SUPPLEMENTARY INFORMATION:

A. Purpose

The General Services Administration will be requesting that the Office of Management and Budget (OMB) review and approve information collection, 3090–0121, concerning industrial funding fee and sales reporting. The information is used primarily by contracting officers to estimate requirements for the subsequent year, evaluate the effectiveness of a schedule, negotiate better prices based on volume and for special reports.

B. Annual Reporting Burden

Respondents: 15,710
Responses Per Respondent: 20
Total Responses: 314,200
Hours Per Response: .0833
Total Burden Hours: 26,173
OBTAINING COPIES OF PROPOSALS: Requesters may obtain a copy of the information collection documents from the General Services Administration, Regulatory Secretariat (VR), 1800 F Street, NW., Room 4035, Washington, DC 20405, telephone (202) 208–7312. Please cite OMB Control No. 3090–0121, Industrial Funding Fee and Sales Reporting, in all correspondence.

Dated: July 26, 2004

RALPH DESTEFANO

(Acting) Director, Contract Policy Division

[FR Doc. 04–17454 Filed 7–30–04; 8:45 am]

BILLING CODE 6820–61–S

DEPARTMENT OF HEALTH AND HUMAN SERVICES

Centers for Disease Control and Prevention

[Program Announcement 04276]

Expansion of HIV/AIDS Surveillance, Monitoring and Evaluation, and Information Management Activities in the Republic of Honduras; Notice of Intent To Fund Single Eligibility Award

A. Purpose

The Centers for Disease Control and Prevention (CDC) announces the intent to fund fiscal year (FY) 2004 funds for a cooperative agreement program to increase the capacity, quality and coverage of HIV/AIDS-related Strategic Information activities undertaken by the Ministry of Health (MOH) as cornerstone components of an expanded national response to HIV/AIDS targeting highly vulnerable populations (HVPs) in the Republic of Honduras. In the context of Honduras, HVPs include prostitutes, men who have sex with men (MSM), persons living with HIV/AIDS (PLWHAs), prisoners, and members of the Garifuna and other ethnic groups. Strategic Information is defined as: programs and activities supporting the implementation of first and second generation epidemiological surveillance survey activities; systems for monitoring and evaluation of the impact of the multi-sectoral national response to HIV/AIDS; and strategic initiatives to improve infrastructure and systems supporting surveillance, prevention, care and treatment, laboratory and information management activities.

The Catalog of Federal Domestic Assistance number for this program is 93.941.

B. Eligible Applicant

Assistance will be provided only to the Ministry of Health (MOH) of the Republic of Honduras.

The Honduras MOH is charged by national law to oversee the national response to health problems that threaten the well being of the country's citizens, including HIV/AIDS. The MOH, as the entity responsible for public health in Honduras, has direct responsibility for overseeing surveillance and the monitoring and evaluation of the national response to HIV/AIDS and HIV-related conditions in the country. The MOH of Honduras has collaborated with HHS/CDC and USAID in the past, including collaborations related to the evaluation of surveillance and laboratory systems related to HIV/AIDS in 1999–2000.

C. Funding

Approximately $300,000 is available in FY 2004 to fund this award. It is expected that the award will begin on or before September 1, 2004, and will be made for a 12-month budget period within a project period of up to 3 years. Funding estimates may change.

D. Where To Obtain Additional Information

For general comments or questions about this announcement, contact: Technical Information Management, CDC Procurement and Grants Office, 2920 Brandywine Road, Atlanta, GA 30341–4146, Telephone: (770) 488–2700.

For technical questions about this program, contact: Edgar Monterroso/Mark Fussell, Co-Project Officers, HHS/CDC AE Guatemala Unit 3321, APO AA 34024, Telephone: (502) 369–0791, Ext 515, E-mail: *em2z@cdc.gov* or *mfzz@cdc.gov*.

Dated: July 27, 2004.

William P. Nichols, MPA,

Acting Director, Procurement and Grants Office, Centers for Disease Control and Prevention.

[FR Doc. 04–17483 Filed 7–30–04; 8:45 am]

BILLING CODE 4163–18–P

DEPARTMENT OF HEALTH AND HUMAN SERVICES

Administration for Children and Families

Agency Recordkeeping/Reporting Requirements Under Emergency Review by the Office of Management and Budget (OMB)

Title: Fourth National Incidence Study of Child Abuse and Neglect (NIS–4).

OMB No.: New Request.

Description: The Department of Health and Human Services intends to issue letters to recruit agencies for participation in the next National Incidence Study of Child Abuse and Neglect (NIS). This will be the fourth cycle of this periodic study. The NIS–1, mandated under Public Law (Pub. L.) 93–247 (1974), was conducted in 1979 and 1980 and reported in 1981. The NIS–2 was mandated under Pub. L. 98–457 (1984), conducted in 1986 and 1987, and reported in 1988. The NIS–3 was mandated under both the Child Abuse Prevention, Adoption and Family Services Act of 1988 (Pub. L. 100–294) and the Child Abuse, Domestic Violence, Adoption and Family Services Act of 1992 (Pub. L. 102–295), conducted between 1993 and 1995, and published in 1996. The NIS–4 is mandated by the Keeping Children and Families Safe Act of 2003 (Pub. L. 108–36).

