

and inadequate responses in educational settings, limiting their educational and social success.

To address these issues, the Tourette Syndrome Association has developed educational workshops and materials to improve the recognition and awareness of TS diagnosis, treatment, co-occurring conditions, and quality of life for those impacted by TS. Health education programs have been developed for 3 groups of audiences: Health professionals, education professionals, and people with TS and their families.

The format includes general education programs for the 3 groups, as well as two more in-depth medical training programs for physicians on TS and on the Comprehensive Behavioral Intervention for Tics (CBIT) treatment. In addition, a range of professional health education materials in various formats have been developed as educational resources and will be disseminated.

CDC requests OMB approval to collect program evaluation information from workshop participants and recipients of

educational materials over a three-year period. Participants of the workshops and recipients of educational resources will be completing program evaluation forms to provide information on whether the workshop or resource met the educational goals. The information will be used to improve future workshops.

There are no costs to respondents other than their time. The total estimated annual burden hours are 277.

ESTIMATED ANNUALIZED BURDEN HOURS

Type of respondent	Form name	Number of respondents	Number of responses/ respondent	Average burden/response (in hours)	Response burden (in hours)
Health professionals	Medical Program Evaluation	1,200	1	2/60	40
Health professionals	Physician Training Retreat Pre-test	50	1	3/60	3
	Physician Training Retreat Post-test	50	1	3/60	3
	Physician Training Retreat 3-Month Follow-up.	30	1	2/60	1
Health professionals	CBIT Program Evaluation	500	1	2/60	17
	CBIT Pre-test	500	1	3/60	25
	CBIT Post-test	500	1	3/60	25
	CBIT Online Program Evaluation	50	1	1/60	1
	CBIT Program 3-Month Follow-up ...	300	1	1/60	5
Health professionals	Medical Resource Dissemination	210	1	2/60	7
Teachers/Educators	Education Program Evaluation	1,200	1	2/60	40
	Education Program Pre-test	800	1	3/60	40
	Education Program Post-test	800	1	3/60	40
	Education Resource Dissemination	210	1	2/60	7
Public	Family/Public Education Program Evaluation.	250	1	2/60	8
	Family/Public Medical Program Evaluation.	250	1	2/60	8
	Family Resource Dissemination	200	1	2/60	7

Dated: July 22, 2011.

Daniel L. Holcomb,

Reports Clearance Officer, Centers for Disease Control and Prevention.

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DEPARTMENT OF HEALTH AND HUMAN SERVICES

Administration for Children and Families

Submission for OMB Review; Comment Request

Title: ACF-535 LIHEAP Quarterly Allocation Estimates.

OMB No.: 0970-0037.

Description: The LIHEAP Quarterly Allocation Estimates, ACF Form-535 is a one-page form that is sent to 50 State grantees and to the District of Columbia. It is also sent to Tribal Government grantees that receive over \$1 million annually for the Low Income Home Energy Assistance Program (LIHEAP). Grantees are asked to complete and submit the form in the 4th quarter of each year. The data collected on the form are grantees estimates of obligations they expect to make each quarter for the upcoming fiscal year for the LIHEAP program. This is the only method used to request anticipated distributions of the grantees LIHEAP

funds. The information is used to develop apportionment requests to OMB and to make grant awards based on grantees anticipated needs. Information collected on this form is not available through any other Federal source. Submission of the form is voluntary.

Respondents: State Governments.

ANNUAL BURDEN ESTIMATES

Instrument	Number of respondents	Number of responses per respondent	Average burden hours per response	Total burden hours
LIHEAP Quarterly Allocation Estimate, ACF-535	55	1	0.25	13.75

Estimated Total Annual Burden Hours: 13.75

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:* 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. 2011-19104 Filed 7-27-11; 8:45 am]

BILLING CODE 4184-01-P

DEPARTMENT OF HEALTH AND HUMAN SERVICES

Administration for Children and Families

Submission for OMB Review; Comment Request

Title: Child Abuse Prevention Program.

OMB No.: 0970-0155.

Description: The Program Instruction, prepared in response to the enactment of the Community-Based Grants for the Prevention of Child Abuse and Neglect (administratively known as the Community Based Child Abuse Prevention Program, (CBCAP), as set forth in Title II of Public Law 111-320,

Child Abuse Prevention and Treatment Act Amendments of 2010, provides direction to the States and Territories to accomplish the purposes of (1) supporting community-based efforts to develop, operate, expand, enhance and coordinate initiatives, programs, and activities to prevent child abuse and neglect; (2) supporting the coordination of resources and activities to better strengthen and support families to reduce the likelihood of child abuse and neglect, and; (3) fostering an understanding, appreciation, and knowledge of diverse populations in order to effectively prevent child abuse and neglect. This Program Instruction contains information collection requirements that are found in Pub. L. 111-320 at sections 201; 202; 203; 205; 206; and pursuant to receiving a grant award. The information submitted will be used by the agency to ensure compliance with the statute, complete the calculation of the grant award entitlement, and provide training and technical assistance to the grantee.

Respondents: State Governments.

ANNUAL BURDEN ESTIMATES

Instrument	Number of respondents	Number of responses per respondent	Average burden hours per response	Total burden hours
Application	52	1	40	2,080
Annual Report	52	1	24	1,248

Estimated Total Annual Burden Hours: 3,328.

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:* 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:

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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.

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DEPARTMENT OF HEALTH AND HUMAN SERVICES

Food and Drug Administration

[Docket No. FDA-2011-N-0510]

Agency Information Collection Activities; Proposed Collection; Comment Request; Substances Prohibited From Use in Animal Food or Feed

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 recordkeeping requirements for substances prohibited for use in animal food or feed.

DATES: Submit electronic or written comments on the collection of information by September 26, 2011.

ADDRESSES: Submit electronic comments on the collection of information to: <http://www.regulations.gov>. 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: Juanmanuel Vilela, Office of