## DEPARTMENT OF HEALTH AND HUMAN SERVICES

### Administration for Children and Families

### Submission for OMB Review; Comment Request

*Title:* Developmental Disabilities Annual Protection and Advocacy Systems Program Performance Report.

*OMB No.:* 0980–0160.

*Description:* This information collection is required by Federal statute. Each State Protection and Advocacy System must prepare and submit a program Performance Report for the preceding fiscal year of activities and accomplishments and of conditions in the State. The information in the Annual Report will be aggregated into a national profile of Protection and Advocacy Systems. It will also provide the Administration on Developmental Disabilities (ADD) with an overview of program trends and achievements and will enable ADD to respond to administration and congressional requests for specific information on program activities. This information will also be used to submit a Centennial Report to Congress as well as to comply with requirements in the Government Performance and Results Act of 1993.

*Respondents:* Protection and Advocacy Entities.

ANNUAL BURDEN ESTIMATES

| Instrument | Number of respondents | Number of responses per respondent | Average burden hours per response | Total burden hours |
|---|---|---|---|---|
| Program Performance Report | 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, Fax: 202–395–7285, E-mail: *OIRA_SUBMISSION@OMB.EOP.GOV,* Attn: Desk Officer for the Administration for Children and Families.

**Robert Sargis,**

*Reports Clearance Officer.*

[FR Doc. 2011–9772 Filed 4–21–11; 8:45 am]

**BILLING CODE 4184–01–P**

## DEPARTMENT OF HEALTH AND HUMAN SERVICES

### Food and Drug Administration

**[Docket No. FDA–2011–N–0033]**

### Withdrawal of Approval of New Animal Drug Applications; Phenylbutazone; Pyrantel; Tylosin; Sulfamethazine; Correction

**AGENCY:** Food and Drug Administration, HHS.

**ACTION:** Notice; correction.

**SUMMARY:** The Food and Drug Administration (FDA) published a document in the **Federal Register** of March 2, 2011 (76 FR 11490), providing notice of the voluntary withdrawal of approval of eight new animal drug applications (NADAs). That document contained an error in the preamble. FDA is correcting the name and address for the sponsor of five of the NADAs. This correction is being made to improve the accuracy of the **Federal Register**.

**FOR FURTHER INFORMATION CONTACT:** George K. Haibel, Center for Veterinary Medicine (HFV–6), Food and Drug Administration, 7519 Standish Pl., Rockville, MD 20855, 240–276–9019, e-mail: *george.haibel@fda.hhs.gov.*

**SUPPLEMENTARY INFORMATION:** In the **Federal Register** of March 2, 2011, in FR Doc. 2011–4545, on page 11490, in the first column, correct the Truow Nutrition, Inc., name and address to read: Trouw Nutrition USA LLC, P.O. Box 219, 115 Executive Dr., Highland, IL 62249.

Dated: April 15, 2011.

**Bernadette Dunham,**

*Director, Center for Veterinary Medicine.*

[FR Doc. 2011–9778 Filed 4–21–11; 8:45 am]

**BILLING CODE 4160–01–P**

## DEPARTMENT OF HEALTH AND HUMAN SERVICES

### Health Resources and Services Administration

### Administration for Children and Families

### Advisory Committee on the Maternal, Infant and Early Childhood Home Visiting Program Evaluation; Notice of Meeting

In accordance with section 10(a)(2) of the Federal Advisory Committee Act (Pub. L. 92–463, 5 U.S.C. appendix 2), notice is hereby given of the following meeting:

*Name:* Advisory Committee on the Maternal, Infant and Early Childhood Home Visiting (MIECHV) Program Evaluation.

*Date and Time:* Thursday, May 5, 2011:
9 a.m.–5:15 p.m. EST. Friday, May 6, 2011:
9 a.m.–2:15 p.m. EST.

*Place:* Hilton Alexandria Old Town, 1767 King Street, Alexandria, Virginia 22314. (703) 837–0440.

This notice announces a forthcoming meeting of a public advisory committee of the Health Resources and Services Administration and the Administration for Children and Families. The meeting will be open to the public. This notice is being published less than 15 days prior to the meeting due to difficulties in securing adequate and accessible space to accommodate the public.

*Meeting Registration:* To register for the meeting, the public can contact Carolyn Swaney at *cswaney@icfi.com.*

*Agenda:* The purpose of this meeting is to gather comments from the Committee on the design of the MIECHV

program evaluation. Topics to be discussed include an overview of the impact and implementation study designs, sampling design, analysis of state needs assessments, and cost effectiveness study.

*Public Comments:* The public can submit comments for the Committee on the design of the national evaluation of the home visiting program to Carlos Cano, Health Resources and Services Administration, at *ccano@hrsa.gov.* Comments should be submitted by May 2, 2011.

