Information Services, 370 L'Enfant Promenade, S.W., Washington, D.C. 20447, Attn: ACF Reports Clearance Officer.

OMB Comment: OMB is required to make a decision concerning the collection of information between 30 and 60 days after publication of this document in the **Federal Register**. Therefore, a comment is best assured of having its full effect if OMB receives it within 30 days of publication. Written comments and recommendations for the proposed information collection should be sent directly to the following: Office of Management and Budget, Paperwork Reduction Project, 725 17th Street, N.W., Washington, D.C. 20503, Attn: Ms. Wendy Taylor.

Dated: February 25, 2000.

Bob Sargis,

Acting Reports Clearance Officer.

[FR Doc. 00–4969 Filed 3–1–00; 8:45 am]

BILLING CODE 4184–01–M

DEPARTMENT OF HEALTH AND HUMAN SERVICES

Administration for Children and Families

Submission for OMB Review; Comment Request

Title: Program Narrative Objective Work Plan (OWP).

OMB No.: 0980–0204.

Description: Program Narrative (OWP) information is collected as part of a competitive, discretionary grant application submitted to the Administration for Native Americans (ANA). Included with the OWP are standard, government-wide Federal assistance application forms (*e.g.,* SF–424, 424A, 424B, Non-Constructions Assurances, and various OMB certifications). The OWP provides information used by legislatively mandated project evaluation panels to compete and rank applications. ANA uses the OWP information to perform legislatively mandated project evaluations supporting the basis for recommendations to award or not award ANA grants. After funding, the OWP is used to reflect funded objectives and to administer and monitor ANA grants.

OWP information presents the grant applicants' locally-determined project objectives and plan to achieve those objectives. Economic development projects may attach a business plan. OWP information is presented as narrative and transcribed onto a government form titled, ''ANA Objective Work Plan''. In the past, ANA used two forms to collect the program narrative; *i.e.,* ''Program Narrative Objective Work Plan'' and ''Program Narrative Approach.'' The new, single form combines the two old forms and eliminates some information items.

Instructions for completing the OWP are provided in the ''Administration for Narrative Americans Application Packet for Financial Assistance.'' Instructions for compiling a complete application are provided in the packet. The OWP and instruction packet are used in all ANA competitive discretionary grant programs such as Social and Economic Development Strategies (SEDS), Native American Languages Preservation, Environmental Regulatory Enhancement, etc.

Resondents: State, Local or Tribal Government.

ANNUAL BURDEN ESTIMATES

Instrument	Number of respondents	Number of responses per respondent	Average burden hours per response	Total burden hours
OWP	650	1	28	18,200

Additional Information: Copies of the proposed collection may be obtained by writing to The Administration for Children and Families, Office of Information Services, 370 L'Enfant Promenade, SW., Washington, DC 20447, Attn: ACF Reports Clearance Officer.

OMB Comment: OMB is required to make a decision concerning the collection of information between 30 and 60 days after publication of this document in the **Federal Register.** Therefore, a comment is best assured of having its full effect if OMB receives it within 30 days of publication. Written comments and recommendations for the proposed information collection should be sent directly to the following: Office of Management and Budget, Paperwork Reduction Project, 725 17th Street, NW., Washington, DC 20503, Attn: Ms. Wendy Taylor.

Dated: February 25, 2000.

Bob Sargis,

Reports Clearance Officer.

[FR Doc. 00–5029 Filed 3–1–00; 8:45 am]

BILLING CODE 4184–01–M

DEPARTMENT OF HEALTH AND HUMAN SERVICES

Food and Drug Administration

[Docket No. 00F–0786]

Eka Chemicals, Inc.; Filing of Food Additive Petition

AGENCY: Food and Drug Administration, HHS.

ACTION: Notice.

SUMMARY: The Food and Drug Administration (FDA) is announcing that Eka Chemicals, Inc., has filed a petition proposing that the food additive regulations be amended to provide for the safe use of chlorine dioxide produced by another method.

DATES: Submit written comments on the petitioner's environmental assessment by April 3, 2000.

ADDRESSES: Submit written comments to the Dockets Management Branch (HFA–305), Food and Drug Administration, 5630 Fishers Lane, rm. 1061, Rockville, MD 20852.

FOR FURTHER INFORMATION CONTACT: Robert L. Martin, Center for Food Safety and Applied Nutrition (HFS–215), Food and Drug Administration, 200 C St. SW., Washington, DC 20204–0001, 202–418–3074.

SUPPLEMENTARY INFORMATION: Under the Federal Food, Drug, and Cosmetic Act (sec. 409(b)(5) (21 U.S.C. 348(b)(5))), notice is given that a food additive petition (FAP 0A4716) has been filed by Eka Chemicals, Inc., c/o Keller and Heckman LLP, 1001 G St. NW., suite 500 West, Washington, DC 20001. The petition proposes to amend the food additive regulations in § 173.300 Chlorine dioxide (21 CFR 173.300) to

provide for the safe use of chlorine dioxide produced by another method.

