getdoc.cgi?dbname=2002__register&docid=02–19871-filed.pdf), soliciting comments on its proposal to recommend approval on HHS support for the research protocol entitled ''Precursors to Diabetes in Japanese American Youth'' (1 R01 DK59234–01). The comment period closed on August 21, 2002. OHRP hereby gives notice that the comment period is being reopened for 30 days, and additional information regarding the protocol is being made available.

OHRP received a number of comments in response to the August 7, 2002, notice. Several who commented stated that (1) the 14-day comment period was too short, and (2) the information about the research protocol that OHRP provided to the public was insufficient to allow for meaningful comment about whether the research was appropriate for HHS to support. In response to these comments, OHRP is reopening the public comment period for 30 days. OHRP also is making available additional information regarding the research protocol, namely: the protocol application reviewed by the Children's Hospital and Regional Medical Center (Seattle) IRB; the assent form; the consent form; and selected parts of the grant application, including the abstract, specific aims, background and significance, discussion of the involvement of human subjects, and literature cited. These materials are available for review on the OHRP Web page at (*http://ohrp.osophs.dhhs.gov/pdjay/pdjayindex.htm*). A paper copy of the information referenced here is available upon request.

FOR FURTHER INFORMATION CONTACT: Requests for additional information about the research proposal may be submitted under the Freedom of Information Act (FOIA). Such requests should be directed to: Ms. Darlene Christian, PHS FOIA Office, Parklawn Building, Room 17A–46, 5600 Fishers Lane, Rockville, MD 20857; telephone (301) 443–5252; fax (301) 443–0925.

Dated: December 12, 2002.

Eve E. Slater,

Assistant Secretary for Health.

[FR Doc. 02–31848 Filed 12–17–02; 8:45 am]

BILLING CODE 4150–28–M

DEPARTMENT OF HEALTH AND HUMAN SERVICES

Centers for Disease Control and Prevention

[30DAY–07–03]

Agency Forms Undergoing Paperwork Reduction Act Review

The Centers for Disease Control and Prevention (CDC) publishes a list of information collection requests under review by the Office of Management and Budget (OMB) in compliance with the Paperwork Reduction Act (44 U.S.C. chapter 35). To request a copy of these requests, call the CDC Reports Clearance Officer at (404) 498–1210. Send written comments to CDC, Desk Officer, Human Resources and Housing Branch, New Executive Office Building, Room 10235, Washington, DC 20503. Written comments should be received within 30 days of this notice.

Proposed Project: Requirement for a Special Permit to Import Cynomolgus, African Green, or Rhesus Monkeys into the United States (0920–0263)—Extension—National Center for Infectious Diseases (NCID), Centers for Disease Control and Prevention (CDC). To receive a special permit to import cynomolgus, African green and/or rhesus monkeys, a registered importer of nonhuman primates must submit to the Director, CDC, a written plan which specifies the steps that will be taken to prevent exposure of persons and animals during the entire importation and quarantine process for the arriving nonhuman primates.

Under the special permit arrangement, registered importers must submit a plan to CDC for the importation and quarantine if they wish to import the specific monkeys covered. The plan must address disease prevention procedures to be carried out in every step of the chain of custody of such monkeys, from embarkation in the country of origin to release from quarantine. Information such as species, origin and intended use for monkeys, transit information, isolation and quarantine procedures, and procedures for testing of quarantined animals is necessary for CDC to make public health decisions. This information enables CDC to evaluate compliance with the standards and determine whether the measures being taken to prevent exposure of persons and animals during importation are adequate. Once CDC is assured, through the monitoring of shipments (normally no more than 2), that the provisions of a special permit plan are being followed by a new permit holder and that the use of adequate disease control practices is being demonstrated, the special permit is extended to cover the receipt of additional shipments under the same plan for a period of 180 days, and may be renewed upon request. This eliminates the burden on importers to repeatedly report identical information, requiring only that specific shipment itineraries and information on changes to the plan which require approval be submitted.

Respondents are commercial or not-for-profit importers of nonhuman primates. The burden represents full submission of information and itinerary/change information respectively. The burden hours are estimated to be approximately 20.

Respondents	Number of respondents	Number of responses per respondents	Avg. burden responses (in hrs.)
Businesses	2	5	30/60
	3	5	10/60
Organizations (limited permit)	15	5	10/60

Dated: December 10, 2002.

Nancy Cheal,

Acting Associate Director for Policy, Planning and Evaluation, Centers for Disease Control and Prevention.

