FEDERAL ACCOUNTING STANDARDS ADVISORY BOARD

Notice of Request for Comments on Proposed Deferred Maintenance and Repairs Standards

AGENCY: Federal Accounting Standards Advisory Board.

ACTION: Notice.

Board Action: Pursuant to 31 U.S.C. 3511(d), the Federal Advisory Committee Act (Pub. L. 92–463), as amended, and the FASAB Rules of Procedure, as amended in October, 2010, notice is hereby given that the Federal Accounting Standards Advisory Board (FASAB) is requesting comments on the Exposure Draft, *Deferred Maintenance and Repairs, Amending Statements of Federal Financial Accounting Standards 6, 14, 29 and 32.*

The Exposure Draft is available on the FASAB home page *http://www.fasab.gov/board-activities/documents-for-comment/exposure-drafts-and-documents-for-comment/*. Copies can be obtained by contacting FASAB at (202) 512–7350. Respondents are encouraged to comment on any part of the exposure draft. Written comments on the Exposure Draft are requested by September 16, 2011. Comments on the Exposure Drafts should be sent to: Wendy M. Payne, Executive Director, Federal Accounting Standards Advisory Board, 441 G Street, NW., Suite 6814, Mail Stop 6K17V, Washington, DC 20548.

FOR FURTHER INFORMATION CONTACT: Wendy Payne, Executive Director, at (202) 512–7350.

Authority: Federal Advisory Committee Act, Pub. L. 92–463.

Dated: June 29, 2011.

Charles Jackson,

Federal Register Liaison Officer.

[FR Doc. 2011–16796 Filed 7–1–11; 8:45 am]

BILLING CODE 1610–02–P

FEDERAL DEPOSIT INSURANCE CORPORATION

Sunshine Act Meeting

Pursuant to the provisions of the ''Government in the Sunshine Act'' (5 U.S.C. 552b), notice is hereby given that the Federal Deposit Insurance Corporation's Board of Directors will meet in open session at 10 a.m. on Wednesday, July 6, 2011, to consider the following matters:

Summary Agenda: No substantive discussion of the following items is anticipated. These matters will be resolved with a single vote unless a member of the Board of Directors requests that an item be moved to the discussion agenda.

Disposition of minutes of previous Board of Directors' Meetings.

Memorandum and resolution re: Final Rule Pursuant to § 742(c) of the Dodd-Frank Wall Street Reform and Consumer Protection Act for the Purpose of Adding 12 CFR part 349 to Regulate FDIC-Supervised Entities Engaged in Retail Forex Transactions.

Memorandum and resolution re: Final Rule on Part 329 & 330 For Interest on Deposits and Deposit Insurance Coverage.

Memorandum and resolution re: Notice of Proposed Rulemaking Regarding Calculating the Maximum Obligation the FDIC May Incur in Liquidating a Covered Financial Company.

Personnel Matters.

Discussion Agenda:

Orderly Liquidation Authority and Priorities and Claims Process under Orderly Liquidation Authority.

Resolution Plans and Credit Exposure Reports.

Special Reporting, Analysis and Contingent Resolution Plans at Certain Insured Depository Institutions.

The meeting will be held in the Board Room on the sixth floor of the FDIC Building located at 550 17th Street, NW., Washington, DC.

This Board meeting will be Webcast live via the Internet and subsequently made available on-demand approximately one week after the event. Visit *http://www.vodium.com/goto/fdic/boardmeetings.asp* to view the event. If you need any technical assistance, please visit our Video Help page at: *http://www.fdic.gov/video.html.*

The FDIC will provide attendees with auxiliary aids (e.g., sign language interpretation) required for this meeting. Those attendees needing such assistance should call 703–562–2404 (Voice) or 703–649–4354 (Video Phone) to make necessary arrangements.

Requests for further information concerning the meeting may be directed to Mr. Robert E. Feldman, Executive Secretary of the Corporation, at 202–898–7043.

Dated: June 29, 2011.

Federal Deposit Insurance Corporation.

Robert E. Feldman,

Executive Secretary.

[FR Doc. 2011–16829 Filed 6–30–11; 11:15 am]

BILLING CODE 6714–01–P

FEDERAL RESERVE SYSTEM

[Change in Bank Control Notices; Acquisitions of Shares of a Bank or Bank Holding Company]

The notificants listed below have 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 shares of a bank or bank holding company. The factors that are considered in acting on the 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. The notices also will 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 July 19, 2011.

