Bartholomew or Jefferson counties, Indiana, that is no longer in operation as a supermarket, except (1) prior to sale, sublease, assignment, or change in occupancy or (2) to relocate such fixtures or equipment in the ordinary course of business to any other supermarket owned or operated by Kroger.

The Proposed Respondents are required to provide to the Commission a report of compliance with the Order within 30 days following the date on which they signed the consent agreement and every 30 days thereafter until the diversitures are completed: and Kroger must report annually for a period of 10 years from the date the proposed order becomes final. The obligations of Group under the Order will terminate upon the date it becomes final.

The Order has been placed on the public record for 60 days for receipt of comments by interested persons. Comments received during this period will become part of the public record. After 60 days, the Commission will again review the agreement and the comments received and will decide whether it should withdraw from the agreement or make the Order final.

By accepting the Order subject to final approval, the Commission anticipates that the competitive problems alleged in the compliant will be resolved. The purpose of this analysis is to invite public comment on the Order, including the proposed sale of supermarkets to Roundy's, in order to aid the Commission in its determination of whether to make the Order final. This analysis is not intended to constitute an official interpretation of the Order nor is it intended to modify the terms of the Order in any way.

By direction of the Commission.

**Benjamin I. Berman,**

*Acting Secretary.*

[FR Doc. 99–22575 Filed 8–30–99; 8:45 am]

**BILLING CODE 6750–01–M**

## DEPARTMENT OF HEALTH AND HUMAN SERVICES

### Centers for Disease Control and Prevention

### The National Center for Environmental Health (NCEH) of the Centers for Disease Control and Prevention (CDC) Announces the Following Meeting

*Name:* Pediatric Centers Directors Meeting.

*Times and Dates:* 8 a.m.–4:30 p.m., October 25, 1999; 8 a.m.–4:30 p.m., October 26, 1999.

*Place:* Sheraton Buckhead, 3405 Lenox Road NE, Atlanta, Georgia 30326, telephone 404–261–9250.

*Status:* Open to the public, limited only by space available. The meeting room accommodates approximately 100 people.

*Purpose:* The purpose of this meeting is to provide a forum for the Directors of the funded Pediatric Centers to review program progress and discuss future plans, prevention issues and concerns.

*Matters to be Discussed:* Agenda items include perspectives of funding agencies and updates from funded centers.

Agenda items are subject to change as priorities dictate.

**CONTACT PERSON FOR MORE INFORMATION:** James Rifenburg, Air Pollution and Respiratory Health Branch, Division of Environmental Hazards and Health Effects, NCEH, CDC, 4770 Buford Highway, NE, M/S F–39, Atlanta, Georgia 30341–3724, telephone 770/488–7320.

The Director, Management Analysis and Services Office, has been delegated the authority to sign **Federal Register** notices pertaining to announcements of meetings and other committee management activities for both CDC and the Agency for Toxic Substances and Disease Registry.

Dated: August 24, 1999.

**John C. Burckhardt,**

*Acting Director, Management Analysis and Services Office, Centers for Disease Control and Prevention.*

[FR Doc. 99–22526 Filed 8–30–99; 8:45 am]

**BILLING CODE 4168–18–P**

## DEPARTMENT OF HEALTH AND HUMAN SERVICES

### Centers for Disease Control and Prevention

### National Vaccine Advisory Committee, Subcommittee on Future Vaccines, Subcommittee on Immunization Coverage, and Subcommittee on Vaccine Safety and Communication Meetings

In accordance with section 10(a)(2) of the Federal Advisory Committee Act (Pub. L. 92–463), the Centers for Disease Control and Prevention (CDC) announces the following Federal advisory committee meetings.

*Name:* National Vaccine Advisory Committee (NVAC).

*Times and Dates:* 9 a.m.–2 p.m., September 16, 1999; 8:30 a.m.–12:45 p.m., September 17, 1999.

*Place:* Hubert H. Humphrey Building, Room 505A, 200 Independence Avenue, SW, Washington, DC 20201.

*Status:* Open to the public, limited only by the space available.

