will discuss its ongoing project ethical and policy issues in international research. Some Commission members may participate by telephone conference. The meeting is open to the public and opportunities for statements by the public will be provided on January 18 from 1:00–1:30 pm.

Dates/times	Location
January 18, 2001, 8:30 am–5 pm.	Sheraton Premiere at Tysons Corner, 8661 Leesburg Pike, Tysons Corner/Vienna, Virginia 22182
January 19, 2001, 8 am–12 pm.	Same Location as Above

SUPPLEMENTARY INFORMATION: The President established the National Bioethics Advisory Commission (NBAC) on October 3, 1999 by Executive Order 12975 as amended. The mission of the NBAC is to advise and make recommendations to the National Science and Technology Council, its Chair, the President, and other entities on bioethical issues arising from the research on human biology and behavior, and from the applications of that research.

Public Participation

The meeting is open to the public with attendance limited by the availability of space on a first come, first serve basis. Members of the public who wish to present oral statements should contact Ms. Jody Crank by telephone, fax machine, or mail as shown below as soon as possible, at least 4 days before the meeting. The Chair will reserve time for presentations by persons requesting to speak and asks that oral statements be limited to five minutes. The order of persons wanting to make a statement will be assigned in the order in which requests are received. Individuals unable to make oral presentations can mail or fax their written comments to the NBAC staff office at least five business days prior to the meeting for distribution to the Commission and inclusion in the public record. The Commission also accepts general comments at its website at bioethics.gov. Persons needing special assistance, such as sign language interpretation or other special accommodations, should contact NBAC staff at the address or telephone number listed below as soon as possible.

FOR FURTHER INFORMATION CONTACT: Ms. Jody Crank, National Bioethics Advisory Commission, 6705 Rockledge Drive, Suite 700, Bethesda, Maryland 20892–7979, telephone (301) 402–4242, fax number (301) 480–6900.

Dated: November 29, 2000.

Eric M. Meslin,

Executive Director, National Bioethics Advisory Commission.

[FR Doc. 00–30871 Filed 12–4–00; 8:45 am]

BILLING CODE 4167–01–U

DEPARTMENT OF HEALTH AND HUMAN SERVICES

Centers for Disease Control and Prevention

Injury Research Grant Review Committee: Conference Call Meeting

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 conference call committee meeting.

Name: Injury Research Grant Review Committee (IRGRC).

Time and Date: 1 p.m.–1:30 p.m., December 19, 2000.

Place: National Center for Injury Prevention and Control (NCIPC), CDC, Koger Center, Vanderbilt Building, 1st Floor, Conference Room 1006, 2939 Flowers Road, South, Atlanta, Georgia 30341. (Exit Chamblee-Tucker Road off I–85.)

Status: Open: 1:00 p.m.–1:10 p.m., December 19, 2000;

Closed: 1:10 p.m.–1:30 p.m., December 19, 2000.

Purpose: This committee is charged with advising the Secretary of Health and Human Services, the Assistant Secretary for Health, and the Director, CDC, regarding the scientific merit and technical feasibility of grant applications received from academic institutions and other public and private profit and nonprofit organizations, including State and local government agencies, to conduct specific injury research that focuses on prevention and control and to support injury prevention research centers.

Matters To Be Discussed: Agenda items include the purpose of the meeting and discussion and vote on site visits to be conducted by IRGRC. Beginning at 1:10 p.m., through 1:30 p.m., December 19, the Committee will discuss and vote on the preliminary evaluation (triage) conducted by IRGRC to determine if a grant application submitted in response to Program Announcement #01007 is of sufficient scientific and technical merit to warrant further review by IRGRC. This portion of the meeting will be closed to the public in accordance with provisions set forth in section 552b(c)(4) and (6), title 5 U.S.C., and the Determination of the Associate Director for Management and Operations, CDC, pursuant to Pub. Law. 92–463.

This notice is published less than 15 days prior to the conference call due to administrative delay.

Agenda items are subject to change as priorities dictate.

Contact Person for More Information: Richard W. Sattin, M.D., Acting Executive Secretary, IRGRC, NCIPC, CDC, 4770 Buford Highway, NE, M/S K58, Atlanta, Georgia 30341–3724, telephone 770/488–4330.

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 the Centers for Disease Control and Prevention and the Agency for Toxic Substances and Disease Registry.

Dated: November 30, 2000.

Julia M. Fuller,

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

[FR Doc. 00–30995 Filed 12–1–00; 11:39 am]

BILLING CODE 4163–18–P

DEPARTMENT OF HEALTH AND HUMAN SERVICES

Administration for Children and Families

Notice of Allotment Percentages to States for Child Welfare Services State Grants

AGENCY: Administration for Children, Youth and Families, Administration for Children and Families, Department of Health and Human Services.

ACTION: Biennial publication of allotment percentages for States under the Title IV–B subpart 1, Child Welfare Services State Grants Program.

SUMMARY: As required by section 421(c) of the Social Security Act (42 U.S.C. 621(c)), the Department is publishing the allotment percentage for each State under the Title IV–B subpart 1, Child Welfare Services State Grants Program. Under section 421(a), the allotment percentages are one of the factors used in the computation of the Federal grants awarded under the Program.

EFFECTIVE DATE: The allotment percentages shall be effective for Fiscal Years 2002 and 2003.

FOR FURTHER INFORMATION CONTACT: Doris Lee, Office of Management Services, Administration for Children, Youth and Families, Administration for Children and Families, 330 C Street, SW., Washington, D.C. 20447.

