418–0530 (voice), (202) 418–0432 (TTY). The complete text of the *Order* may also be purchased from the Commission's duplicating contractor, Best Company and Printing, Inc., Portals II, 445 12th Street, SW., Room CY–B402, Washington, DC 20554, telephone (202) 863–2893, facsimile (202) 863–2898, or by e-mail *fcc@bcpiweb.com*, or via its Web site *http://www.bcpiweb.com*.

Synopsis of Order

- 1. On January 16, 2009, the Commission released its 13th Annual Report to Congress on the status of competition in the market for the delivery of video programming. See Annual Assessment of the Status of Competition in the Market for the Delivery of Video Programming, 13th Annual Report, 24 FCC Rcd 542 (2009). The Report provided information as of 2006. On the same date, the Commission released a Notice of Inquiry (NOI) soliciting 2007 data for the 14th Annual Report. See Annual Assessment of the Status of Competition in the Market for the Delivery of Video Programming, Notice of Inquiry, 24 FCC Rcd 750 (2009), 74 FR 8675, February 11, 2009. The NOI initially set a deadline of February 27, 2009 for comments, and March 28, 2009 for reply comments.
- 2. On February 23, 2009, the Commission issued an extension of time to file comments and reply comments for 2007 data. See Annual Assessment of the Status of Competition in the Market for the Delivery of Video Programming, Order, 24 FCC Rcd 2524 (2009), 74 FR 11102, March 16, 2009. As described in the extension, comments in response to the NOI seeking information as of June 30, 2007, are due on April 28, 2009, and reply comments are due on May 28, 2009. We noted that the establishment of a single set of filing deadlines for 2007 and 2008 data will avoid duplication of effort by commenters and will streamline Commission review of the submitted information, thereby conserving commenters' and Commission resources.
- 3. As described in the Supplemental Notice of Inquiry that the Commission adopted on April 8, 2009, we will release a single report covering the years 2007, 2008, and 2009. Comments for 2008 data are due May 20, 2009, and reply comments are due June 20, 2009.
- 4. To afford commenters the opportunity to file their 2007 and 2008 information concurrently, we are hereby extending the comment periods announced in the Order to coincide with the filing deadlines we establish for the 2008 data.

5. Accordingly, it is ordered that, pursuant to sections 4(i), 4(j), 403 and 628(g) of the Communications Act of 1934, as amended, 47 U.S.C. 154(i), 154(j), 403, and 548(g), and §§ 0.061, 0.204, 0.283, and 1.46 of the rules, 47 CFR 0.061, 0.204, 0.283, and 1.46, the deadlines for responding to the Notice of Inquiry for the 14th Annual Report are extended, sua sponte, until May 20, 2009, for initial comments and June 20, 2009, for reply comments.

Federal Communications Commission,

Marlene H. Dortch,

Secretary.

[FR Doc. E9–9571 Filed 4–24–09; 8:45 am]

FEDERAL RESERVE SYSTEM

Change in Bank Control Notices; Acquisition of Shares of Bank or Bank Holding Companies

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 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 office 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 May 12, 2009.

- A. Federal Reserve Bank of Atlanta (Steve Foley, Vice President) 1000 Peachtree Street, N.E., Atlanta, Georgia 30309
- 1. Jessie Doyle Buffinton; Helmut
 Hans Cawthon; Kenneth Franklin Davis;
 Andrew Charles Heaner; Clinton Gray
 Hubbard; Stephen Curtis Klasson;
 Kimberly Gill Mauer; Laura Hallin
 Mumber; Matthew Peter Mumber; Delos
 Harley Yancey, III; John Demetrius
 Xanthos, all of Rome, Georgia; Andrew
 Charles Heaner, Atlanta, Georgia; and
 Wayne Elm Vick, Armuchee, Georgia;
 collectively to acquire additional voting
 shares of Heritage First Bancshares, Inc.,
 and thereby indirectly acquire
 additional voting shares of Heritage
 First Bank, both of Rome, Georgia.
- B. Federal Reserve Bank of St. Louis (Glenda Wilson, Community Affairs Officer) P.O. Box 442, St. Louis, Missouri 63166–2034:

1. Terry L. Bunnell, individually and in concert with Deborah L. Bunnell, both of Glasgow, Kentucky; to acquire control of Peoples—Marion Bancorp, Inc., and thereby indirectly acquire control of The Peoples Bank, both of Marion, Kentucky.

Board of Governors of the Federal Reserve System, April 22, 2009.

