

Certification Regarding Environmental Tobacco Smoke

Public Law 103-227, Part C—Environmental Tobacco Smoke, also known as the Pro-Children Act of 1994 (Act), requires that smoking not be permitted in any portion of any indoor routinely owned or leased or contracted for by an entity and used routinely or regularly for provision of health, day care, education, or library services to children under the age of 18, if the services are funded by Federal programs either directly or through State or local governments, by Federal grant, contract, loan, or loan guarantee. The law does not apply to children's services provided in private residences, facilities funded solely by Medicare or Medicaid funds, and portions of facilities used for inpatient drug or alcohol treatment. Failure to comply with the provisions of the law may result in the imposition of a civil monetary penalty of up to \$1,000 per day and/or the imposition of an administrative compliance order on the responsible entity.

By signing and submitting this application the applicant/grantee certifies that it will comply with the requirements of the Act. The applicant/grantee further agrees that it will require the language of this certification be included in any subawards which contain provisions for the children's services and that all subgrantees shall certify accordingly.

[FR Doc. 97-8840 Filed 4-7-97; 8:45 am]

BILLING CODE 4184-01-P

DEPARTMENT OF HEALTH AND HUMAN SERVICES**Food and Drug Administration****Grassroots Consumer Participation: Public Meeting**

AGENCY: Food and Drug Administration, HHS.

ACTION: Notice of a public meeting.

SUMMARY: The Food and Drug Administration (FDA) is announcing a grassroots consumer exchange meeting between the general public and FDA officials. The meeting will be chaired by FDA's Cincinnati District Director and is intended to encourage dialogue between consumers and FDA officials, to solicit consumers' concerns about the drug review process, drug clinical trials (including inclusion of women in clinical trials), and to discuss how FDA can improve consumer services.

DATES: The public meeting will be held on Tuesday, April 29, 1997, 9 a.m. to 12 m.

ADDRESSES: The public meeting will be held at the Cincinnati Bell Long Distance Bldg., 36 East 7th St., rm. 1703, Cincinnati, OH.

FOR FURTHER INFORMATION CONTACT: Marilyn R. Zipkes, Food and Drug Administration, 1141 Central Pkwy.,

Cincinnati, OH 45212, 513-684-3501, ext. 110.

SUPPLEMENTARY INFORMATION: The purpose of the meeting is to encourage dialogue between consumers and FDA officials, to identify consumers' current and future health concerns, and to enhance relations between consumers and FDA. There is no registration fee for this meeting. Interested persons are encouraged to register early because space is limited. To register contact the contact person listed above.

Dated: March 31, 1997.

William K. Hubbard,
Associate Commissioner for Policy Coordination.

[FR Doc. 97-8908 Filed 4-7-97; 8:45 am]

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DEPARTMENT OF HEALTH AND HUMAN SERVICES**Food and Drug Administration**

[Docket No. 97M-0126]

CIBA Vision Corp.; Premarket Approval of Unizyme Enzymatic Cleaner

AGENCY: Food and Drug Administration, HHS.

ACTION: Notice.

SUMMARY: The Food and Drug Administration (FDA) is announcing its approval of the application by CIBA Vision Corp., Duluth, GA, for premarket approval, under the Federal Food, Drug, and Cosmetic Act (the act), of Unizyme Enzymatic Cleaner. FDA's Center for Devices and Radiological Health (CDRH) notified the applicant, by letter of January 31, 1997, of the approval of the application.

DATES: Petitions for administrative review by May 8, 1997.

ADDRESSES: Written requests for copies of the summary of safety and effectiveness data and petitions for administrative review to the Dockets Management Branch (HFA-305), Food and Drug Administration, 12420 Parklawn Dr., rm. 1-23, Rockville, MD 20857.

FOR FURTHER INFORMATION CONTACT: James F. Saviola, Center for Devices and Radiological Health (HFZ-460), Food and Drug Administration, 9200 Corporate Blvd., Rockville, MD 20850, 301-594-1744.

SUPPLEMENTARY INFORMATION: On August 15, 1996, CIBA Vision Corp., Duluth, GA, 30155-1518, submitted to CDRH an application for premarket approval of Unizyme Enzymatic Cleaner. The device

is a periodic cleaner and is indicated for use with (hydrogen peroxide), lens care systems in the weekly cleaning of soft (hydrophilic) contact lenses (including daily wear, extended wear, tinted lenses, and lenses prescribed for scheduled replacement).

