs/spinctb1477.pdf>.

If sign language interpretation or other reasonable accommodation for a disability is needed, please contact the Food and Drug Administration (FDA) Office of Equal Employment Opportunity and Diversity Management on (301) 827–4840, no later than July 15, 2011. The agenda, roster, and minutes are available from Ms. Bonnie Campbell, Committee Management Officer, Agency for Healthcare Research and Quality, 540 Gaither Road, Rockville, Maryland, 20850. Ms. Campbell's phone number is (301) 427–1554.

SUPPLEMENTARY INFORMATION:

I. Purpose

The National Advisory Council for Healthcare Research and Quality is authorized by Section 941 of the Public Health Service Act, 42 U.S.C. 299c. In accordance with its statutory mandate, the Council is to advise the Secretary of the Department of Health and Human Services and the Director, Agency for Healthcare Research and Quality (AHRQ), on matters related to AHRQ's conduct of its mission including providing guidance on (A) priorities for health care research, (B) the field of health care research including training needs and information dissemination on health care quality and (C) the role of the Agency in light of private sector activity and opportunities for public private partnerships.

The Council is composed of members of the public, appointed by the Secretary, and Federal ex-officio members specified in the authorizing legislation.

II. Agenda

On Friday, July 22, there will be a subcommittee meeting for the National Healthcare Quality and Disparities Report scheduled to begin at 7:30 a.m. The Council meeting will convene at 8:30 a.m., with the call to order by the Council Chair and approval of previous Council summary notes. The AHRQ Director will present her update on current research, programs, and initiatives. The final agenda will be available on the AHRQ Web site at <http://www.ahrq.gov> no later than July 18, 2011.

This notice is published less than 15 days in advance of the meeting date due to logistical difficulties.

June 5, 2011.

Carolyn M. Clancy,
Director.

[FR Doc. 2011–17310 Filed 7–8–11; 8:45 am]

BILLING CODE 4160–90–M

DEPARTMENT OF HEALTH AND HUMAN SERVICES

Centers for Disease Control and Prevention

National Institute for Occupational Safety and Health, (NIOSH), World Trade Center Health Program Science/Technical Advisory Committee (WTCHP–STAC)

Correction: This notice was published in the **Federal Register** on June 23, 2011, Volume 76, Number 121, Page 36926–36927. The notice for the aforementioned solicitation has been changed to extend the deadline for receiving nominations. Nominations should be submitted (postmarked or received) no later than 5 p.m. EST July 29, 2011.

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 the Centers for Disease Control and Prevention, and the Agency for Toxic Substances and Disease Registry.

Dated: July 1, 2011.

Elizabeth A. Millington,

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

[FR Doc. 2011–17302 Filed 7–8–11; 8:45 am]

BILLING CODE 4163–18–P

DEPARTMENT OF HEALTH AND HUMAN SERVICES

Administration for Children and Families

Submission for OMB Review; Comment Request

Title: Low Income Home Energy Assistance Program (LIHEAP) Household Report.

OMB No. 0970–0060.

Description: This report is an annual activity required by statute (42 U.S.C. 8629) and Federal regulations (45 C.F.R. 96.92) for the Low Income Home Energy Assistance Program (LIHEAP). Submission of the completed report is one requirement for LIHEAP grantees applying for Federal LIHEAP block grant funds. States, the District of Columbia, and the Commonwealth of Puerto Rico are required to report statistics for the previous Federal fiscal year on:

- Assisted and applicant households, by type of LIHEAP assistance;