[FR Doc. 02–31748 Filed 12–17–02; 8:45 am]

BILLING CODE 4163–18–P

DEPARTMENT OF HEALTH AND HUMAN SERVICES

Centers for Disease Control and Prevention

[30DAY–09–03]

Agency Forms Undergoing Paperwork Reduction Act Review

The Centers for Disease Control and Prevention (CDC) publishes a list of information collection requests under review by the Office of Management and Budget (OMB) in compliance with the Paperwork Reduction Act (44 U.S.C. chapter 35). To request a copy of these requests, call the CDC Reports Clearance Officer at (404) 498–1210. Send written comments to CDC, Desk Officer, Human Resources and Housing Branch, New Executive Office Building, Room 10235, Washington, DC 20503. Written comments should be received within 30 days of this notice.

Proposed Project: PHS Supplements to the Application for Federal Assistance SF–424 (0920–0428)—Extension—Office of the Director (OD), Centers for Disease Control and Prevention (CDC) is requesting a three-year extension for continued use of the Supplements to the Request for Federal Assistance Application, SF–424. The Checklist, Program Narrative, and the Public Health System Impact Statement (PHSIS, third party notification form) are a part of the standard application for State and local governments and for private non-profit and for-profit organizations when applying for financial assistance from PHS grant programs. The Checklist assists applicants to ensure that they have included all required information necessary to process the application. The Checklist data helps to reduce the time required to process and review grant applications, expediting the issuance of grant awards. The PHSIS Third Party Notification Form is used to inform State and local health agencies of community-based proposals submitted by non-governmental applicants for Federal funding. The total annualized estimated burden is 42,695 hours.

Respondents	Number of respondents	Number of responses/respondent	Avg. burden/response (in hrs.)
State and local health departments; non-profit and for-profit organizations	7,457	1	5.73

Dated: December 10, 2002.

Nancy E. Cheal,

Acting Associate Director for Policy, Planning and Evaluation, Centers for Disease Control and Prevention (CDC).

[FR Doc. 02–31749 Filed 12–17–02; 8:45 am]

BILLING CODE 4163–18–P

DEPARTMENT OF HEALTH AND HUMAN SERVICES

Centers for Disease Control and Prevention

Interagency Committee on Smoking and Health: Meeting

In accordance with section 10(a)(2) of the Federal Advisory Committee Act (Pub L. 92–463), the National Center for Chronic Disease Prevention and Health Promotion (NCCDPHP) of the Centers for Disease Control and Prevention (CDC) announces the following meeting:

Name: Interagency Committee on Smoking and Health Cessation Subcommittee.

Date and Time: January 16, 2003; 8:30 a.m.–2:30 p.m.

Place: Room 705A, Hubert H. Humphrey Building, 200 Independence Avenue, SW., 7th Floor, Washington, DC 20201.

Status: Open to the public, limited only by the space available. Those who wish to attend are encouraged to register with the contact person listed below. If you will require a sign language interpreter, or have other special needs, please notify the contact person by 4:30 E.S.T. on January 14, 2003.

Purpose: The Interagency Committee on Smoking and Health advises the Secretary, Department of Health and Human Services, and the Assistant Secretary for Health in the: (a) Coordination of all research and education programs and other activities within the Department and with other federal, state, local and private agencies, and (b) establishment and maintenance of liaison with appropriate private entities, federal agencies, and state and local public health agencies with respect to smoking and health activities.

Matters to be discussed: The agenda will focus on developing a plan of action to promote tobacco use cessation to be presented to the Secretary of Health and Human Services.

Contact Person for More Information: Substantive program information as well as summaries of the meeting and roster of committee members may be obtained from the Internet at *http://www.cdc.gov/tobacco* in mid-March 2003, or from Ms. Monica L. Swann, Committee Management Specialist, Interagency Committee on Smoking and Health, Office on Smoking and Health, NCCDPHP, CDC, 200 Independence Avenue, SW., Room 317B, Washington, DC, 20201, telephone (202) 205–8500.

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: December 10, 2002.

Joseph E. Salter,

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

[FR Doc. 02–31774 Filed 12–17–02; 8:45 am]

BILLING CODE 4163–18–P

DEPARTMENT OF HEALTH AND HUMAN SERVICES

Administration for Children and Families

Proposed Information Collection Activity; Comment Request

Title: Child Care and Development Fund Plan for States/Territories.

OMB No.: 0970–0114.

Description: The Child Care and Development Fund (CCDF) Plan for States and Territories is required from the Child Care Lead Agency by section 658E of the Child Care and Development Block Grant Act of 1990 (Pub. L. 101–508), 42 U.S.C. 9858). The implementing regulations for the statutorily required Plan are at 45 CFR 98.10 through 98.18. The Plan, submitted on the ACF–118, is required biennially and remains in effect for two years. This Plan provides ACF and the public with a description of, and assurance about, the State's child care