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 on 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 2006 and 2007.

FOR FURTHER INFORMATION CONTACT: Doris Lee, Grants Fiscal Management Specialist, Office of Grants Management, Office of Administration, Administration for Children and Families, telephone (202) 205–4626.

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 home page on the Internet: *http://www.acf.dhhs.gov/programs/cb/*. The allotment percentage for each State is as follows:

State	Allotment percentage
Alabama	58.78
Alaska	47.24
Arizona	57.42
Arkansas	61.90
California	46.47
Colorado	45.19
Connecticut	31.13
Delaware	48.21
District of Columbia	30.00
Florida	51.94
Georgia	53.36
Hawaii	51.93
Idaho	58.99
Illinois	46.53
Indiana	54.64
Iowa	54.61
Kansas	53.08
Kentucky	58.79
Louisiana	59.22
Maine	54.86
Maryland	41.44
Massachusetts	36.69
Michigan	51.78
Minnesota	46.01
Mississippi	63.49
Missouri	53.96
Montana	59.82
Nebraska	52.37
Nevada	50.48
New Hampshire	44.95
New Jersey	36.08
New Mexico	60.09
New York	41.97
North Carolina	55.13
North Dakota	56.01
Ohio	52.85
Oklahoma	58.07
Oregon	53.45
Pennsylvania	49.80
Rhode Island	50.10
South Carolina	58.79
South Dakota	55.37
Tennessee	55.42
Texas	53.07
Utah	60.24
Vermont	51.92
Virginia	46.92
Washington	47.22
West Virginia	61.73
Wisconsin	51.48
Wyoming	49.48
American Samoa	70.00
Guam	70.00
N. Mariana Islands	70.00
Puerto Rico	70.00
Virgin Islands	70.00

Dated: October 20, 2004.

Joan E. Ohl,

Commissioner, Administration on Children, Youth and Families.

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DEPARTMENT OF HEALTH AND HUMAN SERVICES

Food and Drug Administration

[Docket No. 2004N–0269]

Agency Information Collection Activities; Submission for Office of Management and Budget Review; Comment Request; Radioactive Drug Research Committees

AGENCY: Food and Drug Administration, HHS.

ACTION: Notice.

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

DATES: Fax written comments on the collection of information by December 3, 2004.

ADDRESSES: OMB is still experiencing significant delays in the regular mail, including first class and express mail, and messenger deliveries are not being accepted. To ensure that comments on the information collection are received, OMB recommends that written comments be faxed to the Office of Information and Regulatory Affairs, OMB, Attn: Fumie Yokota, Desk Officer for FDA, FAX: 202–395–6974.

FOR FURTHER INFORMATION CONTACT: Karen L. Nelson, Office of Management Programs (HFA–250), Food and Drug Administration, 5600 Fishers Lane, Rockville, MD 20857, 301–827–1482.

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.

Radioactive Drug Research Committees—(OMB Control Number 0910–0053)

Under sections 201, 505, and 701 of the Federal Food, Drug, and Cosmetic Act (21 U.S.C. 321, 355, and 371), FDA has the authority to issue regulations governing the use of radioactive drugs for basic informational research. Section 361.1 (21 CFR 361.1) sets forth specific regulations regarding the establishment and composition of Radioactive Drug Research Committees and their role in approving and monitoring basic research studies utilizing radiopharmaceuticals. No basic research study involving any administration of a radioactive drug to research subjects is permitted without the authorization of an FDA approved Radioactive Drug Research Committee (§ 361.1(d)(7)). The type of research that may be undertaken with a radiopharmaceutical drug must be intended to obtain basic information and not to carry out a clinical trial. The types of basic research permitted are specified in the regulation, and include studies of metabolism, human physiology, pathophysiology, or biochemistry.

Section 361.1(c)(2) requires that each Radioactive Drug Research Committee shall select a chairman, who shall sign all applications, minutes, and reports of the committee. Each committee shall meet at least once each quarter in which research activity has been authorized or conducted. Minutes shall be kept and shall include the numerical results of votes on protocols involving use in human subjects. Under § 361.1(c)(3), each Radioactive Drug Research Committee shall submit an annual