*Special Accommodations:* Individuals who plan to attend and need special assistance, such as sign language interpretation or other reasonable accommodations, should notify the contact person listed above at least 10 days prior to the meeting.

*For Further Information Contact:* T'Pring Westbrook, Administration for Children and Families, *tpring.westbrook@acf.hhs.gov.*

*Supplementary Information:* The Advisory Committee on the Maternal, Infant and Early Childhood Home Visiting Program Evaluation is authorized by subsection 511(g)(1) of Title V of the Social Security Act (42 U.S.C. 701 *et seq.*) as amended by section 2951 of the Patient Protection and Affordable Care Act of 2010 (Pub. L. 111–148) (the Affordable Care Act). The purpose of the Committee is to advise the Secretary of Health and Human Services on the design, plan, progress, and findings of the evaluation required for the home visiting program under the Affordable Care Act. More specifically, the Committee is to review, and make recommendations on, the design and plan for this evaluation; maintain and advise the Secretary regarding the progress of the evaluation; and comment, if the Committee so desires, on the report submitted to Congress under subsection 511(g)(3) of Title V.

Under a government contract, the MDRC, formerly known as Manpower Demonstration Research Corporation, a nonprofit, nonpartisan education and social policy research organization, developed the design options for the evaluation of the home visiting program. These study design options for this national evaluation will be formally presented to the Committee for review. As specified in the legislation, the evaluation will provide a state-by-state analysis of the needs assessments and the States' actions in response to the assessments. Additionally, as specified in the legislation, the evaluation will provide an assessment of: (a) The effect of early childhood home visiting programs on outcomes for parents, children, and communities with respect to domains specified in the Affordable Care Act (such as maternal and child health status, school readiness, and domestic violence, among others); (b) the effectiveness of such programs on different populations, including the extent to which the ability to improve participant outcomes varies across programs and populations; and (c) the potential for the activities conducted under such programs, if scaled broadly, to enhance health care practices, eliminate health disparities, improve health care system quality, and reduce costs.

Dated: April 18, 2011.

**Mary K. Wakefield,**

*Administrator, Health Resources and Services Administration.*

Dated: April 18, 2011.

**Joan Lombardi,**

*Deputy Assistant Secretary and Inter-Departmental Liaison for Early Childhood Development, Administration for Children and Families.*

[FR Doc. 2011–9756 Filed 4–21–11; 8:45 am]

**BILLING CODE 4165–15–P**

## DEPARTMENT OF HEALTH AND HUMAN SERVICES

### National Institutes of Health

### Proposed Collection; Comment Request; Health Information National Trends Survey 4 (HINTS 4) (NCI)

**SUMMARY:** In compliance with the requirement of Section 3506(c)(2)(A) of the Paperwork Reduction Act of 1995, for opportunity for public comment on proposed data collection projects, the National Cancer Institute (NCI), the National Institutes of Health (NIH) will publish periodic summaries of proposed projects to be submitted to the Office of Management and Budget (OMB) for review and approval.

*Proposed Collection: Title:* Health Information National Trends Survey 4 (HINTS 4) (OMB 0925–0538, Exp 11/30/2008). *Type of Information Collection Request:* Reinstatement with Change. *Need and Use of Information Collection:* HINTS 4 will provide NCI with a comprehensive assessment of the American public's current access to, and use of, information about cancer across the cancer care continuum from cancer prevention, early detection, diagnosis, treatment, and survivorship. The content of the survey will focus on understanding the degree to which members of the general population understand vital cancer prevention messages. More importantly, this NCI survey will couple knowledge-related questions with inquiries into the communication channels through which understanding is being obtained, and assessment of cancer-related behavior. The Public Health Services Act, Sections 411 (42 U.S.C. 285a) and 412 (42 U.S.C. 285a–1.1 and 285a–1.3), outline the research and information dissemination mission of the NCI which authorizes the collection of this information. *Frequency of Response:* Once. *Affected Public:* Individuals. *Type of Respondents:* U.S. adults (persons aged 18+). The annual reporting burden is documented in the table below. There are no Capital Costs, Operating Costs, and/or Maintenance Costs to report.

| Data collection cycle | Type of respondent | Number of respondents | Frequency of response | Average time per response minutes/hour | Annual hour burden |
|---|---|---|---|---|---|
| Cycle 1 | Mail survey | 3,533 | 1 | 30/60 (.5) | 1,766.5 |
| Cycle 2 | Mail survey | 3,533 | 1 | 30/60 (.5) | 1,766.5 |
| Cycle 3 | Mail survey | 3,500 | 1 | 30/60 (.5) | 1,750 |
| Cycle 4 | Mail survey | 3,500 | 1 | 30/60 (.5) | 1,750 |
| Total | | | | | 7,033 |