DEPARTMENT OF HEALTH AND HUMAN SERVICES

Administration for Children and Families

Submission for OMB Review; Comment Request

Title: April 2004 Current Population Survey Supplement on Child Support.

OMB No.: 0992–0003.

Description: Collection of the data will assist legislators and policymakers in determining how effective their policymaking efforts have been over time in applying the various child support legislation to the overall child support enforcement picture. This information will help policymakers determine to what extent individuals on welfare would be able to leave the welfare rolls as a result of more stringent child support enforcement efforts.

Respondents: Individuals and households.

Annual Burden Estimates:

Instrument	Number of respondents	Number of responses per respondent	Average burden hours per response	Total burden hours
Child Support Survey	47,000	1	.0246	1156
Estimated Total Annual Burden Hours:				1156

Additional Information: Copies of the proposed collection may be obtained by writing to the Administration for Children and Families, 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: *rsargis@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, E-mail address: *lauren_wittenberg@omb.eop.gov*.

Dated: January 8, 2003.

Robert Sargis,

Reports Clearance Office.

[FR Doc. 04–775 Filed 1–13–04; 8:45 am]

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DEPARTMENT OF HEALTH AND HUMAN SERVICES

Food and Drug Administration

Animal Drug User Fee Act of 2003; Interim Procedures

AGENCY: Food and Drug Administration, HHS.

ACTION: Notice.

SUMMARY: The Food and Drug Administration (FDA) is announcing interim procedures relating to the Animal Drug User Fee Act (ADUFA) of 2003, which was signed by the President on November 18, 2003. This act amends the Federal Food, Drug, and Cosmetic Act, and authorizes FDA to collect four types of user fees: Application fees, establishment fees, product fees, and sponsor fees. Before FDA can begin collecting these fees, enabling appropriations must be enacted. Until further notice, such fees should not be submitted to FDA. However, sponsors should continue to submit new animal drug applications as in the past until additional direction is provided. Certain types of applications submitted on or after September 1, 2003, will be subject to fees, but an invoice for those fees will not be issued until after enabling appropriations are enacted. FDA will publish another **Federal Register** notice specifying fee amounts and procedures for submitting payments.

ADDRESSES: Visit the FDA Web site that provides further information on ADUFA at: *http://www.fda.gov/oc/adufa.*

FOR FURTHER INFORMATION CONTACT: Robert Miller, Center for Veterinary Medicine (HFV–10), Food and Drug Administration, 7519 Standish Pl., Rockville, MD 20855, 301–827–5436; e-mail: *rmiller2@cvm.fda.gov*. For general questions, you may also contact the Center for Veterinary Medicine at: *mailto:cvmadufa@fda.gov.*

SUPPLEMENTARY INFORMATION: ADUFA authorizes FDA to collect fees for: (1) Certain types of animal drug applications and supplemental animal drug applications submitted on or after September 1, 2003, (2) certain animal drug products, (3) certain establishments where such products are manufactured in final dosage form, and (4) certain sponsors of animal drug applications or investigational animal drug submissions. However, FDA may not begin to collect these fees until enabling appropriations are enacted. After the enactment of enabling appropriations, FDA will publish a **Federal Register** notice with detailed payment procedures.

For FY 2004 through FY 2008, ADUFA establishes overall fee revenue amounts for application fees, establishment fees, product fees, and sponsor fees. Revenue amounts established for years after FY 2004 are subject to annual adjustments for inflation and workload. Fees for applications, establishments, products, and sponsors are to be established each year by FDA so that revenues will approximate the levels established in the statute, after those amounts have been first adjusted for inflation and workload. FDA will publish a **Federal Register** notice with the FY 2004 fee rates and detailed payment instructions.

In an effort to better ensure broad awareness of interim procedures relating to ADUFA, FDA has established a Web site that provides further information at *http://www.fda.gov/oc/adufa*.

Dated: January 7, 2004.

Jeffrey Shuren,

Assistant Commissioner for Policy.

[FR Doc. 04–812 Filed 1–13–04; 8:45 am]

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DEPARTMENT OF HEALTH AND HUMAN SERVICES

Food and Drug Administration

[Docket Number 2003D–0558]

Draft Compliance Policy Guide, Guidance Levels for Radionuclides in Domestic and Imported Foods, Availability; and Draft Supporting Document, Supporting Document for Guidance Levels for Radionuclides in Domestic and Imported Foods, Availability

AGENCY: Food and Drug Administration, HHS.