ACTION: Notice of a Bulletin.


Michael Robertson, Associate Administrator for Governmentwide Policy, Chief Acquisition Officer. [FR Doc. 2010–6610 Filed 3–24–10; 8:45 am]

BILLING CODE 6820–14–P

GENERAL SERVICES ADMINISTRATION

Federal Travel Regulation (FTR): Maximum Per Diem Rates for the States of Kansas, New Mexico, New York, Rhode Island, and Texas

AGENCY: Office of Governmentwide Policy, General Services Administration (GSA).

ACTION: Notice of Per Diem Bulletin 10–03, revised continental United States (CONUS) per diem rates.

SUMMARY: The General Services Administration (GSA) has reviewed the per diem rates for certain locations in the States of Kansas, New Mexico, New York, Rhode Island and Texas and determined that they are inadequate. FOR FURTHER INFORMATION CONTACT: For clarification of content, contact Ms. Jill Denning, Office of Governmentwide Policy, Travel Management Policy, at (202) 208–7642. Please cite FTR Per Diem Bulletin 10–03.

SUPPLEMENTARY INFORMATION:

A. Background

After an analysis of the per diem rates established for FY 2010 (see the Federal Register notice at 74 FR 42898, August 25, 2009, and FTR Bulletin 10–01), the per diem rate is being changed in the following locations:

- State of Kansas
  - Leavenworth County.
- State of New Mexico
  - Dona Ana County.
- State of New York
  - Oswego County.
- State of Rhode Island
  - Bristol County.
- State of Texas
  - Midland County.

Per diem rates are published on the Internet at www.gsa.gov/perdiem as FTR per diem bulletins. This process ensures timely increases or decreases in per diem rates established by GSA for Federal employees on official travel within CONUS. Notices published periodically in the Federal Register, such as this one, now constitute the only notification of revisions in CONUS per diem rates to agencies.

Dated: March 17, 2010.

Becky Rhodes, Deputy Associate Administrator, Office of Travel, Transportation and Asset Management. [FR Doc. 2010–6612 Filed 3–24–10; 8:45 am]

BILLING CODE 6820–14–P

DEPARTMENT OF HEALTH AND HUMAN SERVICES

Decision To Evaluate a Petition To Designate a Class of Employees for Revere Copper and Brass in Detroit, MI, To Be Included in the Special Exposure Cohort

AGENCY: National Institute for Occupational Safety and Health (NIOSH), Department of Health and Human Services (HHS).

ACTION: Notice.

SUMMARY: HHS gives notice as required by 42 CFR 3.12(e) of a decision to evaluate a petition to designate a class of employees for Revere Copper and Brass in Detroit, Michigan, to be included in the Special Exposure Cohort under the Energy Employees Occupational Illness Compensation Program Act of 2000. The initial proposed definition for the class being evaluated, subject to revision as warranted by the evaluation, is as follows:

- Facility: Revere Copper and Brass.
- Location: Detroit, Michigan.
- Job Titles and/or Job Duties: Extruders and Shapes Specialists who worked in the Rod and Shape Mill.
- Period of Employment: January 1, 1943 through December 31, 1984.

FOR FURTHER INFORMATION CONTACT: Stuart L. Hinnefeld, Interim Director, Division of Compensation Analysis and Support, National Institute for Occupational Safety and Health (NIOSH), 4676 Columbia Parkway, MS C–46, Cincinnati, OH 45226, Telephone 877–222–7570. Information requests can also be submitted by e-mail to DCAS@CDC.GOV.

John Howard, Director, National Institute for Occupational Safety and Health. [FR Doc. 2010–6636 Filed 3–24–10; 8:45 am]

BILLING CODE 4163–19–P

DEPARTMENT OF HEALTH AND HUMAN SERVICES

Administration for Children and Families

Proposed Information Collection Activity; Comment Request

Proposed Projects

Title: Data Collection Plan for the Customer Satisfaction Evaluation of Child Welfare Information Gateway. OMB No.: 0970–0303. Description: The National Clearinghouse on Child Abuse and Neglect Information (NCCAN) and the National Adoption Information Clearinghouse (NAIC) received OMB approval to collect data for a customer satisfaction evaluation under OMB control number 0970–0303. On June 20, 2006, NCCAN and NAIC were consolidated into Child Welfare Information Gateway (Information Gateway).

The proposed information collection activities include revisions to the Customer Satisfaction Evaluation approved under OMB control number 0970–0303 to reflect current information needs for providing innovative and useful products and services.

Child Welfare Information Gateway is a service of the Children’s Bureau, a component within the Administration for Children and Families, and Information Gateway is dedicated to the mission of connecting professionals and concerned citizens to information on programs, research, legislation, and statistics regarding the safety, permanency, and well-being of children and families. Information Gateway’s main functions are identifying information needs, locating and acquiring information, creating information, organizing and storing information, disseminating information,
and facilitating information exchange among professionals and concerned citizens. A number of vehicles are employed to accomplish these activities, including, but not limited to, Web site hosting, discussions with customers (e.g., phone, live chat, etc.), and dissemination of publications (both print and electronic).

The Customer Satisfaction Evaluation was initiated in response to Executive Order 12862 issued on September 11, 1993. The Order calls for putting customers first and striving for a customer-driven government that matches or exceeds the best service available in the private sector. To that end, Information Gateway’s evaluation is designed to better understand the kind and quality of services customers want, as well as customers’ level of satisfaction with existing services. The proposed data collection activities for the evaluation include customer satisfaction surveys, customer comment cards, selected publication surveys, and focus groups.


### ANNUAL BURDEN ESTIMATES

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<th>Instrument</th>
<th>Affected public</th>
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<th>Number of responses per respondent</th>
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In compliance with the requirements of Section 3506 (2)(A) of the Paperwork Reduction Act of 1995, the Administration for Children and Families is soliciting public comment on the specific aspects of the information collection described above. Copies of the proposed collection of information can be obtained and comments may be forwarded 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. E-mail address: infocollection@acf.hhs.gov. All requests should be identified by the title of the information collection.

The Department specifically requests comments on: (a) Whether the proposed collection of information is necessary for the proper performance of the functions of the agency, including whether the information shall have practical utility; (b) the accuracy of the agency’s estimate of the burden of the proposed collection of information; (c) the quality, utility, and clarity of the information to be collected; and (d) ways to minimize the burden of the collection of information on respondents, including through the use of automated collection techniques or other forms of information technology. Consideration will be given to comments and suggestions submitted within 60 days of this publication.

Dated: March 18, 2010.

Robert Sargis,
Reports Clearance Officer.
[FR Doc. 2010–6469 Filed 3–24–10; 8:45 am]
BILLING CODE 4184–01–M

DEPARTMENT OF HEALTH AND HUMAN SERVICES

Food and Drug Administration

Determination That DIDREX (Benzphetamine Hydrochloride) Tablets, 25 Milligrams, Were Not Withdrawn From Sale for Reasons of Safety or Effectiveness

AGENCY: Food and Drug Administration, HHS.

ACTIONS: Notice.

SUMMARY: The Food and Drug Administration (FDA) has determined that DIDREX (benzphetamine hydrochloride (HCl)) Tablets, 25 milligrams (mg), were not withdrawn from sale for reasons of safety or effectiveness. This determination will allow FDA to approve abbreviated new drug applications (ANDAs) for benzphetamine HCl 25 mg tablets, if all drug applications (ANDAs) for authorization the approval of duplicate drug products approved under an ANDA procedure. ANDA applicants must, with certain