In accordance with the provisions of section 515(c)(2) of the act (21 U.S.C. 360e(c)(2)) as amended by the Safe Medical Devices Act of 1990, this premarket approval application (PMA) was not referred to the Ophthalmic Devices Panel of the Medical Devices Advisory Committee, an FDA advisory committee, for review and recommendation because the information in the PMA substantially duplicates information previously reviewed by this panel. On January 31, 1997, CDRH approved the application by a letter to the applicant from the Director of the Office of Device Evaluation, CDRH.

A summary of the safety and effectiveness data on which CDRH based its approval is on file in the Dockets Management Branch (address above) and is available from that office upon written request. Requests should be identified with the name of the device and the docket number found in brackets in the heading of this document.

Opportunity for Administrative Review

Section 515(d)(3) of the act authorizes any interested person to petition, under section 515(g) of the act, for administrative review of CDRH's decision to approve this application. A petitioner may request either a formal hearing under 21 CFR part 12 of FDA's administrative practices and procedures regulations or a review of the application and CDRH's action by an independent advisory committee of experts. A petition is to be in the form of a petition for reconsideration under 21 CFR 10.33(b). A petitioner shall identify the form of review requested (hearing or independent advisory committee) and shall submit with the petition supporting data and information showing that there is a genuine and substantial issue of material fact for resolution through administrative review. After reviewing the petition, FDA will decide whether to grant or deny the petition and will publish a notice of its decision in the **Federal Register**. If FDA grants the petition, the notice will state the issue to be reviewed, the form of the review to be used, the persons who may participate in the review, the time and place where the review will occur, and other details.

Petitioners may, at any time on or before May 8, 1997, file with the Dockets Management Branch (address above) two copies of each petition and supporting data and information, identified with the name of the device and the docket number found in brackets in the heading of this document. Received petitions may be seen in the office above between 9 a.m. and 4 p.m., Monday through Friday.

This notice is issued under the Federal Food, Drug, and Cosmetic Act (secs. 515(d), 520(h)(21 U.S.C. 360e(d), 360j(h))) and under authority delegated to the Commissioner of Food and Drugs (21 CFR 5.10) and redelegated to the Director, Center for Devices and Radiological Health (21 CFR 5.53).

Dated: March 4, 1997.

Joseph A. Levitt,

Deputy Director for Regulations Policy, Center for Devices and Radiological Health.

[FR Doc. 97-8853 Filed 4-7-97; 8:45 am]

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DEPARTMENT OF HEALTH AND HUMAN SERVICES

Health Care Financing Administration

[Document Identifier: HCFA-643]

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

In compliance with the requirement of section 3506(c)(2)(A) of the Paperwork Reduction Act of 1995, the Health Care Financing Administration (HCFA), Department of Health and Human Services, has submitted to the Office of Management and Budget (OMB) the following proposal for the collection of information. Interested persons are invited to send comments regarding the burden estimate or any other aspect of this collection of information, including any of the following subjects: (1) The necessity and utility of the proposed information collection for the proper performance of the agency's functions; (2) the accuracy of the estimated burden; (3) ways to enhance the quality, utility, and clarity of the information to be collected; and (4) the use of automated collection techniques or other forms of information technology to minimize the information collection burden.

Type of Information Collection Request: Reinstatement, without change, of previously approved collection for which approval has expired; **Title of Information Collection:** Hospice Survey and Deficiencies Report Form and Supporting Regulations 42 CFR Sections

488.488.26(c), 442.30(a)(4), 442, Subpart B, C, D, E and F; **Form No.:** HCFA-643; **Use:** The survey report form and supporting regulations are needed to ensure provider compliance with Medicare and Medicaid conditions of participation. In order to participate in the Medicare program, a hospice must meet certain Federal health and safety conditions of participation. The survey report form will be used by State surveyors to record data about a hospice's compliance with Federal health and safety conditions in order to initiate the certification or recertification process. **Frequency:** Annually; **Affected Public:** State, Local or Tribal Govt, Federal Govt; **Number of Respondents:** 2,150; **Total Annual Responses:** 2,150; **Total Annual Hours:** 5,375.

