

(SGEs) are generally required to file. Agencies may, if appropriate under the OGE regulation, exclude certain regular employees or SGEs as provided in 5 CFR 2634.905. Reports are normally required to be filed within 30 days of entering a covered position (or earlier if required by the agency concerned), and again annually if the employee serves for more than 60 days in the position. As indicated in § 2634.907 of the OGE regulation, the information required to be collected includes assets and sources of income, gifts and travel reimbursements, liabilities, employment agreements and arrangements, and outside positions, subject to certain thresholds and exclusions.

Most of the persons who file this report form are current executive branch Government employees at the time they complete the forms. However, some filers are private citizens who are asked by their prospective agency to file a new entrant report prior to entering Government service in order to permit advance checking for any potential conflicts of interest and resolution thereof by agreement to recuse, divest, obtaining of a waiver, etc. Based on OGE's annual agency ethics questionnaire responses, approximately 285,000 SF 450 report forms were filed during 1994 throughout the executive branch. Of these, OGE estimates that no more than between 5% and 10%, or some 14,500 to 28,500 per year at most, are filed by private citizens, those potential regular employees whose positions are designated for confidential disclosure filing as well as potential special Government employees whose agencies require that they file their new entrant reports prior to assuming Government responsibilities.

Each filing is estimated to take an average of one and one-half hours. The number of private citizens whose reports are filed each year with OGE is less than 10, but pursuant to 5 CFR 1320.7(s)(1), the lower limit for this general regulatory-based requirement is set at 10 private persons (OGE-processed reports). This yields an annual reporting burden of 15 hours, the same as in the current OMB inventory for this information collection. The remainder of the private citizen reports are filed with other departments and agencies throughout the executive branch.

Public comment is invited on each aspect of the proposed new OGE Form 450 as set forth in this notice, including specifically views on the need for and practical utility of this proposed modified collection of information, the accuracy of OGE's burden estimate, the enhancement of quality, utility and

clarity of the information collected, and the minimization of burden (including the use of information technology).

Comments received submitted in response to this notice will be summarized for, and may be included with, the OGE request for OMB paperwork approval for this modified information collection. The comments will also become a matter of public record.

Approved: August 28, 1995.

Stephen D. Potts,

Director, Office of Government Ethics.

[FR Doc. 95-21753 Filed 8-31-95; 8:45 am]

BILLING CODE 6345-01-U

DEPARTMENT OF HEALTH AND HUMAN SERVICES

Office of the Secretary

Notice of a Meeting of the Commission on Research Integrity

Pursuant to P.L. 92-463, notice is hereby given of a public meeting of the Commission on Research Integrity.

The meeting will be held at the Washington Dulles Airport Marriott Hotel on Monday and Tuesday, September 18-19, from 8:30 a.m. until 5 p.m.

The Commission will be working on its remaining recommendations and final report to congressional oversight committees and the Secretary of the Department of Health and Human Services (HHS) on the administration of Section 493 of the Public Health Service Act, as amended by and added to by Section 161 of the NIH Revitalization Act of 1993. Recommendations include possible administrative actions to improve the HHS scientific misconduct apparatus. Interested parties are advised to call the Executive Secretary, Ms. Henrietta Hyatt-Knorr, shortly before the meeting to verify the date, place, and agenda. Persons wishing to make a presentation should contact her in writing at Rockwall II, Suite 700, 5515 Security Lane, Rockville MD 20852 or by phone at (301) 443-5300, by fax at (301) 443-5351, or via internet at hhyatt@oasch.ssw.dhhs.gov. Ms. Hyatt-Knorr will furnish a meeting agenda upon request. Depending on the number of speakers and other constraints, the Executive Secretary will allocate a timeframe for anyone wishing to make an oral presentation.

Henrietta D. Hyatt-Knorr,

Executive Secretary, Commission on Research Integrity.

[FR Doc. 95-21726 Filed 8-31-95; 8:45 am]

BILLING CODE 4160-17-P

Administration for Children and Families

President's Committee on Mental Retardation; Meeting

Agency Holding the Meeting: President's Committee on Mental Retardation.

Time and Date: Full Committee Meeting, September 7, 1995, 2 p.m.-6 p.m., September 8, 1995, 9 a.m.-5 p.m.

Place: Madison Hotel, 1177 15th Street NW., Washington, DC 20005.

Status: Meetings are open to the public. An interpreter for the deaf will be available upon advance request. All locations are barrier free.

