of current and recent TANF recipients. The survey will gather information from employers on their attitudes, practices, and policies toward TANF recipient and other low-skill hires, including information on worker advancement, the use of work force intermediaries in hiring, and the role that child care plays in worker retention. The survey will allow for comparisons of employers in urban-core areas, suburbs, and exurbs/rural areas. It will also measure employment outcomes for TANF recipients and other low-skilled workers, allowing us to draw connections between employer practices and employee outcomes. In short, this national survey of employers in the low-wage labor market can provide key information on what employer practices and policies are and how they are associated with workplace success for welfare recipients and other less-skilled workers.

Respondents: A nationally representative sample of business establishments having 4 or more workers.

ANNUAL BURDEN ESTIMATES

Instrument	Number of respondents	Number of responses per respondent	Average burden hours per response	Total burden hours
TANF Labor Market Survey	1,300	1	0.33	429

Estimated Total Annual Burden Hours: 429.

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, 725 17th Street, NW., Washington, DC 20503, Attn: Desk Officer for ACF, E-mail address: *Katherine_T._Astrich@omb.eop.gov.*

Dated: August 15, 2006.

Robert Sargis,

Reports Clearance Officer.

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DEPARTMENT OF HEALTH AND HUMAN SERVICES

Food and Drug Administration

[Docket No. 2006N–0320]

Molecular Methods in Immunohematology; Public Workshop

AGENCY: Food and Drug Administration, HHS.

ACTION: Notice of public workshop.

The Food and Drug Administration (FDA) is announcing a public workshop entitled ''Molecular Methods in Immunohematology.'' The purpose of the public workshop is to gather and review current information on scientific developments that might enhance immunohematologic testing of blood donor or patient blood samples as part of pre-transfusion compatibility testing, or in determination and management of feto-maternal blood group incompatibilities.

Date and Time: The public workshop will be held on September 25, 2006, from 8:30 a.m. to 5 p.m., and September 26, 2006, from 8:30 a.m. to 2 p.m.

Location: The public workshop will be held at the Lister Hill Center Auditorium, bldg. 38A, National Institutes of Health, 8800 Rockville Pike, Bethesda, MD 20894.

Contact Person: Rhonda Dawson, Center for Biologics Evaluation and Research (HFM–302), Food and Drug Administration, 1401 Rockville Pike, suite 200N, Rockville, MD 20852–1448, 301–827–6129, FAX: 301–827–2843, e-mail: *rhonda.dawson@fda.hhs.gov*.

Registration: Mail or fax your registration information (including name, title, firm name, address, telephone and fax numbers) to the contact person by September 15, 2006. There is no registration fee for the public workshop. Early registration is recommended because seating is limited. Registration on the day of the public workshop will be provided on a space available basis beginning at 8:00 a.m.

If you need special accommodations due to a disability, please contact Rhonda Dawson (see *Contact Person*) at least 7 days in advance of the workshop.

SUPPLEMENTARY INFORMATION: The public workshop will feature presentations by national and international experts from government, academic institutions, and industry. The main goal of the workshop is to determine potential applications of molecular methods to improve safety in transfusion medicine by overcoming current limitations in the field of immunohematology, namely, the lack of reagent grade antibodies, both polyclonal and monoclonal; variability of reactivity of monoclonal antibodies as compared to polyclonal antibodies; and inherent limitations in the hemagglutination test. Topics to be discussed include the following: (1) Use of molecular methods in platelet and leukocyte typing, (2) use of phage display technology in place of routine hemagglutination tests, (3) potential advantages of using molecular methods in donor screening and patient typing, (4) use of molecular methods to resolve unusual serologic findings, (5) potential use of molecular methods in the manufacture of immunohematology reagents, and (6) current limitations in the use of molecular methods.

Transcripts: Transcripts of the public workshop may be requested in writing from the Freedom of Information Office (HFI–35), Food and Drug Administration, 5600 Fishers Lane, rm. 6–30, Rockville, MD 20857, approximately 15 working days after the public workshop at a cost of 10 cents per page. A transcript of the public workshop will be available on the Internet at *http://www.fda.gov/cber/minutes/workshop-min.htm*.

Dated: August 14, 2006.

Jeffrey Shuren,

Assistant Commissioner for Policy.

[FR Doc. E6–13695 Filed 8–18–06; 8:45 am]

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