To obtain copies of the supporting statement for the proposed paperwork collections referenced above, access HCFA's WEB SITE ADDRESS at <http://www.hcfa.gov/reg/prdact95.htm>, or to obtain the supporting statement and any related forms, E-mail your request, including your address and phone number, to Paperwork@hcfa.gov, or call the Reports Clearance Office on (410) 786-1326. Written comments and recommendations for the proposed information collections must be mailed within 30 days of this notice directly to the HCFA Paperwork Clearance Officer designated at the following address: OMB Human Resources and Housing Branch, Attention: Allison Eydt, New Executive Office Building, Room 10235, Washington, D.C. 20503.

Dated: March 26, 1997.

Edwin J. Glatzel,

Director, Management Analysis and Planning Staff, Office of Financial and Human Resources, Health Care Financing Administration.

[FR Doc. 97-8949 Filed 4-7-97; 8:45 am]

BILLING CODE 4120-03-P

DEPARTMENT OF HEALTH AND HUMAN SERVICES

National Institutes of Health

National Institute of Dental Research; Notice of Closed Meetings

Pursuant to Section 10(d) of the Federal Advisory Committee Act, as amended (5 U.S.C. Appendix 2), notice is hereby given of the following National Institute of Dental Research Special Emphasis Panel (SEP) meetings:

Name of SEP: National Institute of Dental Research Special Emphasis Panel-Review of R03 application (97-32).

Date: April 3, 1997.

Time: 2:30 p.m.

Place: Natcher Building, Rm. 4AN-44F, National Institutes of Health, Bethesda, MD 20892, (teleconference).

Contact Person: Dr. Philip Washko, Scientific Review Administrator, 4500 Center Drive, Natcher Building, Room 4AN-44F, Bethesda, MD 20892, (301) 594-2372.

Purpose/Agenda: To evaluate and review grant applications and/or contract proposals.

Name of SEP: National Institute of Dental Research Special Emphasis Panel-Review of R03 application (97-31).

Date: April 7, 1997.

Time: 10:00 a.m.

Place: Natcher Building, Rm. 4AN-44F, National Institutes of Health, Bethesda, MD 20892, (teleconference).

Contact Person: Dr. Philip Washko, Scientific Review Administrator, 4500 Center Drive, Natcher Building, Room 4AN-44F, Bethesda, MD 20892, (301) 594-2372.

Purpose/Agenda: To evaluate and review grant applications and/or contract proposals.

Name of SEP: National Institute of Dental Research Special Emphasis Panel-Review of R44 application (97-27).

Date: April 8, 1997.

Time: 1:00 p.m.

Place: Natcher Building, Rm. 4AN-44F, National Institutes of Health, Bethesda, MD 20892, (teleconference).

Contact Person: Dr. George Hausch, Chief, Grants Review Section, 4500 Center Drive, Natcher Building, Room 4AN-44F, Bethesda, MD 20892, (301) 594-2372.

Purpose/Agenda: To evaluate and review grant applications and/or contract proposals.

Name of SEP: National Institute of Dental Research Special Emphasis Panel-Review of R03 Applications (97-40).

Date: April 11, 1997.

Time: 2:00 p.m.

Place: Natcher Building, Rm. 4AN-44F, National Institutes of Health, Bethesda, MD 20892, (teleconference).

Contact Person: Dr. George Hausch, Chief, Grants Review Section, 4500 Center Drive, Natcher Building, Room 4AN-44F, Bethesda, MD 20892, (301) 594-2372.

Purpose/Agenda: To evaluate and review grant applications and/or contract proposals.

Name of SEP: National Institute of Dental Research Special Emphasis Panel-Review of R03 application (97-36).

Date: April 15, 1997.

Time: 10:00 a.m.

Place: Natcher Building, Rm. 4AN-44F, National Institutes of Health, Bethesda, MD 20892, (teleconference).

Contact Person: Dr. Philip Washko, Scientific Review Administrator, 4500 Center Drive, Natcher Building, Room 4AN-44F, Bethesda, MD 20892, (301) 594-2372.

Purpose/Agenda: To evaluate and review grant applications and/or contract proposals.

Name of SEP: National Institute of Dental Research Special Emphasis Panel-Review of R03 applications (97-35).

Date: April 24, 1997.

Time: 10:00 a.m.

Place: Natcher Building, Rm. 4AN-44F, National Institutes of Health, Bethesda, MD 20892, (teleconference).

Contact Person: Dr. Philip Washko, Scientific Review Administrator, 4500 Center