

Dated: January 24, 2004.

Georgi Jones,

Director, Office of Policy, Planning, and Evaluation, National Center for Environmental Health/Agency for Toxic Substances and Disease Registry.

[FR Doc. 04-4659 Filed 3-2-04; 8:45 am]

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DEPARTMENT OF HEALTH AND HUMAN SERVICES

Centers for Disease Control and Prevention

Disease, Disability, and Injury Prevention and Control Special Emphasis Panel (SEP): Health Promotion and Disease Prevention Research Centers Cooperative Agreements, Program Announcement Number 04003

In accordance with section 10(a)(2) of the Federal Advisory Committee Act (Pub. L. 92-463), the Centers for Disease Control and Prevention (CDC) announces the following meeting:

Name: Disease, Disability, and Injury Prevention and Control Special Emphasis Panel (SEP): Health Promotion and Disease Prevention Research Centers Cooperative Agreements, Program Announcement Number 04003.

Times and Dates: 8:30 a.m.–9 a.m., March 30, 2004 (open); 9 a.m.–5 p.m., March 30, 2004 (closed); 9 a.m.–5 p.m., March 31, 2004 (closed).

Place: Sheraton Colony Square Hotel, 188 14th Street, NE., Atlanta, GA 30361, telephone 404.892.2004.

Status: Portions of the meeting will be closed to the public in accordance with provisions set forth in section 552b(c) (4) and (6), title 5 U.S.C., and the Determination of the Director, Management Analysis and Services Office, CDC, pursuant to Public Law 92-463.

Matters to be Discussed: The meeting will include the review, discussion, and evaluation of applications received in response to Program Announcement Number 04003.

Contact Person for More Information: Michael N. Waller, Deputy Director, National Center for Chronic Disease Prevention and Health Promotion, CDC, 4770 Buford Highway, NE., MS-K 45, Atlanta, GA 30341, telephone 770.488.5269.

The Director, Management Analysis and Services Office, has been delegated the authority to sign **Federal Register** notices pertaining to announcements of meetings and other committee management activities, for both CDC and the Agency for Toxic Substances and Disease Registry.

Dated: February 24, 2004.

Alvin Hall,

Director, Management Analysis and Services Office, Centers for Disease Control and Prevention.

[FR Doc. 04-4680 Filed 3-2-04; 8:45 am]

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DEPARTMENT OF HEALTH AND HUMAN SERVICES

Administration for Children and Families

Submission for OMB Review; Comment Request

Title: Federal Tax Offset, Administrative Offset, and Passport Denial Program.

OMB No.: 0970-0161.

Description: The Tax Refund Offset and Administrative Offset Programs collect past-due child support by intercepting certain federal payments, including federal tax refunds, of parents who have been ordered to pay child support and are behind in paying the debt. The program is a cooperative effort including the Department of Treasury's Financial Management Service (FMS), the Federal Office of Child Support Enforcement (OCSE) and state Child Support Enforcement (CSE) agencies. The Passport Denial Program reports non-custodial parents who owe arrears above a threshold to the Department of State (DOS), which will then deny passports to these individuals. On an ongoing basis, CSE agencies submit to OCSE the names, Social Security numbers (SSNs) and the amount(s) of past-due child support of people who are delinquent in making child support payments.

Respondents: State IV-D Agencies.

ANNUAL BURDEN ESTIMATES

Instrument	Number of respondents	Number of responses per respondent	Average burden hours per response	Total burden hours
Input Record	54	52	.3 hours ...	842.4 hours.
Output Record	54	52	.46 hours	1292 hours.
Payment File	54	26	.27 hours	379 hours.
Certification Letter	54	1	.4 hours ...	21.6 hours.

Estimated Total Annual Burden Hours: 2535 hours.

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: rsargis@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, Attn: Desk Officer for ACF, E-mail address: katherine_t._astrich@omb.eop.gov.

Dated: February 26, 2004.

Robert Sargis,

Reports Clearance Officer.

[FR Doc. 04-4752 Filed 3-2-04; 8:45 am]

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DEPARTMENT OF HEALTH AND HUMAN SERVICES

Administration for Children and Families

Agency Recordkeeping/Reporting Requirements Under Emergency Review by the Office of Management and Budget (OMB)

Title: ORR Quarterly Performance Report, ORR-6.

