Federal program using the poverty guidelines serves any of those jurisdictions, the Federal office that administers the program is generally responsible for deciding whether to use the contiguous-states-and-DC guidelines for those jurisdictions or to follow some other procedure.

Due to confusing legislative language dating back to 1972, the poverty guidelines have sometimes been mistakenly referred to as the ''OMB'' (Office of Management and Budget) poverty guidelines or poverty line. In fact, OMB has never issued the guidelines; the guidelines are issued each year by the Department of Health and Human Services. The poverty guidelines may be formally referenced as ''the poverty guidelines updated periodically in the **Federal Register** by the U.S. Department of Health and Human Services under the authority of 42 U.S.C. 9902(2).''

Some programs use a percentage multiple of the guidelines (for example, 125 percent or 185 percent of the guidelines), as noted in relevant authorizing legislation or program regulations. Non-Federal organizations that use the poverty guidelines under their own authority in non-federally-funded activities can choose to use a percentage multiple of the guidelines such as 125 percent or 185 percent.

The poverty guidelines do not make a distinction between farm and non-farm families, or between aged and non-aged units. (Only the Census Bureau poverty thresholds have separate figures for aged and non-aged one-person and two-person units.)

Note that this notice does not provide definitions of such terms as ''income'' or ''family.'' This is because there is considerable variation in how different programs that use the guidelines define these terms, traceable to the different laws and regulations that govern the various programs. Therefore, questions about how a particular program applies the poverty guidelines (*e.g.*, Is income before or after taxes? Should a particular type of income be counted? Should a particular person be counted in the family or household unit?) should be directed to the organization that administers the program.

Dated: January 17, 2007.

Michael O. Leavitt,

Secretary of Health and Human Services.

[FR Doc. 07–268 Filed 1–19–07; 8:45 am]

BILLING CODE 4151–05–P

DEPARTMENT OF HEALTH AND HUMAN SERVICES

Centers for Disease Control and Prevention

Disease, Disability, and Injury Prevention and Control Special Emphasis Panel (SEP): NIOSH Occupational Health and Safety Research, Program Announcement Number (PAR) 06–484

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 aforementioned meeting:

Time and Date: 8 a.m.–5 p.m., February 9, 2007 (Closed).

Place: 1750 New York Avenue, NW., Washington, DC 20006.

Status: 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 Director, Management Analysis and Services Office, CDC, pursuant to Public Law 92–463.

Matters To Be Discussed: The SEP meeting will include the review, discussion, and evaluation of applications received in response to ''NIOSH Occupational Health and Safety Research,'' PAR 06–484. The applications being reviewed include information of a confidential nature, including personal information concerning individuals associated with the applications.

Contact Person for More Information: Horace M. Stiles, DDS, PhD, MPH, Designated Federal Officer, 15111 Farm Market Road, Maypearl, Texas 76064–1902, telephone 404.498.2584.

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: January 18, 2007.

Elaine L. Baker,

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

[FR Doc. E7–987 Filed 1–23–07; 8:45 am]

BILLING CODE 4163–18–P

DEPARTMENT OF HEALTH AND HUMAN SERVICES

Administration for Children and Families

Submission for OMB Review; Comment Request

Title: 45 CFR 1304 Head Start Program Performance Standards.

OMB No. 0970–0148.

Description: Head Start Program Performance Standards require Head Start and Early Head Start Programs and Delegate Agencies to maintain program records. The Administration for Children and Families, Office of Head Start, is proposing to renew, without changes, the authority to require certain record keeping in all programs as provided for in 45 CFR part 1304 Head Start Program Performance Standards. These standards prescribe the services that Head Start and Early Head Start programs provide to enrolled children and their families.

Respondents: Head Start and Early Head Start grantees and delegate agencies.

ANNUAL BURDEN ESTIMATES

Instrument	Number of respondents	Number of responses per respondent	Average burden hours per response	Total burden hours
Standard	2,590	16	41.8	1,732,192
Estimated Total Annual Burden Hours:				1,732,192

Additional Information: Copies of the proposed collection may be obtained by writing to the Administration for Children and Families, Office of Administration, Office of Information Services, 370 L'Enfant Promenade, SW., Washington, DC 20447, Attn: ACF Reports Clearance Officer. All requests should be identified by the title of the information collection. E-mail address: *infocollection@acf.hhs.gov.*

OMB Comment: OMB is required to make a decision concerning the collection of information between 30 and 60 days after publication of this document in the **Federal Register**. Therefore, a comment is best assured of having its full effect if OMB receives it within 30 days of publication. Written comments and recommendations for the proposed information collection should be sent directly to the following: Office of Management and Budget, Paperwork

Reduction Project, Attn: Desk Officer for ACF, Fax: 202–395–6974.

Dated: January 18, 2007.

Robert Sargis,

Reports Clearance Officer.