Robert deV. Frierson,

Deputy Secretary of the Board.

[FR Doc. E9–9520 Filed 4–24–09; 8:45 am]

BILLING CODE 6210-01-S

FEDERAL TRADE COMMISSION

[File No. 082 3145]

Kellogg Company; 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 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 May 19, 2009.

ADDRESSES: Interested parties are invited to submit written comments electronically or in paper form.

Comments should refer to "Kellogg, File No. 082 3145" 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 website, 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).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 weblink: (https:// secure.commentworks.com/ftc-Kellogg) (and following the instructions on the web-based form). To ensure that the Commission considers an electronic comment, you must file it on the webbased form at the weblink: (https:// secure.commentworks.com/ftc-Kellogg). If this Notice appears at (http:// www.regulations.gov/search/index.jsp), you may also file an electronic comment through that website. The Commission will consider all comments that regulations.gov forwards to it. You may also visit the FTC website at (http:// www.ftc.gov) to read the Notice and the news release describing it.

A comment filed in paper form should include the "Kellogg, File No. 082 3145" 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-135, 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 website, 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 website. 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:

Heather Hippsley or Kial S. Young, Bureau of Consumer Protection, 600 Pennsylvania Avenue, NW, Washington, D.C. 20580, (202) 326–3285.

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 February 18, 2009), on the World Wide Web, at (http:// www.ftc.gov/os/2009/04/index.htm). A paper copy can be obtained from the FTC Public Reference Room, Room 130-H, 600 Pennsylvania Avenue, NW, Washington, D.C. 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 a consent order from Kellogg Company ("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 and promotion of Kellogg's Frosted Mini-Wheats, a well-known breakfast cereal. According to the FTC complaint, Respondent represented, in various advertisements, that eating a bowl of Kellogg's Frosted Mini-Wheats cereal for breakfast is clinically shown to improve kids' attentiveness by nearly 20%. The complaint alleges that this claim is false or misleading because, in fact, in the clinical study referred to in respondent's advertisements, only about half the kids who ate Frosted Mini-Wheats cereal showed any improvement after three hours as compared to their pre-breakfast baseline. In addition, overall, only one in seven kids who ate the cereal improved their attentiveness by 18% or more, and only about one in nine improved by 20% or more.

The FTC complaint also charges that Respondent represented, in other advertising, that eating a bowl of Kellogg's Frosted Mini-Wheats cereal for breakfast is clinically shown to improve kids' attentiveness by nearly 20% when compared to kids who ate no breakfast. The FTC alleges that this claim is also false or misleading, because in fact, kids in the clinical study who ate Frosted Mini-Wheats had an average of 10.6% better attentiveness three hours later than kids who had skipped breakfast. In addition, relatively few kids experienced better attentiveness near the 20% level.

The proposed consent order contains provisions designed to prevent Respondent from engaging in similar acts and practices in the future. Part I of the proposed order prohibits Respondent from representing that (a) eating a bowl of Kellogg's Frosted Mini-Wheats cereal for breakfast is clinically shown to improve kids' attentiveness by nearly 20%, or any other specific percentage; and (b) eating a bowl of Kellogg's Frosted Mini-Wheats cereal for breakfast is clinically shown to improve kids' attentiveness by nearly 20%, or any other specific percentage, compared to kids who ate no breakfast, unless the representation is true and nonmisleading at the time it is made.

Part II of the proposed order prohibits Respondent from making any representations in advertising for Frosted Mini-Wheats or any other morning food or snack food about the benefits, performance, or efficacy of the product for cognitive function, processes, or health, unless the representation is true and nonmisleading. In addition, Respondent must possess competent and reliable scientific evidence for such claims.

¹ FTC Rule 4.2(d), 16 CFR 4.2(d). 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).

Part III of the proposed order prohibits Respondent from making misrepresentations in advertising for any morning food or snack food about the existence, contents, validity, results, conclusions, or interpretations of any test, study or research.

Part IV of the proposed order 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 issues by the FDA under the Nutrition Labeling and Education Act of 1990.

Parts V through VIII 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 certain of their 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 IX 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 in any way their terms.

By direction of the Commission.

Donald S. Clark

Secretary.

[FR Doc. E9–9484 Filed 4–24–09: 8:45 am] BILLING CODE 6750–01–S

GENERAL SERVICES ADMINISTRATION

[OMB Control No. 3090-0044]

Public Buildings Service; Information Collection; GSA Form 3453, Application/Permit for Use of Space in Public Buildings and Grounds

AGENCY: Public Buildings Service, GSA. **ACTION:** Notice of request for comments regarding a renewal to an existing OMB clearance.

