20472, telephone (202) 646–2756 or by facsimile at (202) 646–4596.

SUPPLEMENTARY INFORMATION: This meeting is open to the public with limited seating available on a first-come, first-served basis. Members of the general public who plan to attend the meeting should contact Ms. Sally P. Magee, Federal Emergency Management Agency, 500 C Street SW., room 442, Washington, DC 20472, telephone (202) 646–8242 or by facsimile at (202) 646–4596 on or before September 6, 1999.

Minutes of the meeting will be prepared and will be available upon request 30 days after they have been approved by the next Technical Mapping Advisory Council meeting.

Dated: November 22, 1999.

Michael J. Armstrong,

Associate Director for Mitigation.

[FR Doc. 99–31093 Filed 11–29–99; 8:45 am]

BILLING CODE 6718–04–P

FEDERAL RESERVE SYSTEM

Sunshine Act Meeting

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

TIME AND DATE: 11:00 a.m., Monday, December 6, 1999.

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.

CONTACT PERSON FOR MORE INFORMATION: Lynn S. Fox, Assistant to the Board; 202–452–3204.

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: November 26, 1999.

Jennifer J. Johnson,

Secretary of the Board.

[FR Doc. 99–31193 Filed 11–26–99; 12:29 pm]

BILLING CODE 6210–01–P

FEDERAL TRADE COMMISSION

[File No. 982 3152]

Quigley Corporation; Analysis 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 of unfair methods of competition. The attached Analysis to Aid Public Comment describes both the allegations in the draft complaint that accompanies the consent agreement 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 January 31, 2000.

ADDRESSES: Comments should be directed to: FTC/Office of the Secretary, Room 159, 600 Pennsylvania Ave., NW, Washington, DC 20580.

FOR FURTHER INFORMATION CONTACT: Daniel Kaufman or Michelle Rusk, FTC/S–4002, 600 Pennsylvania Ave., NW, Washington, DC 20580. (202) 326–2888 or 326–3148.

SUPPLEMENTARY INFORMATION: Pursuant to section 6(f) of the Federal Trade Commission Act, 38 Stat. 721, 15 U.S.C. 46 and § 2.34 of the Commission's 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 sixty (60) 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 November 23, 1999), on the World Wide Web, at ''http://www.ftc.gov/os/actions97.htm.'' A paper copy can be obtained from the FTC Public Reference Room, Room H–130, 600 Pennsylvania Avenue, NW, Washington, DC 20580, either in person or by calling (202) 326–3627.

Public comment is invited. Comments should be directed to: FTC/Office of the Secretary, Room 159, 600 Pennsylvania Ave., NW, Washington, DC 20580. Two paper copies of each comment should be filed, and should be accompanied, if possible, by a 3½ inch diskette containing an electronic copy of the comment. Such comments or views will be considered by the Commission and will be available for inspection and copying at its principal office in accordance with Section 4.9(b)(6)(ii) of the Commission's Rules of Practice (16 CFR 4.9(b)(6)(ii)).

Analysis of Proposed Consent Order To Aid Public Comment

The Federal Trade Commission has accepted, subject to final approval, an agreement containing consent order from respondent the Quigley Corporation (''Quigley'').

The proposed consent order has been placed on the public record for sixty (60) days for reception of comments by interested persons. Comments received during this period will become part of the public record. After sixty (60) days, the Commission will again review the agreement and the comments received and will decide whether it should withdraw from the agreement or make final the agreement's proposed order.

This matter involves alleged deceptive representations for Cold-Eeze Zinc Lozenges and Cold-Eezer Plus Zinc Gluconate Lozenges (hereinafter, collectively ''Cold-Eeze'') and Kids-Eeze Bubble Gum (''Kids-Eeze'').

The Commission's proposed complaint alleges that Quigley made unsubstantiated representations that Cold-Eeze will prevent users from contracting colds and pneumonia; will treat allergies; will reduce the severity of colds in children; and that Kids-Eeze will reduce the severity of cold symptoms in children.

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 the respondent from making the representations about Cold-Eeze and Kids-Eeze challenged in the complaint, unless it possesses and relies upon competent and reliable scientific evidence that substantiates the representation.

Part II of the proposed order prohibits respondent from making any representation that any food, drug, or dietary supplement can or will cure, treat or prevent any disease, or have any effect on the structure or function of the human body, unless it possesses and relies upon competent and reliable scientific evidence that substantiates the representation.

