and based on valid assumptions and methodology; ways to enhance the quality, utility, and clarity of the information to be collected.

*Obtaining Copies of Proposals:* Requesters may obtain a copy of the information collection documents from the General Services Administration, Regulatory Secretariat Division (MVCB), 1800 F Street NW., Washington, DC 20405, telephone 202–501–4755. Please cite OMB Control No. 3090–0221, Civilian Board of Contract Appeals Rules of Procedure, in all correspondence.

Dated: June 4, 2015.

**David A. Shive,**

*Chief Information Officer.*

[FR Doc. 2015–14446 Filed 6–11–15; 8:45 am]

**BILLING CODE 6820–AL–P**

## 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:* Immediate Disaster Case Management Intake Assessment.

*OMB No.:* 0970–New.

**Description**

Section 426 of the Robert T. Stafford Disaster Relief and Emergency Assistance Act (Stafford Act), as amended, 42 U.S.C. 5189d authorizes the Federal Emergency Management Agency (FEMA) and the U.S. Department of Health Services' Administration for Children and Families (ACF) to provide Immediate Disaster Case Management (IDCM) services under the federal Disaster Case Management Program (DCMP).

The use of the Electronic Case Management Record System (ECMRS) is aligned with Executive Order of the President 13589 and the memorandum to the Heads of Executive Departments and Agencies M–12–12 from the Office of Management and Budget to ''Promote Efficient Spending to Support Agency Operations.''

The primary purpose of the information collection pertains to ACF/OHSEPR's initiative to improve the intake process and delivery of case management services to individuals and households impacted by a disaster. Further, the information collection will be used to support ACF/OHSEPR's goal to quickly identify critical gaps, resources, needs, and services to support State, local and non-profit capacity for disaster case management and to augment and build capacity where none exists.

There are two versions of this Paper Reduction Act request: (1) paper intake assessment that will be used until ECMRS is implemented and operational and (2) Electronic Case Record platform. The ECMRS will greatly reduce respondent burden through built-in algorithms that will streamline response options and patterns. All information gathered will be exclusively used to inform the delivery of disaster case management services and programmatic strategies and improvements.

*Respondents:* Individuals impacted by a major disaster.

ANNUAL BURDEN ESTIMATES

| Instrument | Number of respondents | Number of responses per respondent | Average burden hours per response | Total burden hours |
|---|---|---|---|---|
| IDCM Intake Assessment | 3,500 | 1 | 40 minutes | 140,000 minutes |

Estimated Total Annual Burden Hours: 140,000 minutes.

**Additional Information**

ACF is requesting that OMB grant a 180 day approval for this information collection under procedures for emergency processing by June 19, 2015. A copy of this information collection, with applicable supporting documentation, may be obtained by calling the Administration for Children and Families, Reports Clearance Officer, Robert Sargis at (202) 690–7275.

Comments and questions about the information collection described above should be directed to the Office of Information and Regulatory Affairs, Attn: OMB Desk Officer for ACF, Office of Management and Budget, Paperwork Reduction Project, 725 17th Street NW., Washington, DC 20503; FAX: (202) 395–7285; email:*oira_submission@omb.eop.gov.*

**Robert Sargis,**

*Reports Clearance Officer.*

[FR Doc. 2015–14400 Filed 6–11–15; 8:45 am]

**BILLING CODE 4184–01–P**

## DEPARTMENT OF HEALTH AND HUMAN SERVICES

### Food and Drug Administration

**[Docket No. FDA–2008–N–0543]**

### Agency Information Collection Activities; Announcement of Office of Management and Budget Approval; Waivers of Invivo Demonstration of Bioequivalence of Animal Drugs in Soluble Powder Oral Dosage Form and Type A Medicated Articles

**AGENCY:** Food and Drug Administration, HHS.

**ACTION:** Notice.

**SUMMARY:** The Food and Drug Administration (FDA) is announcing that a collection of information entitled, ''Waivers of Invivo Demonstration of Bioequivalence of Animal Drugs in Soluble Powder Oral Dosage Form and Type A Medicated Articles'' 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, 8455 Colesville Rd., COLE–14526, Silver Spring, MD 20993–0002, *PRAStaff@fda.hhs.gov.*

**SUPPLEMENTARY INFORMATION:** On April 24, 2015, the Agency submitted a proposed collection of information entitled, ''Waivers of Invivo Demonstration of Bioequivalence of Animal Drugs in Soluble Powder Oral Dosage Form and Type A Medicated