*Notice:* In the interest of security, the Department has instituted stringent procedures for entrance to the Hubert H. Humphrey Building by non-government employees. Thus, persons without a government identification card should plan to arrive at the building each day either between 8 and 8:30 a.m. or 12:30 and 1 p.m. Entrance to the meeting at other times during the day cannot be assured.

*Purpose:* This committee advises and makes recommendations to the Director of the National Vaccine Program on matters related to the Program responsibilities.

*Matters To Be Discussed:* Agenda items will include: an update on the National Vaccine Program Office (NVPO) activities; a workshop report on thimerasol in vaccines; an update on rotavirus vaccine recommendations; an update on the Vaccine Safety and Communication Subcommittee; decision making and communication issues in vaccine safety; an update on the immunization registries initiative; reports from the Future Vaccines Subcommittee, Vaccine Safety and Communication Subcommittee and the Immunization Coverage Subcommittee; an update on pandemic preparedness planning; and discussions on immunization challenges in Mexico and Canada.

*Name:* Subcommittee on Future Vaccines.

*Time and Date:* 2:45 p.m.–5 p.m., September 16, 1999.

*Place:* Hubert H. Humphrey Building, Room 305A, 200 Independence Avenue, SW, Washington, DC 20201.

*Status:* Open to the public, limited only by the space available.

*Purpose:* This subcommittee develops policy options and guides national activities that lead to accelerated development, licensure, and the best use of new vaccines in the simplest possible immunization schedules.

*Matters To Be Discussed:* Agenda items will include discussions regarding the Institute of Medicine (IOM) report concerning cytomegalovirus vaccines and the IOM report concerning enteric bacteria vaccines.

*Name:* Subcommittee on Immunization Coverage.

*Time and Date:* 2:45 p.m.–5 p.m., September 16, 1999.

*Place:* Hubert H. Humphrey Building, Room 505A, 200 Independence Avenue, SW, Washington, DC 20201.

*Status:* Open to the public, limited only by the space available.

*Purpose:* This subcommittee will identify and propose solutions that provide a multifaceted and holistic approach to reducing barriers that result in low immunization coverage for children.

*Matters To Be Discussed:* Agenda items will include updates on the roll out process for the ''Strategies to Sustain Success'' initiative; the status of the paper on adult immunizations at non-traditional sites and standards for adult immunizations at non-

traditional sites; and, adolescent immunizations. The subcommittee will also discuss revisions of adult immunization standards in conjunction with the National Coalition for Adult Immunization; and assessment of vaccine coverage in view of recent vaccine safety issues.

*Name:* Subcommittee on Vaccine Safety and Communication.

*Time and Date:* 2:45 p.m.–5 p.m., September 16, 1999.

*Place:* Hubert H. Humphrey Building, Room 325A, 200 Independence Avenue, SW, Washington, DC 20201.

*Status:* Open to the public, limited only by the space available.

*Purpose:* This subcommittee reviews issues relevant to vaccine safety and adverse reactions to vaccines.

*Matters To Be Discussed:* Agenda items include a discussion on Thimerosal in vaccines, rotavirus vaccine and intussusception and the private governmental communications/response to acute vaccine safety issues.

Agenda items are subject to change as priorities dictate.

**CONTACT PERSON FOR MORE INFORMATION:** Gloria Sagar, Committee Management Specialist, NVPO, CDC, 1600 Clifton Road, NE, M/S A–11, Atlanta, Georgia 30333, telephone 404/639–4450.

The Director, Management Analysis and Services Office, has been delegated the authority to sign **Federal Register** notices pertaining to announcements of meetings and other committee management activities for both CDC and the Agency for Toxic Substances and Disease Registry.

Dated: August 24, 1999.