A. Federal Reserve Bank of Minneapolis (Jacqueline G. King, Community Affairs Officer) 90 Hennepin Avenue, Minneapolis, Minnesota 55480–0291:

1. *Robert Lynn Nelson,* Cudjoe Key, Florida; to gain control of Kirkwood Bancorporation Co., and thereby indirectly acquire voting shares of Kirkwood Bank and Trust Company, both in Bismarck, North Dakota. In addition, Applicant has also applied to acquire voting shares of Kirkwood Bancorporation of Nevada, Inc., and thereby indirectly acquire voting shares of Kirkwood Bank of Nevada, both in Las Vegas, Nevada.

Board of Governors of the Federal Reserve System, June 29, 2011.

Robert deV. Frierson,

Deputy Secretary of the Board.

[FR Doc. 2011–16726 Filed 7–1–11; 8:45 am]

BILLING CODE 6210–01–P

FEDERAL TRADE COMMISSION

[File No. 092–3194]

Beiersdorf, Inc.; Analysis of Proposed Consent Order To Aid Public Comment

AGENCY: Federal Trade Commission.

ACTION: Proposed consent agreement.

SUMMARY: The consent agreement in this matter settles alleged violations of federal law prohibiting unfair or deceptive acts or practices or unfair methods of competition. The attached Analysis to Aid Public Comment describes both the allegations in the draft complaint and the terms of the

consent order—embodied in the consent agreement—that would settle these allegations.

DATES: Comments must be received on or before July 29, 2011.

ADDRESSES: Interested parties are invited to submit written comments electronically or in paper form. Comments should refer to ''Beiersdorf, File No. 092–3194'' to facilitate the organization of comments. Please note that your comment—including your name and your state—will be placed on the public record of this proceeding, including on the publicly accessible FTC Web site, at *http://www.ftc.gov/os/publiccomments.shtm.*

Because comments will be made public, they should not include any sensitive personal information, such as an individual'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. Comments also should not include any sensitive health information, such as medical records or other individually identifiable health information. In addition, comments should not 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 Commission Rule 4.10(a)(2), 16 CFR 4.10(a)(2). Comments containing material for which confidential treatment is requested must be filed in paper form, must be clearly labeled ''Confidential,'' and must comply with FTC Rule 4.9(c), 16 CFR 4.9(c).[1]

Because paper mail addressed to the FTC is subject to delay due to heightened security screening, please consider submitting your comments in electronic form. Comments filed in electronic form should be submitted by using the following Web link: *https://ftcpublic.commentworks.com/ftc/beiersdorfconsent* and following the instructions on the Web-based form. To ensure that the Commission considers an electronic comment, you must file it on the Web-based form at the Web link *https://ftcpublic.commentworks.com/ftc/beiersdorfconsent.* If this Notice appears at *http://www.regulations.gov/search/index.jsp,* you may also file an electronic comment through that Web site. The Commission will consider all comments that regulations.gov forwards to it. You may also visit the FTC Web site at *http://www.ftc.gov/* to read the Notice and the news release describing it.

A comment filed in paper form should include the ''Beiersdorf, File No. 092–3194'' reference both in the text and on the envelope, and should be mailed or delivered to the following address: Federal Trade Commission, Office of the Secretary, Room H–113 (Annex D), 600 Pennsylvania Avenue, NW., Washington, DC 20580. The FTC is requesting that any comment filed in paper form be sent by courier or overnight service, if possible, because U.S. postal mail in the Washington area and at the Commission is subject to delay due to heightened security precautions.

The Federal Trade Commission Act (''FTC Act'') and other laws 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, whether filed in paper or electronic form. Comments received will be available to the public on the FTC Web site, to the extent practicable, at *http://www.ftc.gov/os/publiccomments.shtm.* As a matter of discretion, the Commission makes every effort to remove home contact information for individuals from the public comments it receives before placing those comments on the FTC Web site. More information, including routine uses permitted by the Privacy Act, may be found in the FTC's privacy policy, at *http://www.ftc.gov/ftc/privacy.shtm.*

FOR FURTHER INFORMATION CONTACT: Evan Rose (415–848–5100), FTC, Western Region, San Francisco, 600 Pennsylvania Avenue, NW., Washington, DC 20580.