The potential environmental impact of this action is being reviewed. To encourage public participation consistent with regulations issued under the National Environmental Policy Act (40 CFR 1501.4(b)), the agency is placing the environmental assessment submitted with the petition that is the subject of this notice on public display at the Dockets Management Branch (address above) for public review and comment. Interested persons may submit to the Dockets Management Branch (address above) written comments by April 3, 2000. Two copies of any comments are to be submitted, except that individuals may submit one copy. Comments are to be identified with the docket number found in brackets in the heading of this document. Received comments may be seen in the office above between 9 a.m. and 4 p.m. Monday through Friday. FDA will also place on public display any amendments to, or comments on, the petitioner's environmental assessment without further announcement in the **Federal Register**. If, based on its review, the agency finds that an environmental impact statement is not required and this petition results in a regulation, the notice of availability of the agency's finding of no significant impact and the evidence supporting that finding will be published with the regulation in the **Federal Register** in accordance with 21 CFR 25.40(c).

Dated: February 14, 2000.

Alan M. Rulis,

Director, Office of Premarket Approval, Center for Food Safety and Applied Nutrition.

[FR Doc. 00–5015 Filed 3–1–00; 8:45 am]

BILLING CODE 4160–01–F

DEPARTMENT OF HEALTH AND HUMAN SERVICES

Food and Drug Administration

[Docket No. 00F–0789]

National Center for Food Safety and Technology; Filing of Food Additive Petition

AGENCY: Food and Drug Administration, HHS.

ACTION: Notice.

SUMMARY: The Food and Drug Administration (FDA) is announcing that the National Center for Food Safety and Technology, Illinois Institute of Technology, has filed a petition proposing that the food additive regulations be amended to expand the conditions of safe use for X-radiation and electron beam energy sources for the treatment of prepackaged foods by irradiation.

FOR FURTHER INFORMATION CONTACT: Mark A. Hepp, Center for Food Safety and Applied Nutrition (HFS–215), Food and Drug Administration, 200 C St. SW., Washington, DC 20204, 202–418–3098.

SUPPLEMENTARY INFORMATION: Under the Federal Food, Drug, and Cosmetic Act (sec. 409(b)(5) (21 U.S.C. 348(b)(5))), notice is given that a food additive petition (FAP 0M4711) has been filed by the National Center for Food Safety and Technology, Illinois Institute of Technology, 6502 South Archer Rd., Summit-Argo, IL 60501–1933. The petition proposes to amend the food additive regulations in § 179.45 *Packaging materials for use during the irradiation of prepackaged foods* (21 CFR 179.45) to expand the conditions of safe use for X-radiation and electron beam energy sources for the treatment of prepackaged foods by irradiation.

The agency has determined under 21 CFR 25.32(j) that this action is of a type that does not individually or cumulatively have a significant effect on the human environment. Therefore, neither an environmental assessment nor an environmental impact statement is required.

Dated: February 14, 2000.

Alan L. Rulis,

Director, Office of Premarket Approval, Center for Food Safety and Applied Nutrition.

[FR Doc. 00–5014 Filed 3–1–00; 8:45 am]

BILLING CODE 4160–01–F

DEPARTMENT OF HEALTH AND HUMAN SERVICES

Food and Drug Administration

Urology Subcommittee of the Advisory Committee for Reproductive Health Drugs; Notice of Meeting

AGENCY: Food and Drug Administration, HHS.

ACTION: Notice.

This notice announces a forthcoming meeting of a public advisory committee of the Food and Drug Administration (FDA). The meeting will be open to the public.

Name of Committee: Urology Subcommittee of the Advisory Committee for Reproductive Health Drugs

General Function of the Committee: To provide advice and recommendations to the agency on FDA's regulatory issues.

Date and Time: The meeting will be held on April 10, 2000, 9 a.m. to 5 p.m.

Location: Holiday Inn, Versailles Ballrooms, 8120 Wisconsin Ave., Bethesda, MD.

Contact Person: Sandra Titus or Jayne E. Peterson or Robin M. Spencer, Center for Drug Evaluation and Research (HFD–21), Food and Drug Administration, 5600 Fishers Lane, Rockville, MD 20857, 301–827–7001, or e-mail: TITUSS@CDER.FDA.GOV, or FDA Advisory Committee Information Line, 1–800–741–8138 (301–443–0572 in the Washington, DC area), code 12537. Please call the Information Line for up-to-date information on this meeting.

Agenda: The subcommittee will consider the safety and efficacy of new drug application 21–118, Uprima™ (apomorphine HCl tablets, sublingual, TAP Holdings) proposed for use in the treatment of erectile dysfunction.

Procedure: Interested persons may present data, information, or views, orally or in writing on issues pending before the subcommittee. Written submissions may be made to the contact person by April 5, 2000. Oral presentations from the public will be scheduled between approximately 1 p.m. and 2 p.m. Time allotted for each presentation may be limited. Those desiring to make formal oral presentations should notify the contact person before April 5, 2000, and submit a brief statement of the general nature of the evidence or arguments they wish to present, the names and addresses of proposed participants, and an indication of the approximate time requested to make their presentation.

Notice of this meeting is given under the Federal Advisory Committee Act (5 U.S.C. app. 2).

Dated: February 18, 2000.

Linda A. Suydam,

Senior Associate Commissioner.

[FR Doc. 00–4947 Filed 3–1–00; 8:45 am]

BILLING CODE 4160–01–F

DEPARTMENT OF HEALTH AND HUMAN SERVICES

National Institutes of Health

Office of the Director, National Institutes of Health; Notice of Meeting

Pursuant to section 10(a) of the Federal Advisory Committee Act, as amended (5 U.S.C. Appendix 2), notice is hereby given of a meeting of the Office of AIDS Research Advisory Council.

The meeting will be open to the public, with attendance limited to space available. Individuals who plan to attend and need special assistance, such as sign language interpretation or other