

role models. Information from the Mentoring Children of Prisoners Online Data Collection is necessary for ACF's reporting and planning under the Government Performance and Results Act (GPRA), and to support evaluation requirements within GPRA. Information collected will be used for accountability monitoring, management improvement,

and research. Data collection ensures that ACF knows if grantees of the MCP program are meeting the established targets (established based on research and benchmarks) recorded in the grant application as required by the GPRA, and that mentoring activities are faithful to characteristics established by research as essential to success. Data

collected will also support grantees as they carry out ongoing responsibilities, maintain program service, and manage information for internal uses.

Respondents: Public, faith-based and community organizations receiving funding to implement the MCP program.

ANNUAL BURDEN ESTIMATES

Instrument	Number of respondents	Number of responses per respondent	Average burden hours per response	Total burden hours
MCP Online Data Collection	238	4	12	11,424

Estimated Total Annual Burden Hours: 11,424.

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.

OMB Comment: OMB is required to make a decision concerning the collection of information between 30 and 60 days after publication of this documentation 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: 202-395-6974,

Attn: Desk Officer for the Administration for Children and Families.

Dated: June 14, 2007.
Robert Sargis,
Reports Clearance Officer.
 [FR Doc. 07-3048 Filed 6-20-07; 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: National Extranet Optimized Runaway and Homeless Youth Management Information System (NEO-RHYMIS).

OMB No.: 0970-0123.
Description: The Runaway and Homeless Youth Act, as amended by

Public Law 106-71 (42 U.S.C. 5701 *et seq.*), mandates that the Department of Health and Human Services (HHS) report regularly to Congress on the status of HHS-funded programs serving runaway and homeless youth. Such reporting is similarly mandated by the Government Performance and Results Act. Organizations funded under the Runaway and Homeless Youth program are required by statute (42 U.S.C. 5712, 42 U.S.C. 5714-2) to meet certain data collection and reporting requirements. These requirements include maintenance of client statistical records on the number and the characteristics of the runaway and homeless youth, and youth at risk of family separation, who participate in the project, and the services provided to such youth by the project.

Respondents: Public and private, community-based nonprofit, and faith-based organizations receiving HHS funds for services to runaway and homeless youth.

ANNUAL BURDEN ESTIMATES

Instrument	Number of respondents	Number of responses per respondent	Average burden hours per response	Total burden hours
Youth Profile	536	153	.25	20,502
Street Outreach Report	141	4211	.02	11,875
Brief Contacts	536	305	.15	24,522
Turnaways	536	13	.1	697
Data Transfer	536	2	.5	536

Estimated Total Annual Burden Hours: 58,132.

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.

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: 202-395-6974, Attn: Desk Officer for the Administration for Children and Families.

Dated: June 14, 2007.

Robert Sargis,

Reports Clearance Officer.

[FR Doc. 07-3049 Filed 6-20-07; 8:45 am]

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DEPARTMENT OF HEALTH AND HUMAN SERVICES

Administration for Children and Families

Statement of Organization, Functions and Delegation of Authority

Notice is hereby given that I have delegated to the Principal Deputy Assistant Secretary, Deputy Assistant Secretaries, Program Directors, Program Commissioners, Deputy Director/Commissioner, Office of Child Support Enforcement, and Staff Office Directors the following authority vested in me by the Secretary of Health and Human Services in the memorandum dated August 20, 1991, Delegations of Authority for Social Security Act Programs; 31 U.S.C. 1535; and HHS General Administrative Manual, Chapter 8-77.

(a) Authorities Delegated

1. Authority to administer approved cooperative research, experimental, pilot or demonstration projects under the provisions of sections 1110 and 1115 of the Social Security Act.

2. Authority to approve interagency agreements to procure, provide or exchange services, supplies or equipment.

(b) Limitations

1. The authority listed in #1 above shall be exercised under the condition that projects may be administered by the Office of Planning, Research and Evaluation (OPRE), by the program/staff office or jointly by OPRE with the program/staff office.

2. Where all or any part of an experimental, pilot, demonstration, or other project is wholly financed with Federal funds made available under sections 1110 or 1115 of the Social Security Act, without any State, local or other non-Federal financial participation, that project must be approved by the Secretary of Health and Human Services.

3. This delegation of authority does not include the authority to approve/disapprove projects under section 1115 of the Social Security Act or approve/disapprove waivers of State Plan requirements or costs that would not otherwise be included as expenditures under the provisions of section

1115(a)(1) and (2) of the Social Security Act.

4. The authority to approve interagency agreements to procure, provide, or exchange services, supplies, or equipment requires the concurrence of the ACF Chief Financial Officer if it exceeds \$250,000 (including amendments) within a fiscal year or if it requires the signature of the Assistant Secretary, ACF, or the Secretary of HHS.

(c) Effective Date

This delegation is effective upon the date of signature.

(d) Effect on Existing Delegations

As related to this delegation of authority, this delegation supersedes all previous delegations of authority involving the administration of the cross-program authorities delegated herein.

I hereby ratify and affirm any actions taken by the Principal Deputy Assistant Secretary, Deputy Assistant Secretaries, Program Directors, Program Commissioners, Deputy Director/Commissioner, Office of Child Support Enforcement, and Staff Office Directors, which involved the exercise of the authority delegated herein prior to the effective date of this delegation.

Dated: June 13, 2007.

Daniel C. Schneider,

Acting Assistant Secretary for Children and Families.

[FR Doc. E7-12019 Filed 6-20-07; 8:45 am]

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DEPARTMENT OF HEALTH AND HUMAN SERVICES

Food and Drug Administration

[Docket No. 2007N-0091]

Agency Information Collection Activities; Submission for Office of Management and Budget Review; Comment Request; Preparing a Claim of Categorical Exclusion or an Environmental Assessment for Submission to the Center for Food Safety and Applied Nutrition

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 July 23, 2007.

ADDRESSES: 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: FDA Desk Officer, FAX: 202-395-6974. All comments should be identified with the OMB control number 0910-0541. Also include the FDA docket number found in brackets in the heading of this document.

FOR FURTHER INFORMATION CONTACT: Jonna Capezzuto, Office of the Chief Information Officer (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.

Preparing a Claim of Categorical Exclusion or an Environmental Assessment for Submission to the Center for Food Safety and Applied Nutrition (OMB Control Number 0910-0541)—Extension

As an integral part of its decisionmaking process, FDA is obligated under the National Environmental Policy Act of 1969 (NEPA) to consider the environmental impact of its actions, including allowing notifications for food contact substances to become effective and approving food additive petitions, color additive petitions, generally recognized as safe affirmation petitions, requests for exemption from regulation as a food additive, and actions on certain food labeling citizen petitions, nutrient content claims petitions, and health claims petitions. In 1997, FDA amended its regulations in part 25 (21 CFR part 25) to provide for categorical exclusions for additional classes of actions that do not individually or cumulatively have a significant effect on the human environment (62 FR 40570, July 29, 1997). As a result of that rulemaking, FDA no longer routinely requires submission of information about the manufacturing and production of FDA-regulated articles. FDA also has eliminated the previously required Environmental Assessment (EA) and abbreviated EA formats from the amended regulations. Instead, FDA has provided guidance that contains sample formats to help industry submit a claim of categorical exclusion or an EA to CFSAN. The guidance document entitled "Preparing a Claim of