1992 (57 FR 48801); February 28, 1994 (59 FR 9486); April 29, 1996 (61 FR 18744; November 17, 1997 (62 FR 61332); October 21, 1999 (64 FR 56792); October 25, 2001 (66 FR 54014) and November 7, 2003 (68 FR 63098).

Notice of the availability of drafts of these six updated and one new toxicological profiles for public review and comment will be published in the **Federal Register** on/or about October 17, 2006, with notice of a 90-day public comment period for each profile, starting from the actual release date. Following the close of the comment period, chemical-specific comments will be addressed, and, where appropriate, changes will be incorporated into each profile.

Development of Toxicological Profiles

This notice announces the development of one new and six updated toxicological profiles of priority hazardous substances comprising the twentieth set prepared by ATSDR.

The following toxicological profiles are now being developed:

SET 20 TOXICOLOGICAL PROFILES

Toxicological profile	CAS number
1. Aluminum	7429–90–5
2. Cresols	1319–77–3
3. Diazinon	0333–41–5
4. Dichloropropene, 1,3	0542–75–6
5. Guthion*	0086–50–0
6. Phenol	0108–95–2
7. Tetrachloroethane, 1,1,2,2-	0079–34–5

* Denotes new profile.

Dated: July 28, 2006.

Ken Rose,

Acting Director, Office of Policy, Planning and Evaluation, National Center for Environmental Health, Agency for Toxic Substances and Disease, Disease Registry.

[FR Doc. E6–12417 Filed 8–1–06; 8:45 am]

BILLING CODE 4163–70–P

DEPARTMENT OF HEALTH AND HUMAN SERVICES

Centers for Disease Control and Prevention

National Center for Environmental Health/Agency for Toxic Substances and Disease Registry; The Program Peer Review Subcommittee of the Board of Scientific Counselors (BSC), Centers for Disease Control and Prevention (CDC), National Center for Environmental Health/Agency or Toxic Substances and Disease Registry (NCEH/ATSDR): Teleconference

In accordance with section 10(a)(2) of the Federal Advisory Committee Act (Pub. L. 92–463), CDC, NCEH/ATSDR announces the following subcommittee meeting:

Name: Program Peer Review Subcommittee (PPRS).

Time and Date: 10 a.m.–12 p.m. Eastern Daylight Savings Time, August 14, 2006.

Place: The teleconference will originate at NCEH/ATSDR in Atlanta, Georgia. To participate, dial 877–315–6535 and enter conference code 383520.

Purpose: Under the charge of the BSC, NCEH/ATSDR, the PPRS will provide the Board with advice and recommendations on NCEH/ATSDR program peer review. They will serve the function of organizing, facilitating, and providing a long-term perspective to the conduct of NCEH/ATSDR program peer review.

Matters to be Discussed: A review of the minutes from the previous meeting; a review and discussion of the draft Program Peer Review Self-Assssment Tool questionnaire; a review and discussion of the partners' and senior management questionnaires; a discussion of the peer review site visit for the Division of Health Assessment and Consultation and the Division of Regional Operation; and a discussion of the peer review timeline.

Due to programmatic matters, this **Federal Register** Notice is being published on less than 15 calendar days notice to the public (41 CFR 102–.3.150(b))

Agenda items are subject to change as priorities dictate.

FOR FURTHER INFORMATION CONTACT: Sandra Malcom, Committee Management Specialist, office of Science, NCEH/ATSDR, M/S E–28, 1600 Clifton Road, NE., Atlanta, Georgia 30333, telephone 404–498–0622.

SUPPLEMENTARY INFORMATION: Public comment period is scheduled for 11:30 a.m.–11:40 a.m.

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 CDC and NCEH/ATSDR. Analysis and Services Office, Centers for Disease Control and Prevention.

Dated: July 28, 2006.

Alvin Hall,

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

[FR Doc. 06–6665 Filed 8–1–06; 8:45 am]

BILLING CODE 4163–18–M

DEPARTMENT OF HEALTH AND HUMAN SERVICES

Administration for Children and Families

[OMB No.: 0970–0207]

Submission for OMB Review; Comment Request; Head Start Program Grant Application and Budget Instrument

Description: The Head Start Bureau is proposing to renew, without changes, the Head Start Grant Application and Budget Instrument, which standardizes the grant application information that is requested from all Head Start and Early Head Start grantees applying for continuation grants. The application and budget forms are available on a data diskette and on the web at *www.acfgabi.com.* Completed applications can be transmitted electronically to Regional and Central Offices. The Administration on Children, Youth and Families believes that this application document makes the process of applying for Head Start program grants more efficient for applicants.

Respondents: Head Start and Early Head Start grantees.

ANNUAL BURDEN ESTIMATES

Instrument	Number of respondents	Number of responses per respondent	Average burden hours per response	Total burden hours
HS grant and budget instrument	1,600	1	33	52,800
Estimated total annual burden hours:				52,800

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]

BILLING CODE 4184–01–M

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.

[FR Doc. 06–6620 Filed 8–1–06; 8:45 am]

BILLING CODE 4184–01–M

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–