

DEPARTMENT OF HEALTH AND HUMAN SERVICES

Administration for Children and Families

Proposed Information Collection Activity; Comment Request

Title: Accomplishments of the Domestic Violence Hotline, Online Connections and Text (ADVHOCaT) Study.

OMB No.: New Collection.

Description: The National Domestic Violence Hotline (NDVH) and the National Dating Abuse Helpline or Love Is Respect (NDAH/LIR), which are supported by the Division of Family Violence Prevention and Services

within the Family and Youth Services Bureau of the Administration for Children and Families (ACF), U.S. Department of Health and Human Services (HHS), serve as partners in the intervention, prevention, and resource assistance efforts of the network of family violence, domestic violence, and dating violence service providers.

In order to describe the activities and accomplishments of the NDVH and NDAH/LIR and develop potential new or revised performance measures, the Office of Planning, Research and Evaluation (OPRE), within ACF/HHS is proposing data collection activity as part of the Accomplishments of the Domestic Violence Hotline, Online Connections and Text (ADVHOCaT) Study.

This study will primarily analyze data previously collected by the NDVH and NDAH/LIR as part of their ongoing program activities and monitoring. ACF proposes to collect additional information, including information about the preferred mode (phone, chat, text), ease of use, and perceived safety of each mode of contact.

This data is to be collected through voluntary web-based surveys that are to be completed by those who access the NDVH and NDAH/LIR Web sites. This information will be critical to informing future efforts to monitor and improve the performance of domestic violence hotlines and provide hotline services.

Respondents: Individuals who access the NDVH and NDAH/LIR Web sites.

ANNUAL BURDEN ESTIMATES

Instrument	Total/annual number of respondents	Number of responses per respondent	Average burden hours per response	Annual burden hours
NDVH/LIR Preference of Use Survey	5000	1	0.041 hours (150 seconds)	205

Estimated Total Annual Burden Hours: 205 hours.

In compliance with the requirements of Section 3506(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: OPRE Reports Clearance Officer. Email address: OPREinfocollection@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.

Karl Koerper,

Reports Clearance Officer.

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DEPARTMENT OF HEALTH AND HUMAN SERVICES

Food and Drug Administration

[Docket No. FDA-2015-N-2406]

Agency Information Collection Activities; Proposed Collection; Comment Request; Market Claims in Direct-to-Consumer Prescription Drug Print Ads

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 and to allow 60 days for public comment in response to the notice. This notice solicits comments on research entitled, "Market Claims in Direct-to-Consumer Prescription Drug Print Ads." This study will examine the

impact of market claim information in direct-to-consumer (DTC) print advertising for prescription drugs.

DATES: Submit either electronic or written comments on the collection of information by September 18, 2015.

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: FDA PRA Staff, Office of Operations, Food and Drug Administration, 8455 Colesville Rd., COLE-14526, Silver Spring, MD 20993-0002, PRAStaff@fda.hhs.gov.

SUPPLEMENTARY INFORMATION: Under the PRA (44 U.S.C. 3501-3520), Federal Agencies must obtain approval from the Office of Management and Budget (OMB) for each collection of information they conduct or sponsor. "Collection of information" is defined in 44 U.S.C. 3502(3) and 5 CFR 1320.3(c) and includes Agency requests or requirements that members of the public submit reports, keep records, or provide information to a third party. Section 3506(c)(2)(A) of the PRA (44 U.S.C. 3506(c)(2)(A)) requires Federal