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. E-mail address: *infocollection@acf.hhs.gov*.

OMB Comment: OMB is required to make a decision concerning the collection 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, 725 17th Street, NW., Washington, DC 20503, Attn: Desk Officer for ACF, E-mail address: *Katherine_T._Astrich@eop.gov*.

Dated: July 27, 2006.

Robert Sargis,

Reports Clearance Officer.

[FR Doc. 06–6619 Filed 8–1–06; 8:45 am]

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DEPARTMENT OF HEALTH AND HUMAN SERVICES

Administration for Children and Families

[OMB No. 0980–0270]

Submission for OMB Review; Comment Request; Developmental Disabilities Protection and Advocacy Statement of Goals and Priorities

Description: Federal statute and regulation require each State Protection and Advocacy (P&A) System to prepare and submit to public comment a Statement of Goals and Priorities (SGP) for the P&A for Developmental Disabilities (PADD) program for each coming fiscal year. While the P&A is mandated to protect and advocate under a range of different Federally authorized disabilities programs, only the PADD program requires an SGP. Following the required public input for the coming fiscal year, the P&As submit the final version of this SGP to the Administration on Developmental Disabilities (ADD). ADD will aggregate the information in the SGPs into a national profile of programmatic emphasis for P&A Systems in the coming year. This aggregation will provide ADD with a tool for monitoring of the public input requirement. Furthermore, it will provide an overview of program direction, and permit ADD to track accomplishments against goals/targets, permitting the formulation of technical assistance and compliance with the Government Performance and Results Act of 1993.

Respondents: State and Tribal Governments.

ANNUAL BURDEN ESTIMATES

Instrument	Number of respondents	Number of responses per respondent	Average burden hours per response	Total burden hours
P&A SGP	57	1	44	2,508
Estimated Total Annual Burden Hours:				2,508

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, 725 17th Street, NW., Washington, DC 20503, Attn: Desk Officer for ACF, E-mail address: *Ketherine__T.__Astrich@omb.eop.gov.*

Dated: July 27, 2007.

Robert Sargis,

Reports Clearance Officer.

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DEPARTMENT OF HEALTH AND HUMAN SERVICES

Food and Drug Administration

Animal Drug User Fee Rates and Payment Procedures for Fiscal Year 2007

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) 2007 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), 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 2007.

For FY 2007, the animal drug user fee rates are: $168,600 for an animal drug application; $84,300 for a supplemental animal drug application for which safety or effectiveness data is required; $4,115 for an annual product fee; $51,350 for an annual establishment fee; and $44,850 for an annual sponsor fee. FDA will issue invoices for FY 2007 product, establishment, and sponsor fees by December 30, 2006, and these invoices will be due and payable by January 31, 2007.

The application fee rates are effective for applications submitted on or after October 1, 2006, and will remain in effect through September 30, 2007. 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 the FDA Web site at *http://www.fda.gov/oc/adufa* or contact Robert Miller, Center for Veterinary Medicine (HFV–