have a significant economic impact on a substantial number of small entities. Therefore, we are not preparing an analysis for the RFA.

In addition, section 1102(b) of the Social Security Act (the Act) requires us to prepare a regulatory impact analysis if a rule may have a significant impact on the operations of a substantial number of small rural hospitals. This analysis must conform to the provisions of section 604 of the RFA. For purposes of section 1102(b) of the Act, we define a small rural hospital as a hospital that is located outside of a Metropolitan Statistical Area and has fewer than 100 beds. We have determined that this notice will not have a significant effect on the operations of a substantial number of small rural hospitals. Therefore, we are not preparing an analysis for section 1102(b) of the Act.

Section 202 of the Unfunded Mandates Reform Act of 1995 also requires that agencies assess anticipated costs and benefits before issuing any rule that may result in expenditures in any 1 year by State, local, or tribal governments, in the aggregate, or by the private sector, of $110 million. This notice has no consequential effect on State, local, or tribal governments or on the private sector.

Executive Order 13132 establishes certain requirements that an agency must meet when it publishes a rule that imposes substantial direct requirement costs on State and local governments, preempts State law, or otherwise has Federalism implications. This notice will not have a substantial effect on State or local governments. There are no other alternatives at this time.

In accordance with the provisions of Executive Order 12866, this notice was reviewed by the Office of Management and Budget.

Authority: Sections 1102 and 1871 of the Social Security Act (42 U.S.C. 1302 and 1395hh).

(Catalog of Federal Domestic Assistance Program No. 93.774, Medicare—Supplementary Medical Insurance Program)

Dated: May 17, 2005.

Mark B. McClellan,

Administrator, Centers for Medicare & Medicaid Services.

Approved: August 25, 2005.

Michael O. Leavitt,

Secretary.

[FR Doc. 05–17278 Filed 8–26–05; 9:46 am]

BILLING CODE 4120–01–P

DEPARTMENT OF HEALTH AND HUMAN SERVICES

Administration for Children and Families

Family and Youth Services Bureau; Positive Youth Development State and Local Collaboration Demonstration Projects

AGENCY: Family and Youth Services Bureau, Administration for Children and Families (ACF), Department of Health and Human Services (HHS).

ACTION: Award announcement.

CFDA#: The Catalog of Federal Domestic Assistance (CFDA) number for this program is 93.623. The title is the Positive Youth Development State and Local Collaboration Demonstration Projects.

Legislative Authority: Grants for Runaway and Homeless Youth programs are authorized by the Runaway and Homeless Youth Act (title III of the Juvenile Justice and Delinquency Prevention Act of 1974), as amended by the Missing, Exploited, and Runaway Children Protection Act of 1999, (Pub. L. 106–71).

Amount of Award: $100,000 per grantee.

Project Period: 9/30/04–9/29/05.

SUMMARY: Notice is hereby given that a noncompetitive grant supplement is being made to the following state agencies: State of Nebraska Health & Human Services, University of Kentucky Research Foundation, State of Oregon, New York Office of Children & Family Services, State of Louisiana, Iowa Dept. of Human Rights Criminal & Juvenile Justice, Commonwealth of Massachusetts, Illinois Department of Human Services, Governor's Office for Children Youth & Families. The purpose of this supplement is to support collaborations between state-level agencies and local community jurisdictions regarding positive development opportunities available to young people as approved in their original planning grant.

FOR FURTHER INFORMATION CONTACT: Administration for Children and Families, Family and Youth Services Bureau, 330 C Street, SW., Washington, DC 20447, Courtney Workman—(202) 205–8657, *cworkman@acf.hhs.gov*.

Dated: August 22, 2005.

Joan E. Ohl,

Commissioner, Administration on Children, Youth and Families.

[FR Doc. 05–17371 Filed 8–31–05; 8:45 am]

BILLING CODE 4184–01–P

DEPARTMENT OF HEALTH AND HUMAN SERVICES

Food and Drug Administration

[Docket No. 2005N–0343]

Agency Emergency Processing Under Office of Management and Budget Review; Guidance for Requesting an Extension to Use Existing Label Stock After the Trans Fat Labeling Effective Date of January 1, 2006

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 emergency processing under the Paperwork Reduction Act of 1995 (the PRA). FDA is preparing a guidance document to notify the public of procedures being implemented by the agency to assist firms that wish to request, on a case-by-case basis upon an appropriate showing, an extension to use existing label stock after the effective date of the *trans* fat labeling final rule. This notice solicits comments on the proposed collection of information associated with the guidance document entitled ''Guidance for Requesting an Extension to Use Existing Label Stock After the *Trans* Fat Labeling Effective Date of January 1, 2006.''

DATES: Fax written comments on the collection of information by October 3, 2005. FDA is requesting approval of this emergency processing by September 8, 2005.

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 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: Peggy Robbins, Office of Management Programs (HFA–250), Food and Drug Administration, 5600 Fishers Lane, Rockville, MD 20857, 301–827–1223.

SUPPLEMENTARY INFORMATION: FDA has requested emergency processing of this proposed collection of information under section 3507(j) of the PRA (44 U.S.C. 3507(j)) and 5 CFR 1320.13. FDA issued a final rule (the *trans* fat final rule) on July 11, 2003 (68 FR 41434) to