DEPARTMENT OF HEALTH AND HUMAN SERVICES

Administration for Children and Families

Notice of the Award of a Single-Source Grant to The WorkPlace, Inc., in Bridgeport, CT

AGENCY: Office of Family Assistance, ACF, HHS.

ACTION: Award of a Single-source Grant to The WorkPlace, Inc., a local workforce investment board located in Bridgeport, CT.

Statutory Authority: Section 2008(a) of Title XX of the Social Security Act, as amended by Section 5507 of the Affordable Care Act (Pub. L. No. 111–148).

SUMMARY: The Administration for Children and Families (ACF), Office of Family Assistance (OFA), Health Profession Opportunity Grants (HPOG) program announces the award of a single-source grant (cooperative agreement) to The WorkPlace, Inc. a local, non-profit workforce investment board located in Bridgeport, CT. Award funds will support a program to provide education and training to Temporary Assistance to Needy Families (TANF) recipients, and other low-income individuals, for occupations in the health care field that pay well and are expected to either experience labor shortages or be in high demand.

The city of Bridgeport, CT, faces high levels of unemployment. The WorkPlace, Inc., proposes working with numerous community partners to coordinate referrals, conduct assessments, and provide remedial and life skills training, supportive services, and occupational skills training.

If performance by the grantee is deemed satisfactory and funds are available, the grantee may be awarded future funding in the form of annual noncompetitive continuation grants.

DATES: The project period for the award is September 30, 2011–September 29, 2012.

FOR FURTHER INFORMATION CONTACT: Stan Koutstaal, Program Manager, Office of Family Assistance, 370 L'Enfant Promenade, SW., Washington, DC 20447. *Telephone:* 202–401–5457; *E-mail: stanley.koutstaal@acf.hhs.gov.*

Dated: September 26, 2011.

Earl S. Johnson,

Director, Office of Family Assistance.

[FR Doc. 2011–27067 Filed 10–18–11; 8:45 am]

BILLING CODE 4184–35–P

DEPARTMENT OF HEALTH AND HUMAN SERVICES

Food and Drug Administration

[Docket No. FDA–2011–N–0494]

Agency Information Collection Activities; Submission for Office of Management and Budget Review; Comment Request; Data To Support Communications To Educate Consumers on How To Safely Purchase Drugs Online

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 November 18, 2011.

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:* 202–395–7285, or e-mailed to *oira_submission@omb.eop.gov.* All comments should be identified with the OMB control number 0910–NEW and title ''Data to Support Communications to Educate Consumers on How to Safely Purchase Drugs Online.'' Also include the FDA docket number found in brackets in the heading of this document.

FOR FURTHER INFORMATION CONTACT: Juanmanuel Vilela, Office of Information Management, Food and Drug Administration, 1350 Piccard Dr., PI50–400B, Rockville, MD 20850, 301–796–7651, *Juanmanuel.vilela@fda.hhs.gov.*

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.

Data To Support Communications To Educate Consumers on How To Safely Purchase Drugs Online—(OMB Control Number 0910–NEW)

FDA has planned an integrated public outreach campaign to improve the safe use of online pharmacies for drug purchases. In order to effectively evaluate this campaign, FDA must understand individuals' knowledge, attitudes, and practices with regard to online pharmacies both at the start of the campaign and on an ongoing basis. This will enable FDA to gauge progress toward educating the public on safely purchasing from online pharmacies. An online survey panel will be employed to collect this information, which serves the need for direct and quantitative measurement of our target population, and which, as a quantitative research tool has some major benefits:

- To focus on our target population of adults who use the Internet.
- To collect data quickly and efficiently with minimal cost to the government.
- To reduce burden to the public by providing a means to complete the survey at a time and place of their choosing.

FDA will use online data collection to establish a baseline and evaluate the success of its messages and distribution methods for its outreach campaign, which educates consumers about how to safely purchase drugs online. Additionally, FDA will use this method to help tailor messages and communications vehicles to have both a more powerful and desired impact on target audiences. The data will not be used for the purposes of making policy or regulatory decisions.

In the **Federal Register** of July 12, 2011 (76 FR 40920), FDA published a 60-day notice requesting public comment on the proposed collection of information. FDA received no comments.