SUPPLEMENTARY INFORMATION: Pursuant to section 6(f) of the Federal Trade Commission Act, 38 Stat. 721, 15 U.S.C. 46(f), and 2.34 the Commission Rules of Practice, 16 CFR 2.34, notice is hereby given that the above-captioned consent agreement containing a consent order to cease and desist, having been filed with and accepted, subject to final approval, by the Commission, has been placed on the public record for a period of thirty (30) days. The following Analysis to Aid Public Comment describes the terms of the consent agreement, and the allegations in the complaint. An electronic copy of the full text of the consent agreement package can be obtained from the FTC Home Page (for June 29, 2011), on the World Wide Web, at *http://www.ftc.gov/os/actions.shtm.* A paper copy can be obtained from the FTC Public Reference Room, Room 130–H, 600 Pennsylvania Avenue, NW., Washington, DC 20580, either in person or by calling (202) 326–2222.

Public comments are invited, and may be filed with the Commission in either paper or electronic form. All comments should be filed as prescribed in the **ADDRESSES** section above, and must be received on or before the date specified in the **DATES** section.

Analysis of Agreement Containing Consent Order To Aid Public Comment

The Federal Trade Commission (''FTC'' or ''Commission'') has accepted, subject to final approval, an Agreement Containing Consent Order from Beiersdorf, Inc. (''respondent''). The proposed consent order has been placed on the public record for thirty (30) days for receipt of comments by interested persons. Comments received during this period will become part of the public record. After thirty (30) days, the Commission will again review the agreement and the comments received, and will decide whether it should withdraw from the agreement and take appropriate action or make final the agreement's proposed order.

This matter involves the advertising, marketing, and sale of ''NIVEA My Silhouette! Redefining Gel-Cream'' (''My Silhouette'') by respondent. Respondent has marketed My Silhouette to consumers through third-party retail outlets.

My Silhouette is a skin cream that contains ''Bio-slim Complex,'' a combination of ingredients that includes white tea and anise. According to the FTC complaint, respondent promoted My Silhouette as able to slim and reshape the body.

Specifically, the FTC complaint alleges that respondent represented, in various advertisements, that regular use of My Silhouette results in significant reductions in body size. The complaint alleges that this claim is false and thus violates the FTC Act.

The proposed consent order contains provisions designed to prevent respondent from engaging in similar acts or practices in the future. Specifically, Part I prohibits respondent from claiming that My Silhouette or any other topically applied product causes substantial weight or fat loss or a substantial reduction in body size.

Part II covers any representation that a drug, dietary supplement, or cosmetic causes weight or fat loss or a reduction in body size. Part II prohibits

[1] The comment must be accompanied by an explicit request for confidential treatment, including the factual and legal basis for the request, and must identify the specific portions of the comment to be withheld from the public record. The request will be granted or denied by the Commission's General Counsel, consistent with applicable law and the public interest. *See* FTC Rule 4.9(c), 16 CFR 4.9(c).

respondent from making such representations unless the representation is non-misleading, and, at the time of making such representation, respondent possesses and relies upon competent and reliable scientific evidence that substantiates that the representation is true. For purposes of Part II, the proposed order defines ''competent and reliable scientific evidence'' as at least two randomized, double-blind, placebo-controlled human clinical studies that are conducted by independent, qualified researchers and that conform to acceptable designs and protocols, and whose results, when considered in light of the entire body of relevant and reliable scientific evidence, are sufficient to substantiate that the representation is true.

Part III of the proposed order prohibits respondent from making representations, other than representations covered under Parts I or II, about the health benefits of any drug, dietary supplement, or cosmetic, unless the representation is non-misleading, and, at the time of making such representation, respondent possesses and relies upon competent and reliable scientific evidence that is sufficient in quality and quantity based on standards generally accepted in the relevant scientific fields, when considered in light of the entire body of relevant and reliable scientific evidence, to substantiate that the representation is true. For purposes of Part III, the proposed order defines ''competent and reliable scientific evidence'' as ''tests, analyses, research, or studies that have been conducted and evaluated in an objective manner by qualified persons, and that are generally accepted in the profession to yield accurate and reliable results.''

