

2. *By regular mail.* You may mail written comments to the following address: CMS, Office of Strategic Operations and Regulatory Affairs, Division of Regulations Development, Attention: Document Identifier/OMB Control Number, Room C4-26-05, 7500 Security Boulevard, Baltimore, Maryland 21244-1850.

Dated: September 30, 2010.

**Michelle Shortt,**

*Director, Regulations Development Group, Office of Strategic Operations and Regulatory Affairs.*

[FR Doc. 2010-25053 Filed 10-7-10; 8:45 am]

**BILLING CODE 4120-01-P**

**DEPARTMENT OF HEALTH AND HUMAN SERVICES**

**Substance Abuse and Mental Health Services Administration**

**Agency Information Collection Activities: Proposed Collection; Comment Request**

In compliance with Section 3506(c)(2)(A) of the Paperwork Reduction Act of 1995 concerning opportunity for public comment on proposed collections of information, the Substance Abuse and Mental Health Services Administration (SAMHSA) will publish periodic summaries of proposed projects. To request more information on the proposed projects or to obtain a copy of the information collection plans, call the SAMHSA Reports Clearance Officer on (240) 276-1243.

*Comments are invited on:* (a) Whether the proposed collections of information

are 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) ways to enhance 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.

**Proposed Project: 2011-2014 National Survey on Drug Use and Health: Methodological Field Tests (OMB No. 0930-0290-Revision)**

The National Survey on Drug Use and Health (NSDUH) is a survey of the civilian, non-institutionalized population of the United States 12 years old and older. The data are used to determine the prevalence of use of tobacco products, alcohol, illicit substances, and illicit use of prescription drugs. The results are used by SAMHSA, ONDCP, Federal government agencies, and other organizations and researchers to establish policy, direct program activities, and better allocate resources.

In March 2008, SAMHSA received a three-year renewal of its generic clearance for methodological field tests. This will be a request for another renewal of the generic approval to continue methodological tests over the next three years, with conditions similar to the previous clearance. These methodological tests will continue to be designed to examine the feasibility, quality, and efficiency of new

procedures or revisions to existing survey protocol. Specifically, the tests will measure the reliability and validity of certain questionnaire sections and items through multiple measurements on a set of respondents; assess new methods for gaining cooperation and participation of respondents with the goal of increasing response and decreasing potential bias in the survey estimates; and assess the impact of new sampling techniques and technologies on respondent behavior and reporting. Research will involve focus groups, cognitive laboratory testing, field tests, and customer surveys.

The next wave of methodological tests will continue to examine ways to increase data quality, lower operating costs, and gain a better understanding of sources and effects of nonsampling error on the NSDUH estimates. Particular attention will be given to minimizing the impact of design changes so that survey data continue to remain comparable over time. If these tests provide successful results, current procedures or data collection instruments may be revised.

The number of respondents to be included in each field test will vary, depending on the nature of the subject being tested and the target population. However, the total estimated response burden is 8,251 hours. The exact number of subjects and burden hours for each test are unknown at this time, but will be clearly outlined in each individual submission. The table below, however, describes the anticipated burden for each of the major testing activities for which generic approval is being tested.

**ESTIMATED BURDEN FOR NSDUH METHODOLOGICAL FIELD TESTS**

Activity	Number of respondents	Responses per respondent	Total number of responses	Average burden per response (hrs.)	Total burden (hrs.)
a. Focus Groups .....	270	1	270	2.0	540
b. Cognitive laboratory testing .....	200	1	200	1.0	200
c. Field Tests .....	6,600	1	6,600	1.0	6,600
d. Customer Satisfaction Surveys .....	300	1	300	0.25	75
Household screening for c .....	8,910	1	8,910	0.083	740
Screening Verification for c .....	445	1	445	0.067	30
Interview Verification for c .....	990	1	990	0.067	66
<b>Total .....</b>	<b>17,715</b>	<b>.....</b>	<b>17,715</b>	<b>.....</b>	<b>8,251</b>
<b>Annual Average (Total divided by 3 years) .....</b>	<b>5,905</b>	<b>.....</b>	<b>5,905</b>	<b>.....</b>	<b>2,750</b>

Send comments to Summer King, SAMHSA Reports Clearance Officer, Room 8–1099, One Choke Cherry Road, Rockville, MD 20857 and e-mail a copy to [summer.king@samhsa.hhs.gov](mailto:summer.king@samhsa.hhs.gov). Written comments should be received within 60 days of this notice.

Dated: September 30, 2010.

**Elaine Parry,**

Director, Office of Management, Technology and Operations.

[FR Doc. 2010–25439 Filed 10–7–10; 8:45 am]

**BILLING CODE 4162–20–P**

**DEPARTMENT OF HEALTH AND HUMAN SERVICES**

**Administration for Children and Families**

**Submission for OMB Review; Comment Request**

*Title:* Financial Institution Data Match.

*OMB No.:* 0970–0196.

*Description:* Section 466(a)(17) of the Social Security Act (the Act) requires States to establish procedures under which the State Child Support

Enforcement IV–D agencies shall enter into agreements with financial institutions doing business in States for the purpose of securing information leading to the enforcement of child support orders. Under 452(l) and 466(a)(17)(A)(i) of the Act, the Secretary may aid State agencies conducting data matches with financial institutions doing business in multiple States by centrally matching through the Federal Parent Locator Service.

Respondents: Financial institutions doing business in two or more States.

**ANNUAL BURDEN ESTIMATES**

Instrument	Number of respondents	Number of responses per respondent	Average burden hours per response	Total burden hours
Financial Data Match Result File .....	259	4	0.33	341.88
Election Form .....	122	1	0.50	61
Estimated Total Annual Burden Hours .....				402.88

*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](mailto: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](mailto:OIRA_SUBMISSION@OMB.EOP.GOV)  
*Attn:* Desk Officer for the Administration for Children and Families.

Dated: October 5, 2010.

**Robert Sargis,**

Reports Clearance Officer.

[FR Doc. 2010–25414 Filed 10–7–10; 8:45 am]

**BILLING CODE 4184–01–P**

**DEPARTMENT OF HEALTH AND HUMAN SERVICES**

**Food and Drug Administration**

[Docket No. FDA–2010–N–0493]

**Agency Information Collection Activities; Proposed Collection; Comment Request; Additional Criteria and Procedures for Classifying Over-the-Counter Drugs as Generally Recognized as Safe and Effective and Not Misbranded**

**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 the additional criteria and procedures for classifying over-the-counter (OTC) drugs as generally recognized as safe and effective and not misbranded.

**DATES:** Submit either electronic or written comments on the collection of information by December 7, 2010.

**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:**

Elizabeth Berbakos, Office of Information Management, Food and Drug Administration, 1350 Piccard Dr., PI50–400B, Rockville, MD 20850, 301–796–3792, [Elizabeth.Berbakos@fda.hhs.gov](mailto:Elizabeth.Berbakos@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 agencies to provide a 60-day notice in the **Federal Register** concerning each proposed collection of information before submitting the collection to OMB for approval. To comply with this requirement, FDA is publishing notice of the proposed collection of information set forth in this document.

With respect to the following collection of information, FDA invites comments on these topics: (1) Whether the proposed collection of information