SUMMARY: Under the provisions of the Paperwork Reduction Act of 1995 (44 U.S.C. Chapter 35), the General Services Administration will be submitting to the Office of Management and Budget (OMB) a request to review and approve a renewal of a currently approved information collection requirement regarding GSA Form 3453, Application/Permit for Use of Space in Public Buildings and Grounds. The clearance currently expires on April 30, 2009.

Public comments are particularly invited on: Whether this collection of information is necessary and whether it will have practical utility; whether our estimate of the public burden of this collection of information is accurate, and based on valid assumptions and methodology; ways to enhance the quality, utility, and clarity of the information to be collected.

DATES: Submit comments on or before: June 26, 2009.

FOR FURTHER INFORMATION CONTACT:

Frank Giblin, Public Buildings Service, at telephone (202) 501–1856, or via e-mail to frank.giblin@gsa.gov.

ADDRESSES: Submit comments regarding this burden estimate or any other aspect of this collection of information, including suggestions for reducing this burden, to the Regulatory Secretariat (VPR), General Services Administration, Room 4041, 1800 F Street, NW., Washington, DC 20405. Please cite OMB Control No. 3090–0044, GSA Form 3453, Application/Permit for Use of Space in Public Buildings and Grounds, in all correspondence.

SUPPLEMENTARY INFORMATION:

A. Purpose

The general public uses GSA Form 3453, Application/Permit for Use of Space in Public Buildings and Grounds, to request the use of public space in Federal buildings and on Federal grounds for cultural, educational, or recreational activities. A copy, sample, or description of any material or item proposed for distribution or display must also accompany this request.

B. Annual Reporting Burden

Respondents: 8,000.

Responses per Respondent: 1.

Hours per Response: 0.05.

Total Burden Hours: 400.

Obtaining Copies of Proposals: Requesters may obtain a copy of the information collection documents from the General Services Administration, Regulatory Secretariat (VPR), 1800 F Street, NW., Room 4041, Washington, DC 20405, telephone (202) 501–4755. Please cite OMB Control No. 3090–0044, GSA Form 3453, Application/Permit for Use of Space in Public Buildings and Grounds, in all correspondence.

Dated: April 21, 2009.

Philip E. Klokis,

Acting Chief Information Officer.
[FR Doc. E9–9490 Filed 4–24–09; 8:45 am]
BILLING CODE 6820–YT–P

DEPARTMENT OF HEALTH AND HUMAN SERVICES

Food and Drug Administration [Docket No. FDA-2009-D-0179]

Draft Guidance for Industry and Food and Drug Administration Staff: Technical Considerations for Pen, Jet, and Related Injectors Intended for Use With Drugs and Biological Products; Availability

AGENCY: Food and Drug Administration,

HHS.

ACTION: Notice.

SUMMARY: The Food and Drug Administration (FDA) is announcing the availability of a draft guidance document entitled "Technical Considerations for Pen, Jet, and Related Injectors Intended for Use with Drugs and Biological Products." The draft guidance document provides technical and scientific information for sponsors to consider in developing information to support a marketing application for a pen, jet, or related injector device intended for use with drugs or biological products. The marketing application would typically be a premarket notification submission (510(k)) or a premarket approval (PMA) application for the injector alone. For a combination product that includes the injector, the marketing application would typically be a new drug application (NDA) or a biological licensing application (BLA).

DATES: Although you can comment on any guidance at any time (see 21 CFR 10.115(g)(5)), to ensure that the agency considers your comment on this draft guidance before it begins work on the final version of the guidance, submit written or electronic comments on the draft guidance by July 27, 2009.

ADDRESSES: Submit written requests for single copies of the draft guidance to the Office of Combination Products, 15800 Crabbs Branch Way, Rockville, MD 20855. Send one self-addressed adhesive label to assist the office in processing your requests. The draft guidance may also be obtained by mail by calling the Office of Combination Products at 301–427–1934 or by e-mail to combination@fda.gov. See the SUPPLEMENTARY INFORMATION section for

SUPPLEMENTARY INFORMATION section for electronic access to the draft guidance document.

Submit written comments on the draft guidance to the Division of Dockets Management (HFA–305), Food and Drug Administration, 5630 Fishers Lane, rm. 1061, Rockville, MD 20852. Submit electronic comments to http://www.regulations.gov.