To Be Considered: The Committee plans to discuss critical issues concerning Federal Policy, Federal Research and Demonstration, State Policy Collaboration, Minority and Cultural Diversity and Mission and Public Awareness.

The PCMR acts in an advisory capacity to the President and the Secretary of the U.S. Department of Health and Human Services on a broad range of topics relating to programs and services for persons with mental retardation. The Committee, by Executive Order, is responsible for evaluating the adequacy of current practices in programs for persons with mental retardation, and for reviewing legislative proposals that impact the quality of life that is experienced by citizens with mental retardation and their families.

Contact Person for More Information: Gary H. Blumenthal, Wilbur J. Cohen Building, Room 5325, 330 Independence Avenue SW., Washington, DC 20201-0001, (202) 619-0634.

Dated: August 29, 1995.

Gary H. Blumenthal,

Executive Director, PCMR.

[FR Doc. 95-21782 Filed 8-31-95; 8:45 am]

BILLING CODE 4184-01-M

Agency for Health Care Policy and Research

Notice of Advisory Committee Meetings

In accordance with section 10(a) of the Federal Advisory Committee Act (5 U.S.C., Appendix 2) announcement is made of the following advisory committees scheduled to meet during the months of October and November 1995:

Name: Health Services Developmental Grants Review Subcommittee.

Date and Time: October 18-19, 1995 8:00 a.m.

Place: The DoubleTree, 1750 Rockville Pike, Conference Room TBA, Rockville, Maryland 20852.

Open October 18, 8:00 a.m. to 9:00 a.m.
Closed for remainder of meeting.

Purpose: The Subcommittee is charged with the initial review of grant applications proposing experimental, analytical and theoretical research on costs, quality, access, effectiveness, and efficiency of the delivery of health services for the research grant program administered by the Agency for Health Care Policy and Research (AHCPR).

Agenda: The open session of the meeting on October 18 from 8:00 a.m. to 9:00 a.m. will be devoted to a business meeting covering administrative matters and reports. During the closed session, the Subcommittee will be reviewing and discussing grant applications dealing with health services research issues. In accordance with the Federal Advisory Committee Act, section 10(d) of 5 U.S.C., Appendix 2 and 5 U.S.C., 552b(c)(6), it has been determined that this latter session will be closed because the discussions are likely to reveal personal information concerning individuals associated with the applications. This information is exempt from mandatory disclosure.

Anyone wishing to obtain a roster of members, minutes of the meeting, or other relevant information should contact J. Terrell Hoffeld, D.D.S., Ph.D., Scientific Review Administrator, Office of Scientific Affairs, Agency of Health Care Policy and Research, Suite 400, Executive Office Center, 2101 East Jefferson Street, Rockville, Maryland 20852, Telephone (301) 594-1449.

Name: Health Services Research Review Subcommittee.

Date and Time: October 18-20, 1995, 8:00 a.m.

Place: Ramada Inn, 1775, Rockville Pike, Conference Room TBA, Rockville, Maryland 20852.

Open October 18, 8:00 a.m. to 8:45 a.m.
Closed for remainder of meeting.

Purpose: The Subcommittee is charged with the initial review of grant applications proposing analytical and theoretical research on costs, quality, access, and efficiency of the delivery of health services for the research grant program administered by the Agency for Health Care Policy and Research (AHCPR).

Agenda: The open session of the meeting on October 18, from 8:00 a.m. to 8:45 a.m. will be devoted to a business meeting covering administrative matters and reports. During the closed session, the Subcommittee will be reviewing and discussing grant applications dealing with health services research issues. In accordance with the Federal Advisory Committee Act, section 10(d) of 5 U.S.C., Appendix 2 and 5 U.S.C., 552b(c)(6), it has been determined that this latter session will be closed because the discussions are likely to reveal personal information concerning individuals associated with the applications. This information is exempt from mandatory disclosure.

Anyone wishing to obtain a roster of members, minutes of the meeting, or other relevant information should contact Patricia G. Thompson, Ph.D., Scientific Review Administrator, Office of Scientific Affairs, Agency for Health Care Policy and Research, Suite 400, Executive Office Center, 2101 East Jefferson Street, Rockville, Maryland 20852, Telephone (301) 594-1451.

Name: Health Services Research Dissemination Study Section.

Date and Time: October 23-24, 1995, 8:00 a.m.