The NIS is unique in that it goes beyond the abused and neglected children who come to the attention of the Child Protective Services (CPS) system. In contrast to the National Child Abuse and Neglect Data Systems (NCANDS), which rely solely on reported cases, the NIS design assumes that reported children represent only a portion of the children who actually are maltreated. Following the implications of its assumption, the NIS estimates the scope of the maltreated child population by combining information about reported cases with data on maltreated children identified by professionals (called ''sentinels'') who encountered them during the normal course of their work in a wide range of agencies in representative communities. These professionals are asked to remain on the lookout for children they believe are

maltreated during the study reference period and to provide information about those children. Children identified by sentinels and those who alleged maltreatment is investigated by CPS during the same period are evaluated against standardized definitions and only children who meet the study standards are used to develop the study estimates. The study estimates are couched in terms of numbers of maltreated children, with data unduplicated so a given child is counted only once. Confidentiality of all participants is carefully protected.

A nationally representative sample of 120 counties will be selected and all local child protective service (CPS) agencies serving the selected counties will be identified. Plans will be developed to obtain data on cases investigated during the study reference period, September 4 to December 3, 2005. Sentinels in the selected counties will be identified through samples of agencies in 11 categories: county juvenile probation departments, sheriff (and/or state police) departments, public health departments, public housing departments, municipal police departments, hospitals, schools, day care centers, social service agencies, mental health agencies, and shelters for battered women or runaway/homeless youth. A total of approximately 1,600 sentinel agencies will be sampled. Plans will be developed to identify staff in these agencies who have direct contact with children to serve as sentinels during the study by submitting data on maltreated children they encounter during the study referenced period. in preparation for the study, letters will be sent to the directors of the selected agencies asking them to permit their agencies to participate in the NIS–4, and describing the general nature of the data collection effort. DHHS will issue subsequent notice of proposed data collection for this study after data collection plans are developed.

Respondents:

ANNUAL BURDEN ESTIMATES

Instrument	Number of respondents	Number of responses per respondent	Average burden hours per response	Total burden hours
Letters to CPS Agencies	120	1	.20	24
Letter to Sentinel Agencies	1600	1	.20	320
Letter to Sentinels	12000	1	.20	2400
Estimated Total Annual Burden Hours:	.....	.....	.20	2744

Additional Information: ACF is requesting that OMB grant a 180 day approval for this information collection under procedures for emergency processing by August 15, 2004. A copy of this information collection, with applicable supporting documentation, may be obtained by calling the Administration for Children and Families, Mary Bruce Webb at (202) 205–8628. In addition, a request may be made by sending an e-mail request to: *mbwebb@acf.hhs.gov.*

Comments and questions about the information collection described above should be directed to the following address by August 15, 2004: Office of Information and Regulatory Affairs, Attn: OMB Desk Officer for ACF, Office of Management and Budget, Paper Reduction Project, 725 17th Street, NW., Washington, DC 20503. E-mail address: *Katherine _T._Astrich@omb. eop. gov.*

Dated: July 26, 2004.

Robert Sargis,

Reports Clearance Officer.

[FR Doc. 04–17453 Filed 7–30–04; 8:45 am]

BILLING CODE 4184–01–M

DEPARTMENT OF HEALTH AND HUMAN SERVICES

Food and Drug Administration

Establishment of Animal Drug User Fee Rates and Payment Procedures for Fiscal Year 2005

AGENCY: Food and Drug Administration, HHS.

ACTION: Notice.

SUMMARY: The Food and Drug Administration (FDA) is announcing the rates and payment procedures for fiscal year (FY) 2005 animal drug user fees. The Federal Food, Drug, and Cosmetic Act (the act), as amended by the Animal Drug User Fee Act of 2003 (ADUFA), Public Law 108–130, authorizes FDA to collect user fees for certain animal drug applications, on certain animal drug products, on certain establishments where such products are made, and on certain sponsors of such animal drug applications and/or investigational animal drug submissions. This notice establishes the fee rates for FY 2005.

For FY 2005, the animal drug user fee rates are: $119,300 for an animal drug application; $59,650 for a supplemental animal drug application for which safety or effectiveness data is required; $3,085 for an annual product fee; $42,600 for an annual establishment fee; and $32,150 for an annual sponsor fee. FDA will issue invoices for FY 2005 product, establishment and sponsor fees by December 30, 2004, and these invoices will be due and payable by January 31, 2005.

The application fee rates are effective for applications submitted on or after October 1, 2004, and will remain in effect through September 30, 2005. Applications will not be accepted to review until FDA has received full payment of application fees and any other animal drug user fees owed.

FOR FURTHER INFORMATION CONTACT: Visit FDA's Web site at: *http://www.fda.gov/oc/adufa* or contact Robert Miller, Center for Veterinary Medicine (HFV–10), Food and Drug Administration, 7519 Standish Pl., Rockville, MD 20855, 301–827–5436. For general questions, you may also e-mail the Center for Veterinary Medicine (CVM) at: *cvmadufa@fda.gov.*

SUPPLEMENTARY INFORMATION:

I. Background

Section 740 of the act (21 U.S.C. 379j–12) establishes four different kinds of user fees: (1) Fees for certain types of animal drug applications and supplements, (2) annual fees for certain animal drug products, (3) annual fees for certain establishments where such products are made, and (4) annual fees for certain sponsors of animal drug applications and/or investigational animal drug submissions. (See 21 U.S.C.