OMB No.: 0970-0036.

Description: We ask for the information on this form in order to determine the effectiveness of the state cash and medical assistance, social services, and targeted assistance

programs as required by 412(e) of the Immigration and Naturalization Act. We also calculate state-by-state Refugee Cash Assistance and Refugee Medical

Assistance utilization rates for use in formulating program initiatives, priorities, standards, budget requests, and assistance policies. The Office of

Refugee Resettlement regulations require that this form be completed in order to participate in the program.
Respondents: States.

ANNUAL BURDEN ESTIMATES

Instrument	Number of respondents	Number of responses per respondent	Average burden hours per response	Total burden hours
ORR-6	48	4	3.875	744

Estimated Total Annual Burden Hours: 744 hours.

Additional Information: The Administration for Children and Families (ACF) is requesting that OMB grant a 180-day approval for this information collection under procedures for emergency processing by March 5, 2004. A copy of this information collection, with applicable supporting documentation, may be obtained by calling the ACF Reports Clearance Officer, Robert Sargis at (202) 690-7275. In addition, a request may be made by sending an e-mail request to: rsargis@acf.hhs.gov.

Comments and questions about the information collection described above should be directed to the following address by March 5, 2004: Office of Information and Regulatory Affairs, Attn: OMB Desk Officer for ACF, Office of Management and Budget, Paperwork Reduction Project, Washington, DC. E-mail address: katherine_t_astrich@omb.eop.gov.

Dated: February 26, 2004.

Robert Sargis,

Reports Clearance Officer.

[FR Doc. 04-4753 Filed 3-2-04; 8:45 am]

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DEPARTMENT OF HEALTH AND HUMAN SERVICES

Food and Drug Administration

[Docket No. 2003N-0508]

Agency Information Collection Activities; Submission for Office of Management and Budget Review; Comment Request; Focus Groups as Used by the Food and Drug Administration

AGENCY: Food and Drug Administration, HHS.

ACTION: Notice.

SUMMARY: The Food and Drug Administration (FDA) is announcing that a proposed collection of information has been submitted to the Office of Management and Budget (OMB) for review and clearance under the Paperwork Reduction Act of 1995.

DATES: Fax written comments on the collection of information by April 2, 2004.

ADDRESSES: OMB is still experiencing significant delays in the regular mail, including first class and express mail, and messenger deliveries are not being accepted. To ensure that comments on the information collection are received, OMB recommends that written comments be faxed to the Office of Information and Regulatory Affairs, OMB, Attn: Fumie Yokota, Desk Officer for FDA, FAX: 202-395-6974.

FOR FURTHER INFORMATION CONTACT: JonnaLynn P. Capezuto, Office of Management Programs (HFA-250), Food and Drug Administration, 5600 Fishers Lane, Rockville, MD 20857, 301-827-4659.

SUPPLEMENTARY INFORMATION: In compliance with 44 U.S.C. 3507, FDA has submitted the following proposed

collection of information to OMB for review and clearance.

Focus Groups as Used by the Food and Drug Administration—(OMB Control Number 0910-0497)—Extension

FDA will collect and use information gathered through the focus group vehicle. This information will be used to develop programmatic proposals, and as such, compliments other important research findings to develop these proposals. Focus groups provide an important role in gathering information because they allow for a more in depth understanding of consumers' attitudes, beliefs, motivations, and feelings than quantitative studies.

Also, information from these focus groups will be used to develop policy and redirect resources, when necessary, to our constituents. If this information is not collected, a vital link in information gathering by FDA to develop policy and programmatic proposals will be missed causing further delays in policy and program development.

FDA estimates the burden for completing the forms for this collection of information in table 1 of this document. The total annual estimated burden imposed by this collection of information is 2,830 hours annually.

Annually, FDA projects about 28 focus group studies using 186 focus groups lasting an average of 1.78 hours each. FDA has allowed burden for unplanned focus groups to be completed so as not to restrict the agency's ability to gather information on public sentiment for its proposals in its regulatory as well as other programs.

In the **Federal Register** of November 24, 2003 (68 FR 65938), FDA published a 60-day notice requesting public comment on the information collection provisions. No comments were received.