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DEPARTMENT OF HEALTH AND HUMAN SERVICES

Administration for Children and Families

Proposed Information Collection Activity; Comment Request

Title: Federal Strategic Action Plan on
Services for Victims of Human
Trafficking: Enhancing the Health Care
System's Response to Human
Trafficking

OMB No.: New Collection

Description:

In 2013, the U.S. Department of Health and Human Services co-chaired an inter-agency process with the Departments of Justice and Homeland Security to create the first Federal Strategic Action Plan on Services for Victims of Human Trafficking in the United States. The Plan addresses the needs for the implementation of coordinated, effective, culturally appropriate and trauma informed care for victims of human trafficking. The purpose of this initiative is to develop a pilot training project that will strengthen the health systems' response to human trafficking in four key ways

1. Increase knowledge about human trafficking among health care providers;
2. Build the capacity of health care providers to deliver culturally

appropriate and trauma-informed care to victims of human trafficking;

3. Increase the identification of victims of human trafficking; and

4. Increase services to survivors of human trafficking.

The evaluation will measure immediate outcomes, e.g., from pre-intervention to post-intervention, as well as intermediate outcomes at a 3 month post intervention.

Respondents:

The target audience for training and evaluation will be 200 health care providers from hospitals, clinics, and private health practices. The health care providers will be from federal, state/territorial, and local health departments, the Veterans' Administration, professional associations, and tribal institutions.

ANNUAL BURDEN ESTIMATES

Instrument	Number of respondents	Number of responses per respondent	Average burden hours per response	Total burden hours
Pre-training survey	200	1	0.40	80.00
Post-training survey	200	1	0.40	80.00
Email Follow-up	200	1	0.40	80.00
Telephone Follow-up	40	1	0.40	16.00
.....	256.00

Estimated Total Annual Burden
Hours: 256

In compliance with the requirements of Section 506(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 Planning, Research and Evaluation, 370 L'Enfant Promenade, SW., Washington, DC 20447, Attn: ACF Reports Clearance Officer. Email 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 the 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.

Robert Sargis,

Reports Clearance Officer.

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DEPARTMENT OF HEALTH AND HUMAN SERVICES

Food and Drug Administration

[Docket No. FDA-2013-N-0878]

Agency Information Collection Activities; Announcement of Office of Management and Budget Approval; Premarket Notification for a New Dietary Ingredient

AGENCY: Food and Drug Administration, HHS.

ACTION: Notice.

SUMMARY: The Food and Drug Administration (FDA) is announcing

that a collection of information entitled "Premarket Notification for a New Dietary Ingredient" has been approved by the Office of Management and Budget (OMB) under the Paperwork Reduction Act of 1995.

FOR FURTHER INFORMATION CONTACT: FDA PRA Staff, Office of Operations, Food and Drug Administration, 1350 Piccard Dr., PI50-400B, Rockville, MD 20850, PRASStaff@fda.hhs.gov.

SUPPLEMENTARY INFORMATION: On November 25, 2013, the Agency submitted a proposed collection of information entitled "Premarket Notification for a New Dietary Ingredient" to OMB for review and clearance under 44 U.S.C. 3507. An Agency may not conduct or sponsor, and a person is not required to respond to, a collection of information unless it displays a currently valid OMB control number. OMB has now approved the information collection and has assigned OMB control number 0910-0330. The approval expires on February 28, 2015. A copy of the supporting statement for this information collection is available on the Internet at <http://www.reginfo.gov/public/do/PRAMain>.