Place: Holiday Inn Chevy Chase, 5520 Wisconsin Avenue, Conference Room TBA, Chevy Chase, Maryland 20815.

Open October 23, 1995, 8:00 a.m.
Closed for remainder of meeting.

Purpose: The Study Section is charged with the review of and making recommendations on grant applications for Federal support of conferences, workshops, meetings, or projects related to dissemination and utilization of research findings, and AHCPR liaison with health care policy makers, providers, and consumers.

Agenda: The open session of the meeting on October 23 from 8:00 a.m. to 8:30 a.m. will be devoted to general business matters. During the closed portions of the meeting, the Study Section will be reviewing and discussing grant applications dealing with health services research issues. In accordance with the Federal Advisory Committee Act, section 10(d) of 5 U.S.C., Appendix 2 and 5 U.S.C. 552b(c)(6), it has been determined that this latter session will be closed because the discussions are likely to reveal personal information concerning individuals associated with the grant applications. This information is exempt from mandatory disclosure.

Anyone wishing to obtain a roster of members, minutes of the meeting, or other relevant information should contact Linda Blankenbaker, Scientific Review Administrator, Office of Scientific Affairs, Agency for Health Care Policy and Research, Suite 400, 2101 East Jefferson Street, Rockville, Maryland 20852, Telephone (301) 594-1438.

Name: Health Care Technology Study Section.

Date and Time: November 6-7, 1995, 8:00 a.m.

Place: Hyatt Regency, One Bethesda Metro Center, Conference Room TBA, Bethesda, Maryland 20814.

Open November 6, 8:00 a.m. to 9:00 a.m.
Closed for remainder of meeting.

Purpose: The Study Section is charged with conducting the initial review of health services research grant applications concerned with medical decisionmaking, computers in health care delivery, and the utilization and effects of health care technologies and procedures.

Agenda: The open session on November 6 from 8:00 a.m. to 9:00 a.m. will be devoted to a business meeting covering administrative matters and reports. The closed session of the meeting will be devoted to reviewing and discussing grant applications dealing with health services research issues. In accordance with the Federal Advisory Committee Act, section 10(d) of 5 U.S.C., Appendix 2 and 5

U.S.C., 552b(c)(6), it has been determined that this latter session will be closed because the discussions are likely to reveal personal information concerning individuals associated with the applications. This information is exempt from mandatory disclosure.

Anyone wishing to obtain a roster of members, minutes of the meeting, or other relevant information should contact Karen Rudzinski, Ph.D., Scientific Review Administrator, Office of Scientific Affairs, Agency for Health Care Policy and Research, Suite 400, Executive Office Center, 2101 East Jefferson Street, Rockville, Maryland 20852, Telephone (301) 594-1437.

Agenda items for all meetings are subject to change as priorities dictate.

Dated: August 24, 1995.

Clifton R. Gaus,

Administrator.

[FR Doc. 95-21816 Filed 8-31-95; 8:45 am]

BILLING CODE 4160-90-M

Food and Drug Administration

[Docket No. 95C-0286]

Ebonex Corp.; Filing of a Color Additive Petition

AGENCY: Food and Drug Administration, HHS.

ACTION: Notice.

SUMMARY: The Food and Drug Administration (FDA) is announcing that Ebonex Corp. has filed a petition proposing that the color additive regulations be amended to provide for the safe use of bone black as a color additive in cosmetics, including cosmetics intended for use in the eye area.

DATES: Written comments on the petitioner's environmental assessment by October 2, 1995.

ADDRESSES: Submit written comments to the Dockets Management Branch (HFA-305), Food and Drug Administration, rm. 1-23, 12420 Parklawn Dr., Rockville, MD 20857.

FOR FURTHER INFORMATION CONTACT: Mitchell A. Cheeseman, Center for Food Safety and Applied Nutrition (HFS-217), Food and Drug Administration, 200 C St. SW., Washington, DC 20204, 202-418-3083.

SUPPLEMENTARY INFORMATION: Under the Federal Food, Drug, and Cosmetic Act (sec. 721(d)(1) (21 U.S.C. 379e(d)(1))), notice is given that a color additive petition (CAP 5C0247) has been filed by Ebonex Corp., P.O. Box 3247, Melvindale, MI 48122. The petition proposes to amend the color additive regulations to provide for the safe use of bone black as a color additive in cosmetics, including cosmetics intended for use in the eye area.