TABLE 1.—ESTIMATED ANNUAL REPORTING BURDEN[1]

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

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

TABLE 2.—ESTIMATED ANNUAL RECORDKEEPING BURDEN[1]

21 CFR Section	No. of Recordkeepers	Annual Frequency per Recordkeeping	Total Annual Records	Hours per Recordkeeper	Total Hours
1210.15	8	1	8	.05	.4

[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. 02–8840 Filed 4–11–02; 8:45 am]

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DEPARTMENT OF HEALTH AND HUMAN SERVICES

Food and Drug Administration

Drug Manufacturing Inspections; Public Workshops; Correction

AGENCY: Food and Drug Administration, HHS.

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]

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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.