[FR Doc. 07–272 Filed 1–23–07; 8:45 am]

BILLING CODE 4184–01–M

DEPARTMENT OF HEALTH AND HUMAN SERVICES

Administration for Children and Families

Submission for OMB Review; Comment Request

Title: Annual Maintenance-of-Effort (MOE) Report.

OMB No. 0970–0248.

Description: The Administration for Children and Families (ACF) is requesting a three-year extension of the ACF–204 (Annual MOE Report). The report is used to collect descriptive program characteristics information on the programs operated by States and Territories in association with their Temporary Assistance for Needy Families (TANF) programs. All State and Territory expenditures claimed toward States' and Territories' MOE requirements must be appropriate, i.e., meet all applicable MOE requirements. The Annual MOE Report provides the ability to learn about and to monitor the nature of State and Territory expenditures used to meet State's and Territories' MOE requirements, and it is an important source of information about the different ways that States and Territories are using their resources to help families attain and maintain self-sufficiency.

In addition, the report is used to obtain State and Territory program characteristics for ACF's annual report to Congress, and the report serves as a useful resource to use in Congressional hearings about how TANF programs are evolving, in assessing State the Territory MOE expenditures, and in assessing the need for legislative changes.

Respondents: The 50 States of the United States, the District of Columbia, Guam, Puerto Rico, and the Virgin Islands.

ANNUAL BURDEN ESTIMATES

Instrument	Number of respondents	Number of responses per respondent	Average burden hours per response	Total burden hours
ACF–204	54	1	128	6,912
Estimated Total Annual Burden Hours:				6,912

OMB Comment: OMB required to make a decision concerning the collection of information between 30 and 60 days after publication of this document in the **Federal Register**. Therefore, a comment is best assured of having its full effect if OMB receives it within 30 days of publication. Written comments and recommendations for the proposed information collection should be sent directly to the following: Office of Management and Budget, Paperwork Reduction Project, Attn: Desk Officer for ACF, Fax: 202–395–6974.

Dated: September 18, 2007.

Robert Sargis,

Reports Clearance Officer.

[FR Doc. 07–273 Filed 1–23–07; 8:45 am]

BILLING CODE 4184–01–M

DEPARTMENT OF HEALTH AND HUMAN SERVICES

Food and Drug Administration

[Docket No. 2007D–0017]

Guidance for Industry: Certain Human Cells, Tissues, and Cellular and Tissue-Based Products Recovered From Donors Who Were Tested for Communicable Diseases Using Pooled Specimens or Diagnostic Tests; Availability

AGENCY: Food and Drug Administration, HHS.

ACTION: Notice.

SUMMARY: The Food and Drug Administration (FDA) is announcing the availability of a document entitled ''Guidance for Industry: Certain Human Cells, Tissues, and Cellular and Tissue-Based Products (HCT/Ps) Recovered From Donors Who Were Tested for Communicable Diseases Using Pooled Specimens or Diagnostic Tests'' dated January 2007. The guidance document provides establishments that make HCT/P donor eligibility determinations with recommendations concerning the donor eligibility requirements contained in 21 CFR part 1271, subpart C, which became effective on May 25, 2005. The guidance applies only to certain HCT/Ps that were not regulated as HCT/Ps before May 25, 2005, and that were recovered from donors beginning on or after the May 25, 2005, and within 30 days of the date of publication of this document in the **Federal Register**. This guidance has an immediate implementation date because FDA has determined that prior public participation is not feasible or appropriate. In certain cases, donor retesting needs to be initiated quickly, and the availability of certain HCT/Ps may be critical to their intended recipients.

DATES: Submit written or electronic comments on agency guidances at any time.

ADDRESSES: Submit written requests for single copies of the guidance to the Office of Communication, Training, and Manufacturers Assistance (HFM–40), Center for Biologics Evaluation and Research (CBER), Food and Drug Administration, 1401 Rockville Pike, suite 200N, Rockville, MD 20852–1448. Send one self-addressed adhesive label to assist the office in processing your requests. The guidance may also be obtained by mail by calling CBER at 1–800–835–4709 or 301–827–1800. See the **SUPPLEMENTARY INFORMATION** section for electronic access to the guidance document.

Submit written comments on the guidance to the Division of Dockets Management (HFA–305), Food and Drug Administration, 5630 Fishers Lane, rm. 1061, Rockville, MD 20852. Submit electronic comments to *http://www.fda.gov/dockets/ecomments.*

FOR FURTHER INFORMATION CONTACT: Valerie A. Butler, Center for Biologics Evaluation and Research (HFM–17), Food and Drug Administration, 1401 Rockville Pike, suite 200N, Rockville, MD 20852–1448, 301–827–6210.

SUPPLEMENTARY INFORMATION:

I. Background

FDA is announcing the availability of a document entitled ''Guidance for Industry: Certain Human Cells, Tissues, and Cellular and Tissue-Based Products (HCT/Ps) Recovered From Donors Who