and public health advisories by surveying a sample of recipients. Subjects will receive a questionnaire to be completed and returned to FDA. The information to be collected will address how clearly actions for reducing risk are explained, the timeliness of the information, and whether the reader has taken any action to eliminate or reduce risk as a result of information in the alert. Subjects will also be asked whether they wish to receive future alerts electronically, as well as how the safety alert program might be improved.

The information collected will be used to shape FDA’s editorial policy for the safety alerts and public health advisories. Understanding how target audiences view these publications will aid in deciding what changes should be considered in their content, format, and method of dissemination.

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

<table>
<thead>
<tr>
<th>No. of Respondents</th>
<th>Annual Frequency per Response</th>
<th>Total Annual Records</th>
<th>Hours per Response</th>
<th>Total Hours</th>
</tr>
</thead>
<tbody>
<tr>
<td>308</td>
<td>3</td>
<td>924</td>
<td>.17</td>
<td>157</td>
</tr>
</tbody>
</table>

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

Based on the history of the safety alert and public health advisory program, it is estimated that an average of three collections will be conducted a year. The total burden of response time is estimated at 10 minutes per survey. This was derived by CDRH staff completing the survey and through discussions with the contacts in trade organizations.


Jeffrey Shuren,
Assistant Commissioner for Policy.

[FR Doc. 03–11773 Filed 5–12–03; 8:45 am]
CVM, and (3) in the laboratory trial of methods submitted to CVM. The document should also help in making decisions about appropriate methodology in various regulatory situations and ensuring consistency in work done for CVM’s purposes.

Information collection provisions described in this guidance have been approved under OMB control numbers 0910–0032 and 0910–0325.

II. Significance of Guidance

This level 1 guidance is being issued consistent with FDA’s good guidance practices regulation (21 CFR 10.115). The guidance represents the agency’s animal drug residues. The document does not create or confer any rights for or on any person and does not operate to bind FDA or the public. Alternative methods may be used as long as they satisfy the requirements of the applicable statutes and regulations.

III. Comments

As with all of FDA’s guidance, the public is encouraged to submit written or electronic comments with new data or other new information pertinent to this guidance. FDA periodically will review the comments in the docket and, where appropriate, will amend the guidance. The public will be notified of any such amendments through a notice in the Federal Register.

Interested persons may submit to the Dockets Management Branch (see ADDRESSES) written or electronic comments with new information which:

- Evaluate whether the proposed collection of information is necessary for the proper performance of the functions of the agency, including whether the information will have practical utility;
- Evaluate the accuracy of the agency’s estimate of the burden of the proposed collection of information, including the validity of the methodology and assumptions used;
- Enhance the quality, utility, and clarity of the information to be collected; and
- Minimize the burden of the collection of information on those who are to respond, including through the use of appropriate automated, electronic, mechanical, or other technological collection techniques or other forms of information technology, e.g., permitting electronic submissions of responses.

Analysis

Title: Vendor Information Site.
OMB Number: 1600—new collection.
Frequency: On occasion.

Affected Public: Individuals or households; businesses or other for-profit; not-for-profit institutions; farms; State, local or tribal government.
Number of Respondents: 20,000.
Estimated Time Per Respondent: 30 minutes for startup; 30 minutes for maintaining.
Total Burden Hours: 20,000.
Total Burden Cost (capital/startup): $25.00 per respondent; $500,000 annually.
total Burden Cost (operating/ maintaining): $25.00 per respondent, $500,000 annually.

Description: This web-based Vendor Information Site information collection will provide a uniform voluntary way companies can provide descriptions of their product-and-service ideas to DHS for enhancing homeland security.

Steve I. Cooper,
Chief Information Officer.
[FR Doc. 03–11855 Filed 5–8–03; 12:16 pm]
BILLING CODE 4410–10–M

DEPARTMENT OF HOMELAND SECURITY

Federal Emergency Management Agency

[FEMA–1459–DR]

Mississippi; Major Disaster and Related Determinations


ACTION: Notice.

SUMMARY: This is a notice of the Presidential declaration of a major disaster for the State of Mississippi (FEMA–1459–DR), dated April 24, 2003, and related determinations.


FOR FURTHER INFORMATION CONTACT:

SUPPLEMENTARY INFORMATION: Notice is hereby given that, in a letter dated April 24, 2003, the President declared a major disaster under the authority of the Robert T. Stafford Disaster Relief and Emergency Assistance Act, 42 U.S.C. 5121–5206 (the Stafford Act), as follows:

I have determined that the damage in certain areas of the State of Mississippi, resulting from severe storms, tornadoes, and flooding on April 6–14, 2003, is of sufficient severity and magnitude to warrant a major disaster declaration under the Robert T. Stafford Disaster Relief and Emergency Assistance Act, 42 U.S.C. 5121–5206 (the