Office, 150 William Street, Suite 1300, New York, NY 10038. (212) 264–1207.

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 accompanying complaint. An electronic copy of the full text of the consent agreement package can be obtained from the Commission Actions section of the FTC Home Page (for October 29, 1997), 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, Sixth Street and Pennsylvania Avenue, N.W., Washington, D.C. 20580, either in person or by calling (202) 326–3627. Public comment is invited. 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 to a proposed consent order from Venegas Inc. (''Venegas'') and Angel Venegas.

The proposed consent order has been placed on the public record for sixty (60) days for receipt 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 comments received and will decide whether it should withdraw from the agreement or make final the agreement's proposed order.

This matter concerns print advertisements for proposed respondents' Alen, a powdered nutritional supplement that contains wheat germ, wheat bran, soybean extract, and seaweed extract. The Commission's complaint alleges that the proposed respondents made unsubstantiated representations that Alen: increases life expectancy; delays the aging process; eliminates anemia; increases the immune system's defenses; increases memory or scholastic performance; helps diabetics naturally produce insulin; reduces the pain of rheumatism or migraines; lowers blood pressure; helps heal ulcers; increases muscle bulk; controls addictions to excess fat and sweets; and protects against infections and increases and enhances the healing process.

The proposed order contains provisions designed to remedy the violations charged and to prevent proposed respondents from engaging in similar acts in the future.

Paragraph I of the proposed order prohibits proposed respondents from representing that Alen or any other product: Increases life expectancy; delays the aging process; eliminates anemia; increases the immune system's defenses; increases memory or scholastic performance; helps diabetics naturally produce insulin; reduces the pain of rheumatism or migraines; lowers blood pressure; helps heal ulcers, increases muscle bulk; controls addictions to excess fat and sweets; or protects against infections and increases and enhances the healing process, unless at the time the representation is made, respondents possess and rely upon competent and reliable scientific evidence that substantiates the representation.

Paragraph II of the proposed order prohibits proposed respondents from making any representation about the benefits, performance, or efficacy of Alen, or any food, dietary supplement, or drug, unless, at the time the representation is made, proposed respondents possess and rely upon competent and reliable scientific evidence that substantiates the representation.

Paragraph III of the proposed order provides that nothing in this order shall prohibit proposed respondents from making any representation for any product permitted by the Food and Drug Administration. Paragraph IV of the proposed order provides that nothing in this order shall prohibit proposed respondent from making any representation for any drug permitted by the Food and Drug Administration.

Paragraph V of the proposed order requires the proposed respondents to keep and maintain all advertisements and promotional materials containing any representation, and all materials that were relied upon in disseminating the representations, covered by the proposed order. Additionally, Paragraph VI requires distribution of a copy of the consent order to current and future officers and agents. Further, Paragraph VII provides for Commission notification upon a change in the corporate respondent, and Paragraph VIII requires Commission notification when the individual respondent changes his present business or employment. Paragraph IX requires proposed respondents to file compliance reports with the Commission. Lastly, Paragraph X provides for the termination of the order after twenty (20) years under certain circumstances.

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.

Donald S. Clark,

Secretary.

[FR Doc. 97–29279 Filed 11–4–97; 8:45 am]

BILLING CODE 6750–01–M

GENERAL ACCOUNTING OFFICE

Federal Accounting Standards Advisory Board

AGENCY: General Accounting Office.

ACTION: Cancellation of November Meeting.

Cancellation

The previously announced meeting (**Federal Register** of October 30) on Friday, November 7, 1997, is hereby cancelled. Due notice will be given for the next meeting, to be held on December 19.

FOR FURTHER INFORMATION CONTACT: Wendy Comes, Executive Director, 441 G St., N.W., Room 3B18, Washington, D.C. 20548, or call (202) 512–7350.

Authority: Federal Advisory Committee Act. Pub. L. No. 92–463, Section 10(a)(2), 86 Stat. 770, 774 (1972) (Current version at 5 U.S.C. app. section 10(a)(2) (1988); 41 CFR 101–6.1015 (1990).

Dated: October 31, 1997.

Wendy M. Comes,

Executive Director.

[FR Doc. 97–29299 Filed 11–4–97; 8:45 am]

BILLING CODE 1610–01–M

DEPARTMENT OF HEALTH AND HUMAN SERVICES

Advisory Commission on Consumer Protection and Quality in the Health Care Industry's Ad Hoc Work Group on Respect and Nondiscrimination; Notice of Public Meeting

In accordance with Section 10(a)(2) of the Federal Advisory Committee Act, Public Law 92–463, notice is hereby given of the meeting of the Advisory Commission on Consumer Protection and Quality in the Health Care

Industry's Ad Hoc Work Group on Respect and Nondiscrimination. This meeting will be open to the public, limited only by the space available.