FDA estimates the burden of this collection of information as follows:

TABLE 1—ESTIMATED ANNUAL REPORTING BURDEN [1]

Activity	Number of respondents	Number of responses per respondent	Total annual responses	Average burden per response	Total hours
Survey Study	5,000	1	5,000	.33 (20 min.)	1,650

[1] There are no capital costs or operating and maintenance costs associated with this collection of information.

Annually, FDA projects one survey study. FDA is requesting this data collection burden so as not to restrict the Agency's ability to gather information on public sentiment for its proposals in its regulatory and communications programs.

Dated: October 14, 2011.

Leslie Kux,

Acting Assistant Commissioner for Policy.

[FR Doc. 2011–27019 Filed 10–18–11; 8:45 am]

BILLING CODE 4160–01–P

DEPARTMENT OF HEALTH AND HUMAN SERVICES

Food and Drug Administration

[Docket No. FDA–2011–N–0510]

Agency Information Collection Activities; Submission for Office of Management and Budget Review; Comment Request; Substances Prohibited From Use in Animal Food or Feed

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 November 18, 2011.

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: 202–395–7285, or e-mailed to *oira_submission@omb.eop.gov.* All comments should be identified with the OMB control number 0910–0627. Also include the FDA docket number found in brackets in the heading of this document.

FOR FURTHER INFORMATION CONTACT: Juanmanuel Vilela, Office of Information Management, Food and Drug Administration, 1350 Piccard Dr., PI50–400B, Rockville, MD 20850, 301–796–7651, *juanmanuel.vilela@fda.hhs.gov*.

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.

Substances Prohibited From Use in Animal Food or Feed—21 CFR Part 589 (OMB Control Number 0910–0627)—(Extension)

The final rule on bovine spongiform encephalopathy (BSE) (73 FR 22720, April 25, 2008) prohibits the use of certain cattle origin materials in the food or feed of all animals to help prevent the spread of BSE in U.S. cattle. BSE is a progressive and fatal neurological disorder of cattle that results from an unconventional transmissible agent. BSE belongs to the family of diseases known as transmissible spongiform encephalopathies (TSEs). All TSEs affect the central nervous system of infected animals. These measures will further strengthen existing safeguards against BSE.

In the **Federal Register** of July 28, 2011 (76 FR 45259), FDA published a 60-day notice requesting public comment on the proposed collection of information. No comments were received on the information collection.

Description of Recordkeeping for Respondents: Rendering facilities, medicated feed manufacturers, livestock feeders.

FDA estimates the burden of this collection of information as follows:

TABLE 1—ESTIMATED ANNUAL RECORDKEEPING BURDEN [1]

21 CFR Section	Number of recordkeepers	Number of records per recordkeeper	Total annual records	Average burden per recordkeeper	Total hours	Total operating & maintenance costs
589.2001 (c)(2)(vi) and (c)(3)(i)	175	1	175	20	3,500	$59,500
589.2001 (c)(2)(ii)	50	1	50	20	1,000	17,000
589.2001 (c)(3)(i)(A)	175	1	175	26	4,550	80,580
Total					9,050	157,080

[1] There are no capital costs associated with this collection of information.

The number of recordkeepers times the number of records per recordkeeper equals total annual records. Total annual records times average burden per recordkeeper equals total hours.

Description of Respondents for Reporting: The final rule on BSE (73 FR 22720) included a provision that exempts cattle materials prohibited in animal feed (CMPAF) from designated countries from the prohibition on its use in animal feed (21 CFR 589.2001(b)(1)(vi)). A foreign country seeking this designation will submit a written request to FDA that includes a variety of information about the countries' BSE status (21 CFR 589.2001(f)). FDA estimates that 10 countries could submit a request to FDA to be exempted from CMPAF restrictions.

FDA estimates the reporting burden for this information collection as follows:

TABLE 2—ESTIMATED ONE-TIME AND RECURRING REPORTING BURDEN [1]

21 CFR Section	Number of respondents	Number of responses per respondent	Total annual responses	Average burden per response	Total hours
589.2001(b)(1) [2]	10	1	10	80	800
589.2001(f)	10	1	10	26	260

[1] There are no capital costs or operating costs associated with the collection of information.

[2] One-time burden.