Applicants must provide a certification concerning Lobbying. Prior to receiving an award in excess of $100,000, applicants shall furnish an executed copy of the lobbying certification. Applicants must sign and return the certification with their applications.

Requirements, Debarment and Other Responsibilities and Environmental Tobacco Smoke Certifications. A signature on the application constitutes an assurance that the applicant will comply with the pertinent Departmental regulations contained in 45 CFR part 74.

5. Documents of Support. The maximum number of pages for supporting documentation is 10 pages, double-spaced, exclusive of letters of support or agreement. These documents must be numbered and might include resumes, photocopies of news clippings, evidence of the program's efforts to coordinate child care services at the local level, etc. Documentation over the ten-page limit will not be reviewed. The applicant may, however, include as many letters of support or agreement as are appropriate.

B. *Application Submission:* To be considered for funding, the applicant must submit one signed original and two additional copies of the application, including all attachments, to the application receipt point specified above. The original copy of the application must have original signatures, signed in black ink. Each copy must be stapled (back and front) in the upper left corner. All copies of an application must be submitted in a single package.

Because each application will be duplicated, do not use or include separate covers, binders, clips, tabs, plastic inserts, maps, brochures or any other items that cannot be processed easily on a photocopy machine with an automatic feed. Do not bind, clip, staple, or fasten in any way separate subsections of the application, including supporting documentation. Applicants are advised that the copies of the application submitted, not the original, will be reproduced by the Federal government for review.

Dated: April 5, 2001.

James A. Harrell,

Acting Commissioner, Administration on Children, Youth and Families.

[FR Doc. 01–8994 Filed 4–11–01; 8:45 am]

BILLING CODE 4184–01–P

DEPARTMENT OF HEALTH AND HUMAN SERVICES

Food and Drug Administration

[Docket No. 01P–0150]

Salad Dressing Deviating From Identity Standard; Temporary Permit for Market Testing

AGENCY: Food and Drug Administration, HHS.

ACTION: Notice.

SUMMARY: The Food and Drug Administration (FDA) is announcing that a temporary permit has been issued to Kraft Foods, Inc., to market test a product designated as ''salad dressing'' that deviates from the U.S. standard of identity for salad dressing. The purpose of the temporary permit is to allow the applicant to measure consumer acceptance of the product, identify mass production problems, and assess commercial feasibility, in support of a petition to amend the standard of identity for salad dressing.

DATES: This permit is effective for 15 months, beginning on the date the food is introduced or caused to be introduced into interstate commerce, but not later than July 11, 2001.

FOR FURTHER INFORMATION CONTACT: Loretta A. Carey, Center for Food Safety and Applied Nutrition (HFS–822), Food and Drug Administration, 200 C St. SW., Washington, DC 20204, 202–205–4168.

SUPPLEMENTARY INFORMATION: In accordance with 21 CFR 130.17 concerning temporary permits to facilitate market testing of foods deviating from the requirements of the standards of identity issued under section 401 of the Federal Food, Drug, and Cosmetic Act (21 U.S.C. 341), FDA is giving notice that a temporary permit has been issued to Kraft Foods, Inc., Three Lakes Dr., Northfield, IL 60093–2753.

The permit covers limited interstate marketing tests of products identified as ''salad dressing'' that deviate from the U.S. standard of identity for salad dressing (21 CFR 169.150) by adding potassium sorbate, which is not permitted under the current standard, and by reducing the amount of egg 2 percent below the amount required by the current standard. The test product meets all the requirements of the standard with the exception of the reduced amount of egg level in the product and the addition of potassium sorbate. Because test preferences vary by area, along with social and environmental differences, the purpose of this permit is to test the product throughout the United States.

Under this temporary permit, the salad dressing will be test marketed as ''salad dressing.''

This permit provides for the temporary marketing of 150 million pounds of product during the entire 15-month period. The test product will be manufactured by Kraft Foods, Inc., at 2340 Forest Lane, Garland, TX 75040; 1701 West Bradley Ave., Champaign, IL 61821; and 7352 Industrial Blvd., Allentown, PA 18106. The product will be distributed throughout the United States.

The information panel of the labels will bear nutrition labeling in accordance with 21 CFR 101.9. Each of the ingredients used in the food must be declared on the labels as required by the applicable sections of 21 CFR part 101. This permit is effective for 15 months, beginning on the date the food is introduced or caused to be introduced into interstate commerce, but not later than July 11, 2001.

Dated: April 3, 2001.

Christine J. Lewis,

Director, Office of Nutritional Products Labeling and Dietary Supplements, Center for Food Safety and Applied Nutrition.

[FR Doc. 01–8978 Filed 4–11–01; 8:45 am]

BILLING CODE 4160–01–S

DEPARTMENT OF HEALTH AND HUMAN SERVICES

Food and Drug Administration

Science Board to the Food and Drug Administration Advisory Committee; Amendment of Notice

AGENCY: Food and Drug Administration, HHS.

ACTION: Notice.

SUMMARY: The Food and Drug Administration (FDA) is announcing an amendment to the notice of a public meeting of the Science Board to the Food and Drug Administration Advisory Committee. The meeting was announced in the **Federal Register** of March 23, 2001 (65 FR 16253). The amendment is being made to reflect changes in the *Agenda* portion of the meeting notice. The time for the open public hearing and open committee discussion has been changed. This meeting is open to the public. There are no other changes.

FOR FURTHER INFORMATION CONTACT: Susan Bond, Office of the Commissioner, Office of Science Coordination and Communication (HF–33), Food and Drug Administration, 5600 Fishers Lane, rm. 17–35, Rockville, MD 20857, 301–827–6687, or FDA