Place of meeting: Hubert H. Humphrey Building, Room 800; 200 Independence Avenue, S.W. Washington, D.C. 20201.

Time and Dates: 10:00 a.m.–2:00 p.m., Monday, November 3, 1997.

Purpose/Agenda: To discuss issues related to a draft Consumer Bill of Rights and Responsibilities. Agenda items are subject to change as priorities dictate.

Contact Person: For more information, including substantive program information and summaries of the meeting, please contact: Edward (Chip) Malin, Hubert H. Humphrey Building, Room 118F, 200 Independence Avenue, S.W., Washington, DC 20201; [202/205–3038].

Dated: October 29, 1997.

Janet Corrigan,

Executive Director, Advisory Commission on Consumer Protection and Quality in the Health Care Industry.

[FR Doc. 97–29205 Filed 10–31–97; 10:05 am]

BILLING CODE 4110–60–M

DEPARTMENT OF HEALTH AND HUMAN SERVICES

Health Resources and Services Administration

Agency Information Collection Activities: Submission for OMB Review; Comment Request

Periodically, the Health Resources and Services Administration (HRSA) publishes abstracts of information collection requests under review by the Office of Management and Budget, in compliance with the Paperwork Reduction Act of 1995 (44 U.S.C. Chapter 35). To request a copy of the clearance requests submitted to OMB for review, call the HRSA Reports Clearance Office on (301)-443–1129.

The following request has been submitted to the Office of Management and Budget for review under the Paperwork Reduction Act of 1995:

Proposed Project: Evaluation of the National Health Service Corps —New— The National Health Service Corps (NHSC) was established in 1971 to help correct the maldistribution of health care personnel and to improve the delivery of services in areas with shortages of health care professionals. Through the Scholarship and Loan Repayment Programs the NHSC recruits health clinicians and places them in areas designated as health professional shortage areas.

The evaluation of this program will include three mail surveys, two directed at scholarship and loan repayment program clinicians (physicians, dentists, physician assistants, nurse practitioners and nurse midwives), and one directed at site administrators currently employing NHSC clinicians. The Survey of NHSC Alumni (clinicians who began service on January 1, 1980 and terminated their service before March 14, 1997) will assess alumni attitudes about the NHSC experience including recruitment, placement, and service contributions to the site and community (for example, expanding clinical services, serving in clinical leadership positions, participating in quality improvement activities and initiating community primary care initiatives). In addition, the survey will examine various measures of clinician retention in underserved areas. The Survey of NHSC Clinicians (current) will also assess attitudes about the NHSC experience including recruitment, placement and service contribution to the site and community. The Survey of Administrators in Sites with NHSC Clinicians will assess sites' experiences with NHSC clinicians and will provide an assessment of their service contributions to the site and community. The data collected through the surveys will be used to formulate programmatic and policy recommendations designed to strengthen the NHSC program and increase its effectiveness.

The estimated burden is as follows:

Type of respondent	Number of respondents	Response per respondent	Hours per response	Total burden hours
Eligible Alumni Clinicians	1,555	1	.50	778
Ineligible Alumni Clinicians	173	1	.07	12
Eligible Current Clinicians	965	1	.50	483
Ineligible Current Clinicians	51	1	.07	4
Eligible Site Administrators	251	1	.50	126
Ineligible Site Administrators	13	1	.07	1
Total	3,008			1404

Written comments and recommendations concerning the proposed information collection should be sent within 30 days of this notice to: Laura Oliven, Human Resources and Housing Branch, Office of Management and Budget, New Executive Office Building, Room 10235, Washington, D.C. 20503.

Dated: October 30, 1997.

Jane Harrison,

Acting Director, Division of Policy Review and Coordination.

[FR Doc. 97–29259 Filed 11–4–97; 8:45 am]

BILLING CODE 4160–15–P

DEPARTMENT OF HEALTH AND HUMAN SERVICES

National Institutes of Health

Submission for OMB Review; Comment Request; Gila River Indian Community Demographic Information

SUMMARY: Under the provisions of section 3506(c)(2)(A) of the Paperwork Reduction Act of 1995, the National Institute of Diabetes and Digestive and Kidney Diseases, the National Institutes of Health has submitted to the Office of Management and Budget (OMB) a request to review and approve the information collection listed below. This proposed information collection was previously published in the **Federal Register** on July 11, 1997, on page number 37269 and allowed 60 days for public comment. No public comments were received during the comment period. The purposes of this notice is to allow an additional 30 days for public comment. the National Institutes of Health may not conduct or sponsor, and the respondent is not required to respond to, an information collection that has been extended, revised, or implemented on or after October 1,