Part III of the proposed order allows the respondent to make any representations for any drug that are permitted in labeling for the drug under any tentative final or final Food and Drug Administration (''FDA'') standard or under any new drug application approved by the FDA.

Part IV of the proposed order allows the respondent to make 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.

Parts V through VIII require the respondent to keep copies of advertisements making representations covered by the order; to keep records concerning those representations, including material that they relied upon when making the representations; to provide copies of the order to certain of the respondents' personnel; to notify the Commission of changes in corporate structure; and to file compliance reports with the Commission.

Part IX of the proposed order is a ''sunset'' provision, dictating that the order will terminate twenty years from the date it is issued or twenty years after a complaint is filed in federal court, but either the United States or the FTC, alleging any violation of the order.

The purpose of this analysis is to facilitate public comment on the proposed order. 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. 99–31055 Filed 11–29–99; 8:45 am]

BILLING CODE 6750–01–M

FEDERAL TRADE COMMISSION

[File No. 982 3152]

QVC, Inc.; Analysis 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 that accompanies the consent agreement 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 January 31, 2000.

ADDRESSES: Comments should be directed to: FTC/Office of the Secretary, Room 159, 600 Pennsylvania Ave., NW, Washington, DC 20580.

FOR FURTHER INFORMATION CONTACT: Daniel Kaufman or Michelle Rusk, FTC/S–4002, 600 Pennsylvania Ave., NW, Washington, DC 20580. (202) 326–2888 or 326–3148.

SUPPLEMENTARY INFORMATION: Pursuant to section 6(f) of the Federal Trade Commission Act, 38 Stat. 721, 15 U.S.C. 46 and § 2.34 of the Commission's 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 sixty (60) 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 November 23, 1999), on the World Wide Web, at ''http://www.ftc.gov/os/actions97.htm.'' A paper copy can be obtained from the FTC Public Reference Room, Room H–130, 600 Pennsylvania Avenue, NW, Washington, DC 20580, either in person or by calling (202) 326–3627.

Public comment is invited. Comments should be directed to: FTC/Office of the Secretary, Room 159, 600 Pennsylvania Ave., NW, Washington, DC 20580. Two paper copies of each comment should be filed, and should be accompanied, if possible, by a 3½ inch diskette containing an electronic copy of the comment. Such comments or views will be considered by the Commission and will be available for inspection and copying at its principal office in accordance with § 4.9(b)(6)(ii) of the Commission's rules or practice (16 CFR 4.9(b)(6)(ii).

Analysis of Proposed Consent Order To Aid Public Comment

The Federal Trade Commission has accepted, subject to final approval, an agreement containing consent order from respondent QVC, Inc. (''QVC'').

The proposed consent order has been placed on the public record for sixty (60) days for reception of comments by interested persons. Comments received during this period will become part of the public record. After sixty (60) days, the Commission will again review the agreement and the comments received and will decide whether it should withdraw from the agreement or make final the agreement's proposed order.

This matter involves alleged deceptive representations for Cold-Eeze Zinc Lozenges and Cold-Eezer Plus Zinc Gluconate Lozenges (hereinafter, collectively ''Cold-Eeze'').

The Commission's proposed complaint alleges that QVC made unsubstantiated representations that Cold-Eeze will prevent users from contracting colds and pneumonia; will treat allergies; and will reduce the severity of colds in children.

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 the respondent from making the representations about Cold-Eeze challenged in the complaint, unless it possesses and relies upon competent and reliable scientific evidence that substantiates the representation.

Part II of the proposed order prohibits the respondent from making any representation that any dietary supplement can or will cure, threat or prevent any disease, or have any effect on the structure or function of the human body, unless it possesses and relies upon competent and reliable scientific evidence that substantiates the representation.

Part III of the proposed order allows the respondent to make any representations for any drug that are permitted in labeling for the drug under any tentative final or final Food and Drug Administration (''FDA'') standard or under any new drug application approved by the FDA.

Part IV of the proposed order allows the respondent to make 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.

Parts V through VIII require the respondent to keep copies of advertisements making representations covered by the order; to keep records concerning those representations, including material that they relied upon when making the representations; to provide copies of the order to certain of the respondents' personnel; to notify the Commission of changes in corporate structure; and to file compliance reports with the Commission.

Part IX of the proposed order is a ''sunset'' provision, dictating that the order will terminate twenty years from the date it is issued or twenty years after a complaint is filed in federal court, by either the United States or the FTC, alleging any violation of the order.