SUPPLEMENTARY INFORMATION: The allotment percentage for each State is determined on the basis of paragraphs (b) and (c) of section 421 of the Act. These figures are available on the ACF homepage on the internet: *http://www.acf.dhhs.gov/programs/cb/*. The allotment percentage for each State is as follows:

State	Allotment percentage
Alabama	59.48
Alaska	48.91
Arizona	55.89
Arkansas	61.01
California	48.04
Colorado	45.34
Connecticut	31.28
Delaware	46.32
District of Columbia	30.47
Florida	50.80
Georgia	52.27
Hawaii	50.75
Idaho	59.88
Illinois	45.27
Indiana	54.08
Iowa	54.58
Kansas	52.98
Kentucky	59.13
Louisiana	59.38
Maine	56.91
Maryland	43.39
Massachusetts	38.66
Michigan	50.76
Minnesota	46.26
Mississippi	63.55
Missouri	53.49
Montana	61.15
Nebraska	52.59
Nevada	45.54
New Hampshire	45.92
New Jersey	37.46
New Mexico	61.30
New York	40.91
North Carolina	53.73
North Dakota	59.09
Ohio	52.15
Oklahoma	59.47
Oregon	52.39
Pennsylvania	49.81
Rhode Island	48.60
South Carolina	58.73
South Dakota	56.50
Tennessee	55.06
Texas	52.96
Utah	59.18
Vermont	54.70
Virginia	48.07
Washington	47.49
West Virginia	62.93
Wisconsin	52.03
Wyoming	54.03
American Samoa	70.00
Guam	70.00
N. Mariana Islands	70.00
Puerto Rico	70.00
Virgin Islands	70.00

Dated: November 27, 2000.

Patricia Montoya,

Commissioner, Administration for Children, Youth and Families.

[FR Doc. 00–30887 Filed 12–4–00; 8:45 am]

BILLING CODE 4184–01–P

DEPARTMENT OF HEALTH AND HUMAN SERVICES

Food and Drug Administration

[Docket No. 00N–1575]

Agency Information Collection Activities; Proposed Collection; Comment Request; Nutrition Labeling; Declaration of Caloric Amounts and Serving Sizes for Breath Mints

AGENCY: Food and Drug Administration, HHS.

ACTION: Notice.

SUMMARY: The Food and Drug Administration (FDA) is announcing an opportunity for public comment on the proposed collection of certain information by the agency. Under the Paperwork Reduction Act of 1995 (the PRA), Federal agencies are required to publish notice in the **Federal Register** concerning each proposed collection of information, including each proposed extension of an existing collection of information, and to allow 60 days for public comment in response to the notice. This notice solicits comments on information collection requirements regarding the nutrition labeling of breath mints.

DATES: Submit written or electronic comments on the collection of information by February 5, 2001.

ADDRESSES: Submit electronic comments on the collection of information to http://www.accessdata.fda.gov/scripts/oc/dockets/edockethome.cfm. Submit written comments on the collection of information to the Dockets Management Branch (HFA–305), Food and Drug Administration, 5630 Fishers Lane, rm. 1061, Rockville, MD 20852. All comments should be identified with the docket number found in brackets in the heading of this document.

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: Under the PRA (44 U.S.C. 3501–3520), Federal agencies must obtain approval from the Office of Management and Budget (OMB) for each collection of information they conduct or sponsor. ''Collection of information'' is defined in 44 U.S.C. 3502(3) and 5 CFR 1320.3(c) and includes agency requests or requirements that members of the public submit reports, keep records, or provide information to a third party. Section 3506(c)(2)(A) of the PRA (44 U.S.C. 3506(c)(2)(A)) requires Federal agencies to provide a 60-day notice in the **Federal Register** concerning each proposed collection of information, including each proposed extension of an existing collection of information, before submitting the collection to OMB for approval. To comply with this requirement, FDA is publishing notice of the proposed collection of information set forth in this document.

With respect to the following collection of information, FDA invites comments on: (1) Whether the proposed collection of information is necessary for the proper performance of FDA's functions, including whether the information will have practical utility; (2) the accuracy of FDA's estimate of the burden of the proposed collection of information, including the validity of the methodology and assumptions used; (3) ways to enhance the quality, utility, and clarity of the information to be collected; and (4) ways to minimize the burden of the collection of information on respondents, including through the use of automated collection techniques, when appropriate, and other forms of information technology.

Nutrition Labeling; Declaration of Caloric Amounts and Serving Sizes for Breath Mints—21 CFR 101.9(b) and 101.9(c)(1) (OMB Control Number 0910–0364)—Extension

Section 403(q) of the Federal Food, Drug, and Cosmetic Act (the act) (21 U.S.C. 343(q)) requires that the label or labeling of a food bear nutrition information, including information on: (1) The serving size and number of servings per container, and (2) the number of calories present in a serving of the food. Under FDA's nutrition labeling regulations in § 101.9(d)(3) (21 CFR 101.9(d)(3)), the nutrition facts panel of the food label must disclose the serving size of the food product and the number of servings in each package. Under § 101.9(c)(1), the nutrition facts panel must disclose the number of calories present in a serving of the food.

In the **Federal Register** of December 30, 1997 (62 FR 67775), FDA published a proposed rule to amend the nutrition labeling regulations by changing the label serving size for the product category ''Hard candies, breath mints'' to one unit. FDA proposed this change in response to a petition to provide a serving size for breath mints that more accurately reflects the amount customarily consumed per eating occasion. In a related issue, FDA also proposed to: (1) Modify the rounding rules for calories to allow the declaration of caloric amounts of less than 5 calories on the nutrition label,