NOTIFICATION PROCEDURE:

For the purpose of access, the subject individual should write to the system manager who will require the system name, address, age, gender type, and, for verification purposes, the subject individual's name (woman's maiden name, if applicable).

RECORD ACCESS PROCEDURE:

For purpose of access, use the same procedures outlined in Notification Procedures above. Requestors should also reasonably specify the record contents being sought. (These procedures are in accordance with Department regulation 45 CFR 5b.5(a)(2)).

CONTESTING RECORD PROCEDURES:

The subject individual should contact the system manager named above, and reasonably identify the record and specify the information to be contested. State the corrective action sought and the reasons for the correction with supporting justification. (These procedures are in accordance with Department regulation 45 CFR 5b.7).

SYSTEMS EXEMPTED FROM CERTAIN PROVISIONS OF THE ACT:

None

[FR Doc. 05–19676 Filed 9–30–05; 8:45 am] BILLING CODE 4120-03-P

DEPARTMENT OF HEALTH AND HUMAN SERVICES

Administration for Children and Families

Administration on Children, Youth, and Families; Notice of Award of Non-**Competitive Grant**

AGENCY: Administration on Children, Youth and Families, ACF, HHS. **ACTION:** Award announcement.

SUMMARY: The Administration on Children, Youth and Families herein announces an urgent grant award to the National Association of Child Care **Resource and Referral Agencies** (NACCRRA) to provide technical assistance to reestablish the operations of the resource and referral agencies in Mississippi and Louisiana whose operations have been disrupted by Hurricane Katrina. This grant will help to re-establish child care referral services so that families along the Gulf Coast can find child care. This grant will also support local and Statewide inventories of child care need and availability.

The amount of the proposed grant to NACCRRA is \$99,500 in FY 2005 child care funds. The duration of the grant is 12 months.

Statutory Authority: This award will be made pursuant to the Child Care and Development Block Grant Act of 1990 as amended (CCDBG Act); section 418 of the Social Security Act; Consolidated Appropriations Act, 2001 (Pub. L. 106-554). FOR FURTHER INFORMATION CONTACT:

Shannon Rudisill, Director of Technical Assistance, Child Care Bureau, at 202-205-8051.

Dated: September 27, 2005.

Joan E. Ohl,

Commissioner, Administration on Children, Youth and Families.

[FR Doc. 05-19650 Filed 9-30-05; 8:45 am] BILLING CODE 4184-01-P

DEPARTMENT OF HEALTH AND HUMAN SERVICES

Food and Drug Administration

[Docket No. 2005P-0376]

Iceberg Water 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 Canada's Original ICEBERG Water Corp., to market a product designated as "Canada's Original Iceberg Water" that deviates from the U.S. standard of identity for bottled water. 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.

DATES: This permit is effective for 15 months, beginning on the date the permit holder introduces or causes the introduction of the test product into interstate commerce, but not later than January 3, 2006.

FOR FURTHER INFORMATION CONTACT: Loretta A. Carey, Center for Food Safety and Applied Nutrition (HFS-820), Food and Drug Administration, 5100 Paint Branch Pkwy., College Park, MD 20740, 301-436-2371.

SUPPLEMENTARY INFORMATION: Inaccordance 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 Canada's Original ICEBERG Water Corp., 23 Lesmill Rd., suite 304, Toronto, Ontario Canada, M3B-3P6.

The permit covers limited interstate marketing tests of products identified as "Canada's Original Iceberg Water" that deviate from the U.S. standard of identity for bottled water (§165.110 (21 CFR 165.110)) in that the source of the water is an iceberg. The test product meets all the requirements of the standard with the exception of the source definition. The purpose of this permit is to test the product throughout the United States, in order to allow the applicant to measure consumer acceptance of the product, identify mass production problems, and assess commercial feasibility.

This permit provides for the temporary marketing of 500,000 cases of the 24 x 500 milliliter bottles and 500,000 cases of the 12 x 1 liter bottles, totaling 1 million cases per year. The total fluid quantity covered by this application is 12 million liters (3,170,065 gallons). The test product will be manufactured for Canada's Original ICEBERG Water Corp., by Discovery Springs, Daniel's Point Rd., Trepassey, Newfoundland, Canada A0Å–4B0. Canada's Original ICEBERG Water Corp. will distribute the test products throughout the United States. The information panel of the labels will bear nutrition labeling in accordance with 21 CFR 101.9. The bottled water will be manufactured in accordance with the quality standards in §165.110 and the requirements for processing and bottling of bottled drinking water in 21 CFR part 129. 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 January 3, 2006.

Dated: September 22, 2005.

Barbara Schneeman,

Director, Office of Nutritional Products, Labeling, and Dietary Supplements, Center for Food Safety and Applied Nutrition. [FR Doc. 05-19728 Filed 9-30-05; 8:45 am] BILLING CODE 4160-01-S

DEPARTMENT OF HEALTH AND **HUMAN SERVICES**

Food and Drug Administration

[Docket No. 2005N-0394]

Food and Drug Administration's **Communication of Drug Safety** Information; Public Hearing

AGENCY: Food and Drug Administration, HHS.