Estimated Total Annual Burden Hours: 297.

In compliance with the requirements of Section 3506(c)(2)(A) of the Paperwork Reduction Act of 1995, the Administration for Children and Families is soliciting public comment on the specific aspects of the information collection described above. Copies of the proposed collection of information can be obtained and comments may be forwarded 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. E-mail address: *infocollection@acf.hhs.gov.* All requests should be identified by the title of the information collection.

The Department specifically requests comments on: (a) Whether the proposed collection of information is necessary for proper performance of the functions of the agency, including whether the information shall have practical utility; (b) the accuracy of the agency's estimate of the burden of the proposed collection of information; (c) the quality, utility, and clarity of the information to be collected; and (d) ways to minimize the burden of the collection of information on respondents, including through the use of automated collection techniques or other forms of information technology. Consideration will be given to comments and suggestions submitted within 60 days of this publication.

Dated: October 17, 2005.

Robert Sargis,

Reports Clearance Officer.

[FR Doc. 05–21163 Filed 10–21–05; 8:45 am]

BILLING CODE 4184–01–M

DEPARTMENT OF HEALTH AND HUMAN SERVICES

Administration for Children and Families

Delegation of Authority

Notice is hereby given that, under the authority vested in me by the Secretary of Health and Human Services, I have redelegated to the Commissioner, Administration on Children, Youth and Families, with the authority to further redelegate, the authority to continue the administration of grants and contracts initially awarded in the Fiscal Years 2002, 2003 and 2004 under the Special Projects of Regional and National Significance (SPRANS) Community-based Abstinence Education Program, pursuant to Title V, section 501(a)(2) of the Social Security Act, as amended. The SPRANS Community-based Abstinence Education Program includes Community-based Abstinence Education grants, Abstinence Education Special Congressional Initiative Project grants, and the Abstinence Education Technical Assistance contract with the National Abstinence Clearinghouse. This delegation permits the Commissioner, Administration on Children, Youth and Families, to administer FY 2002, 2003 and FY 2004 SPRANS abstinence education grants under the terms and conditions of the initial awards, thereby allowing the continuation of the existing grants consistent with recent appropriations enactments (Pub. L. 108–477).

This delegation shall be exercised under the Department's policy on regulations and the existing delegation of authority to approve and issue regulations. This delegation excludes the authority to issue reports to Congress, to take final action to withhold funds from States and to act under the nondiscrimination provisions of the Social Security Act.

This delegation also supersedes all prior delegations of authority to the extent that they are inconsistent with the provisions of this delegation.

I hereby ratify any actions taken by the Commission, Administration on Children Youth and Families, or any other Administration on Children, Youth and Families official, which, in effect, involved the exercise of this authority prior to the effective date of this delegation.

This delegation is effective on the date of signature.

Dated: October 6, 2005.

Wade. F. Horn,

Assistant Secretary for Children & Families.

[FR Doc. 05–21162 Filed 10–21–05; 8:45am]

BILLING CODE 4184–01–M

DEPARTMENT OF HEALTH AND HUMAN SERVICES

Food and Drug Administration

[Docket No. 2005N–0395]

Agency Information Collection Activities: Proposed Collection; Comment Request; Guidance for Industry on Formal Meetings With Sponsors and Applicants for Prescription Drug User Fee Act Products

AGENCY: Food and Drug Administration, HHS.

ACTION: Notice.

SUMMARY: The Food and Drug Administration (FDA) is announcing an opportunity for public comment on the proposed collection of certain information by the agency. Under the Paperwork Reduction Act of 1995 (the PRA), Federal agencies are required to publish notice in the **Federal Register** concerning each proposed collection of information, including each proposed extension of an existing collection of information, and to allow 60 days for public comment in response to the notice. This notice solicits comments on the information collection contained in the guidance for industry on formal meetings with sponsors and applicants for Prescription Drug User Fee Act (PDUFA) products.

DATES: Submit written or electronic comments on the collection of information by December 23, 2005.

ADDRESSES: Submit electronic comments on the collection of information to *http://www.fda.gov/dockets/ecomments*. Submit written comments on the collection of information to the Division of Dockets Management (HFA–305), Food and Drug Administration, 5630 Fishers Lane, rm. 1061, Rockville, MD 20852. All comments should be identified with the docket number found in brackets in the heading of this document.

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

SUPPLEMENTARY INFORMATION: Under the PRA (44 U.S.C. 3501–3520), Federal agencies must obtain approval from the Office of Management and Budget (OMB) for each collection of information they conduct or sponsor. ''Collection of information'' is defined in 44 U.S.C. 3502(3) and 5 CFR 1320.3(c) and includes agency requests or requirements that members of the public submit reports, keep records, or provide information to a third party. Section 3506(c)(2)(A) of the PRA (44 U.S.C. 3506(c)(2)(A)) requires Federal agencies to provide a 60-day notice in the **Federal Register** concerning each proposed collection of information, including each proposed extension of an existing collection of information, before submitting the collection to OMB for approval. To comply with this requirement, FDA is publishing notice of the proposed collection of information set forth in this document.

With respect to the following collection of information, FDA invites comments on these topics: (1) Whether the proposed collection of information is necessary for the proper performance of FDA's functions, including whether the information will have practical