Part IV of the proposed order states that the order does not prohibit respondent from making representations for any drug that are permitted in labeling for that drug under any tentative or final standard promulgated by the Food and Drug Administration (''FDA''), or under any new drug application approved by the FDA. This part of the proposed order also states that the order does not prohibit respondent from making representations for any product that are specifically permitted in labeling for that product by regulations issued by the FDA under the Nutrition Labeling and Education Act of 1990.

Part V of the proposed order requires respondent to pay nine hundred thousand dollars ($900,000) to the Commission to be used for equitable relief, including restitution, and any attendant expenses for the administration of such equitable relief.

Parts VI, VII, VIII, and IX of the proposed order require respondent to keep copies of relevant advertisements and materials substantiating claims made in the advertisements; to provide copies of the order to its personnel; to notify the Commission of changes in corporate structure that might affect compliance obligations under the order; and to file compliance reports with the Commission. Part X provides that the order will terminate after twenty (20) years, with certain exceptions.

The purpose of this analysis is to facilitate public comment on the proposed order, and it is not intended to constitute an official interpretation of the agreement and proposed order or to modify their terms in any way.

By direction of the Commission.

Donald S. Clark,

Secretary.

[FR Doc. 2011–16739 Filed 7–1–11; 8:45 am]

BILLING CODE 6750–01–P

DEPARTMENT OF HEALTH AND HUMAN SERVICES

Meeting of the Advisory Committee on Minority Health

AGENCY: Office of Minority Health, Office of the Assistant Secretary for Health, Office of the Secretary, Department of Health and Human Services.

ACTION: Notice of meeting.

SUMMARY: As stipulated by the Federal Advisory Committee Act, the Department of Health and Human Services (DHHS) is hereby giving notice that the Advisory Committee on Minority Health (ACMH) will hold a meeting. This meeting is open to the public. Preregistration is required for both public attendance and comment. Any individual who wishes to attend the meeting and/or participate in the public comment session should e-mail *acmh@osophs.dhhs.gov.*

DATES: The meeting will be held on Monday, August 29, 2011 from 9 a.m. to 5 p.m. and Tuesday, August 30, 2011 from 9 a.m. to 1 p.m.

ADDRESSES: The meeting will be held at the Doubletree Hotel, 1515 Rhode Island Ave., NW., Washington, DC 20005.

FOR FURTHER INFORMATION CONTACT: Ms. Monica A. Baltimore, Tower Building, 1101 Wootton Parkway, Suite 600, Rockville, Maryland 20852. Phone: 240–453–2882 Fax: 240–453–2883.

SUPPLEMENTARY INFORMATION: In accordance with Public Law 105–392, the ACMH was established to provide advice to the Deputy Assistant Secretary for Minority Health in improving the health of each racial and ethnic minority group and on the development of goals and specific program activities of the Office of Minority Health.

Topics to be discussed during this meeting will include increasing the health care workforce and strategies to improve the health of racial and ethnic minority populations through the development of health policies and programs that will help eliminate health disparities, as well as other related issues.

Public attendance at the meeting is limited to space available. Individuals who plan to attend and need special assistance, such as sign language interpretation or other reasonable accommodations, should notify the designated contact person at least fourteen (14) business days prior to the meeting. Members of the public will have an opportunity to provide comments at the meeting. Public comments will be limited to three minutes per speaker. Individuals who would like to submit written statements should mail or fax their comments to the Office of Minority Health at least seven (7) business days prior to the meeting. Any members of the public who wish to have printed material distributed to ACMH committee members should submit their materials to the Executive Secretary, ACMH, Tower Building, 1101 Wootton Parkway, Suite 600, Rockville, Maryland 20852, prior to close of business August 24, 2011.

Dated: June 20, 2011.

Garth N. Graham,

Deputy Assistant Secretary for Minority Health, Office of Minority Health, Office of the Assistant Secretary for Health, Office of the Secretary, U.S. Department of Health and Human Services.

[FR Doc. 2011–16744 Filed 7–1–11; 8:45 am]

BILLING CODE 4150–29–P

DEPARTMENT OF HEALTH AND HUMAN SERVICES

HIT Standards Committee's Workgroup Meetings; Notice of Meetings

AGENCY: Office of the National Coordinator for Health Information Technology, HHS.

ACTION: Notice of meetings.

This notice announces forthcoming subcommittee meetings of a Federal advisory committee of the Office of the National Coordinator for Health Information Technology (ONC). The