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.


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.

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 November 18, 2011.

ADDRESSES: To ensure that comments on the information collection are received, FDA 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>
<thead>
<tr>
<th>Activity</th>
<th>Number of respondents</th>
<th>Number of responses per respondent</th>
<th>Total annual responses</th>
<th>Average burden per response</th>
<th>Total hours</th>
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<tbody>
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<td>Survey Study</td>
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<td>1</td>
<td>5,000</td>
<td>.33 (20 min.)</td>
<td>1,650</td>
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</tbody>
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1 There are no capital costs or operating and maintenance costs associated with this collection of information.