study will be made available to the public at the conclusion of the evaluation. Construction and mining companies who participate in the study will be randomly assigned to receive eight weekly toolbox safety training sessions that use either a case-study narrative or conventional instructional approach. The training sessions are designed to last fifteen minutes. The impact of these materials will be evaluated through the examination of changes in employee knowledge gains, attitudes toward safety practices, and the use of safety behaviors prior to and following their participation in the safety training program. Trainers will complete brief response cards each week. A sample of trainers will participate in structured interviews. Findings of the study will be reported to participants and in the literature. The total annual burden for this data collection is 233 hours.

<table>
<thead>
<tr>
<th>Respondents</th>
<th>Number of respondents</th>
<th>Number of responses/responder</th>
<th>Average burden/response (in hours)</th>
</tr>
</thead>
<tbody>
<tr>
<td>Worker Pre-training Survey (attitude survey)</td>
<td>412</td>
<td>1</td>
<td>15/60</td>
</tr>
<tr>
<td>Worker Post-training Survey (attitude survey)</td>
<td>412</td>
<td>1</td>
<td>15/60</td>
</tr>
<tr>
<td>Instructor Feedback Cards</td>
<td>41</td>
<td>8</td>
<td>5/60</td>
</tr>
</tbody>
</table>


Thomas Bartenfeld,
Acting Associate Director for Policy, Planning and Evaluation, Centers for Disease Control and Prevention.

[FR Doc. 03–2277 Filed 1–30–03; 8:45 am]
BILLING CODE 4163–18–P

DEPARTMENT OF HEALTH AND HUMAN SERVICES

Centers for Medicare and Medicaid Services


Agency Information Collection Activities: Proposed Collection; Comment Request

AGENCY: Centers for Medicare and Medicaid Services, HHS.

In compliance with the requirement of section 3506(c)(2)(A) of the Paperwork Reduction Act of 1995, the Centers for Medicare and Medicaid Services (CMS) (formerly known as the Health Care Financing Administration (HCFA)), Department of Health and Human Services, is publishing the following summary of proposed collections for public comment. Interested persons are invited to send comments regarding this burden estimate or any other aspect of this collection of information, including any of the following subjects: (1) The necessity and utility of the proposed information collection for the proper performance of the agency’s functions; (2) the accuracy of the estimated burden; (3) ways to enhance the quality, utility, and clarity of the information to be collected; and (4) the use of automated collection techniques or other forms of information technology to minimize the information collection burden.

(1) Type of Information Collection Request: Extension of a currently approved collection; Title of Information Collection: Refinement of RHC Certification and QAPI and Supporting Regulations in 42 CFR 491.8 and 491.11; Form No.: CMS–R–242 (OMB# 0938–0792); Use: This collection contains information collection requirements concerning requests for additional waivers of staffing requirements and documentation of quality assessment and performance improvement programs; Frequency: Annually; Affected Public: Business or other for-profit; Number of Respondents: 3,528; Total Annual Responses: 3,573; Total Annual Hours: 3,663.

(2) Type of Information Collection Request: Extension of a currently approved collection; Title of Information Collection: Medicare Waiver Demonstration Application; Form No.: CMS–10069 (OMB# 0938–0880); Use: The Medicare Waiver Demonstration Application will be used to collect standard information needed to implement Congressionally mandated and administration high priority demonstrations. The application will be used to gather information about the characteristics of the applicant’s organization, benefits, and services they propose to offer, success in operating the model, and evidence that the model is likely to be successful in the Medicare program. The standard application will be used for all waiver demonstrations and will reduce the burden on applicants, provide for consistent and timely information collections across demonstration, and provide a user-friendly format for respondents; Frequency: On occasion; Affected Public: Business or other for-profit and not-for-profit institutions; Number of Respondents: 75; Total Annual Responses: 75; Total Annual Hours: 1600.

