TABLE 1. — ESTIMATED ANNUAL REPORTING BURDEN

<table>
<thead>
<tr>
<th>FDA Form No.</th>
<th>21 CFR Section</th>
<th>No. of Respondents</th>
<th>Annual Frequency per Response</th>
<th>Total Annual Responses</th>
<th>Hours per Response</th>
<th>Total Hours</th>
</tr>
</thead>
<tbody>
<tr>
<td>FDA 1815/Permits granted on certificates</td>
<td>1210.23</td>
<td>8</td>
<td>1</td>
<td>8</td>
<td>0.5</td>
<td>4</td>
</tr>
<tr>
<td>FDA 1993/Application of permit</td>
<td>1210.20</td>
<td>8</td>
<td>1</td>
<td>8</td>
<td>0.5</td>
<td>4</td>
</tr>
<tr>
<td>FDA 1994/Tuberculin test</td>
<td>1210.13</td>
<td>1</td>
<td>1</td>
<td>1</td>
<td>1</td>
<td>1</td>
</tr>
<tr>
<td>FDA 1995/Physical examination of cows</td>
<td>1210.12</td>
<td>1</td>
<td>1</td>
<td>1</td>
<td>1</td>
<td>1</td>
</tr>
<tr>
<td>FDA 1996/Sanitary inspection of dairy farms</td>
<td>1210.11</td>
<td>8</td>
<td>200</td>
<td>1,600</td>
<td>1.5</td>
<td>2,400</td>
</tr>
<tr>
<td>FDA 1997/Sanitary inspections of plants</td>
<td>1210.14</td>
<td>8</td>
<td>1</td>
<td>8</td>
<td>2.0</td>
<td>16</td>
</tr>
<tr>
<td><strong>Total</strong></td>
<td></td>
<td><strong>2,426</strong></td>
<td></td>
<td><strong>2,426</strong></td>
<td></td>
<td></td>
</tr>
</tbody>
</table>

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

TABLE 2. — ESTIMATED ANNUAL RECORDKEEPING BURDEN

<table>
<thead>
<tr>
<th>21 CFR Section</th>
<th>No. of Recordkeepers</th>
<th>Annual Frequency per Recordkeeping</th>
<th>Total Annual Records</th>
<th>Hours per Recordkeeper</th>
<th>Total Hours</th>
</tr>
</thead>
<tbody>
<tr>
<td>1210.15</td>
<td>8</td>
<td>1</td>
<td>8</td>
<td>.05</td>
<td>.4</td>
</tr>
</tbody>
</table>

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

These estimates are based on the number of current permit holders and the number of inquiries that FDA has received regarding requests for applications in the next 3 years.

No burden has been estimated for the tagging requirement in § 1210.22 because the information on the tag is either supplied by FDA (permit number) or is disclosed to third parties as a usual and customary part of the shipper’s normal business activities (type of product and shipper’s name and address). Under 5 CFR 1320.3(c)(2), the public disclosure of information originally supplied by the Federal Government to the recipient for the purpose of disclosure to the public is not a collection of information. Under 5 CFR 1320.3(b)(2), the time, effort, and financial resources necessary to comply with a collection of information are excluded from the burden estimate if the reporting, recordkeeping, or disclosure activities needed to comply are usual and customary because they would occur in the normal course of activities. The Secretary of Health and Human Services has the discretion to allow Form FDA 1815, a duly certified statement signed by an accredited official of a foreign government, to be submitted in lieu of Forms FDA 1994 and 1995.

Dated: April 5, 2002.

Margaret M. Dotzel,
Associate Commissioner for Policy.

[FR Doc. 02–7579 Filed 4–11–02; 8:45 am]
BILLING CODE 4160–01–S

DEPARTMENT OF HEALTH AND HUMAN SERVICES

Food and Drug Administration

Drug Manufacturing Inspections; Public Workshops; Correction

AGENCY: Food and Drug Administration, HHHS.

ACTION: Notice of public workshops; correction.

SUMMARY: The Food and Drug Administration (FDA) is correcting a notice that appeared in the Federal Register of March 29, 2002 (67 FR 15210). The document announced a series of workshops to discuss the application of a systems-based approach to drug manufacturing inspections. The document was published with an inadvertent error. This document corrects that error.

FOR FURTHER INFORMATION CONTACT: Doris B. Tucker, Office of Policy (HF–27), Food and Drug Administration, 5600 Fishers Lane, Rockville, MD 20857.

SUPPLEMENTARY INFORMATION: In FR Doc. 02–7579, appearing on page 15210 in the Federal Register of Friday, March 29, 2002, the following correction is made:

1. On page 15211, in the first column, in the fifth line from the bottom and also in the last line of the document “http://www.fda.gov.cder/calendar” is corrected to read “http://www.fda.gov/cder/workshop.htm.”

Dated: April 5, 2002.

Margaret M. Dotzel,
Associate Commissioner for Policy.

[FR Doc. 02–8839 Filed 4–11–02; 8:45 am]
BILLING CODE 4160–01–S

DEPARTMENT OF HEALTH AND HUMAN SERVICES

Health Resources and Services Administration

HRSA AIDS Advisory Committee; Notice of Meeting

In accordance with section 10(a)(2) of the Federal Advisory Committee Act (Public Law 92–463), announcement is made of the following National Advisory body scheduled to meet during the month of May 2002.

Name: HRSA AIDS Advisory Committee (HAAC).

Date and Time: May 30, 2002; 8:30 a.m.–5 p.m.; May 31, 2002; 8:30 a.m.–3:30 p.m.