**John C. Burckhardt,**

*Acting Director, Management Analysis and Services Office, Centers for Disease Control and Prevention.*

[FR Doc. 99–22525 Filed 8–30–99; 8:45 am]

**BILLING CODE 4163–18–P**

## DEPARTMENT OF HEALTH AND HUMAN SERVICES

### Administration for Children and Families

### Program Enhancement Supplement of National Adoption Information Exchange

**AGENCY:** Administration for Children and Families (ACYF)—Children's Bureau, ACF, DHHS.

**ACTION:** Notice.

**SUMMARY:** Notice is hereby given concerning the intention of the Children's Bureau to award a noncompetitive supplement of the National Adoption Information Exchange (NAE) project of $500,000 for an additional year—not to extend beyond September 29, 2000 to the National Adoption Center, Philadelphia, Pennsylvania. This noncompetitive supplement will allow for the continuation of a current NAE in order to facilitate the adoption of children with special needs in preparation for full competition of an expanded effort in FY 2000.

On November 24, 1998, the President issued an Executive Memorandum to the Secretary of Health and Human Services, directing the Secretary to develop a plan to expand the use of the Internet as a tool for finding homes for children waiting to be adopted from the public child welfare system. The plan calls for an extensive consultation process with state social services agencies, courts, private agencies and other interested stakeholders, to identify and address important issues and strategies and build on promising existing efforts to create an effective national registry system.

The Administration on Children, Youth and Families is awarding this supplement until the expanded scope of the project has been established and all eligible organizations will have a fair and equitable opportunity to compete for the multiyear project. This will ensure that the project offered to the field for competition is the project described by the Executive Memorandum. It will also ensure that the successful applicant is the best qualified to perform the activities called for by the expanded scope.

**Authority:** The award will be made under the authority of the Adoption Opportunities Program, Title II of the Child Abuse Prevention and Treatment and Adoption Reform Act of 1978, as amended [42 U.S.C. 5111].

**FOR FURTHER INFORMATION CONTACT:** Geneva Ware-Rice, Children's Bureau, Administration on Children, Youth and Families, 330 C Street, SW, Room 2424, Washington, DC 20447; Telephone: (202) 205–8305.

Dated: August 25, 1999.

**Patricia Montoya,**

*Commissioner, Administration on Children, Youth and Families.*

[FR Doc. 99–22606 Filed 8–30–99; 8:45 am]

**BILLING CODE 4184–01–M**

## DEPARTMENT OF HEALTH AND HUMAN SERVICES

### Food and Drug Administration

**[Docket No. 99N–1522]**

### Agency Information Collection Activities; Submission for OMB Review; Comment Request; Temporary Marketing Permit Applications

**AGENCY:** Food and Drug Administration, HHS.

**ACTION:** Notice.

**SUMMARY:** The Food and Drug Administration (FDA) is announcing that the proposed collection of information listed below has been submitted to the Office of Management and Budget (OMB) for review and clearance under the Paperwork Reduction Act of 1995 (the PRA).

**DATES:** Submit written comments on the collection of information by September 30, 1999.

**ADDRESSES:** Submit written comments on the collection of information to the Office of Information and Regulatory Affairs, OMB, New Executive Office Bldg., 725 17th St. NW., rm. 10235, Washington, DC 20503, Attn: Wendy Taylor, Desk Officer for FDA.

**FOR FURTHER INFORMATION CONTACT:** Peggy Schlosburg, Office of Information Resources Management (HFA–250), Food and Drug Administration, 5600 Fishers Lane, Rockville, MD 20857, 301–827–1223.

**SUPPLEMENTARY INFORMATION:** In compliance with 44 U.S.C. 3507, FDA has submitted the following proposed collection of information to OMB for review and clearance.

### Temporary Marketing Permit Applications—21 CFR 130.17(c) and (I) (OMB Control Number 0910–0133)—Extension

Section 401 of the Federal Food, Drug, and Cosmetic Act (the act) (21 U.S.C. 341) directs FDA to issue regulations establishing definitions and standards of identity for food ''whenever * * * such action will promote honesty and fair dealing in the interest of consumers.'' Under section 403(g) of the act (21 U.S.C. 343(g)), a food that is subject to a definition and standard of identity prescribed by regulation is misbranded if it does not conform to such definition and standard of identity. Section 130.17 (21 CFR 130.17) provides for the issuance by FDA of temporary marketing permits that enable the food industry to test consumer acceptance and measure the technological and commercial feasibility in interstate