(3) Type of Information Collection Request: Extension of a currently approved collection; Title of Information Collection: Matching Grants to States for the Operation of High Risk Pools; Form No.: CMS–10078 (OMB# 0938–0887); Use: HHS/CMS is requiring this information as a condition of eligibility for grants that were authorized in the Trade Act of 2002 (Pub. L. 107–210). The information is necessary to determine if a state applicant meets the necessary eligibility criteria for a grant as required by the law. The respondents will be states that have a high risk pool as defined in section 2744(c)(2) of the Public Health Service Act. The grants will provide matching funds to states that incur losses in the operation of high risk pools. High risk pools are set up by states to provide health insurance to individuals that cannot obtain health insurance in the private market because of a history of illness; Frequency: On occasion; Affected Public: State, local, or tribal government; Number of Respondents: 20; Total Annual Responses: 20; Total Annual Hours: 800.

(4) Type of Information Collection Request: Extension of a currently approved collection; Title of Information Collection: Conditions of Coverage of Suppliers of End Stage Renal Disease (ESRD); Form No.: CMS–R–52 (OMB# 0938–0386); Use: This package is needed to encourage proper distribution and effective utilization of ESRD treatment sources while maintaining and improving the efficient delivery of care by physicians and dialysis facilities; Frequency: Annually; Affected Public: Business or other for-profit and Federal Government; Number of Respondents: 4,297; Total Annual Responses: 4,297; Total Annual Hours: 148,785.

(5) Type of Information Collection Request: Revision of a currently approved collection; Title of Information Collection: Information Collection Requirements in the Hospice Conditions Coverage. The following
DEPARTMENT OF HEALTH AND HUMAN SERVICES

Food and Drug Administration

Advisory Committees; Filing of Annual Reports

AGENCY: Food and Drug Administration, HHS.

ACTION: Notice.

SUMMARY: The Food and Drug Administration (FDA) is announcing that, as required by the Federal Advisory Committee Act, the agency has filed with the Library of Congress the annual reports of those FDA advisory committees that held closed meetings during fiscal year 2002.

ADDRESSES: Copies are available from the Dockets Management Branch (HFA–305), Food and Drug Administration, 5630 Fishers Lane, rm. 1061, Rockville, MD 20852.


Linda Arey Skladany, Associate Commissioner for External Relations. [FR Doc. 03–2294 Filed 1–30–03; 8:45 am] BILLING CODE 4160–01–S

Draft Guidance for Industry on Estrogen and Estrogen/Progestin Drug Products to Treat Vasomotor Symptoms and Vulvar and Vaginal Atrophy Symptoms—Recommendations for Clinical Evaluation; Availability

AGENCY: Food and Drug Administration, HHS.

ACTION: Notice.

SUMMARY: The Food and Drug Administration (FDA) is announcing the availability of a draft guidance for industry entitled “Estrogen and Estrogen/Progestin Drug Products to Treat Vasomotor Symptoms and Vulvar and Vaginal Atrophy Symptoms—Recommendations for Clinical Evaluation.” The agency is revising its guidance for industry entitled “Guidance for Clinical Evaluation of Combination Estrogen/Progestin-Containing Drug Products Used for Hormone Replacement Therapy of Postmenopausal Women,” which was issued in March 1995. Once finalized, this guidance will replace the 1995 guidance.

DATES: Submit written or electronic comments on the draft guidance by April 1, 2003. General comments on agency guidance documents are welcome at any time.

ADDRESSES: Submit written requests for single copies of the draft guidance to the Division of Drug Information (HFD–240), Center for Drug Evaluation and Research, Food and Drug Administration, 5600 Fishers Lane, Rockville, MD 20857. Send on self-addressed adhesive label to assist that office in processing your requests. Submit written comments on the draft guidance to the Dockets Management Branch (HFA–305), Food and Drug Administration, 5630 Fishers Lane, rm. 1061, Rockville, MD 20852. Submit electronic comments to http://