6. Assurance that the firm or individual representing the firm and submitting a certificate for signature to FDA is aware of and knows that they are subject to the provisions of section 1001 of Title 18, United States Code. This law provides that it is a criminal offense to knowingly and willfully make a false statement or alter or counterfeit documents in a matter within the jurisdiction of a U.S. agency.

In the **Federal Register** of February 28, 2001 (66 FR 12802), the agency requested comments on the proposed collection of information. One comment was received. In this comment there were two concerns regarding burden. The first was that States may incur more than ''information'' burden. The impact on a few States has been to retrieve inspection reports from FDA contracted inspections or from a State inspection. The second concern was that FDA ''assumed no operating or maintenance costs''. The burden on a company for placement on an EC required list is only the initial information asked for in the **Federal Register** notice. A company may inquire about the status during the review process for placement on the list but this is of their choosing.

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

TABLE 1.—ESTIMATED ANNUAL REPORTING BURDEN[1]

Products	No. of Respondents	Annual Frequency per Response	Total Annual Responses	Hours per Response	Total Hours
Shell eggs	10	1	10	0.25	2.5
Dairy	100	1	100	0.25	25
Game meat and meat products	10	1	10	0.25	2.5
Animal casings	15	1	15	0.25	3.75
Gelatin	6	1	6	0.25	1.5
Total					35.25

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

The estimated number of respondents is based on the volume of exports and responses received to date. The estimated number of yearly responses has decreased from the estimate in FDA's previous notice seeking comment for this collection of information (63 FR 29738, June 1, 1998) because the actual number of responses has been decreasing. Companies do not need to reapply unless they have a compliance problem. An estimate for processors that export gelatin also has been added because these processors are now being included in the listing process.

TABLE 2.—ESTIMATED ANNUAL RECORDKEEPING BURDEN[1]

Respondents	No. of Record-keepers	Annual Frequency of Recordkeeping	Total Annual Records	Hours per Record-keeper	Total Hours
Trade association	15	1	15	8	120
State	50	1	50	8	400
Total					520

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

The burden estimated for the trade associations assumes the trade associations will disseminate FDA's information request through mass mailings to their membership or publish it in their trade magazine or newsletter. The burden estimated for State authorities assumes dissemination of information to the processors or dissemination of information about processors to FDA.

Dated: May 29, 2001.

Margaret M. Dotzel,

Associate Commissioner for Policy.

[FR Doc. 01–13985 Filed 6–4–01; 8:45 am]

BILLING CODE 4160–01–S

DEPARTMENT OF HEALTH AND HUMAN SERVICES

Indian Health Service

Indians Into Medicine Program; Correction

AGENCY: Indian Health Services, HHS.

ACTION: Notice; correction.

SUMMARY: The Indian Health Service published a document in the **Federal Register** on May 18, 2001, concerning an application deadline of June 1, 2001, for the Indians Into Medicine Program. The document contained an incorrect deadline date.

FOR FURTHER INFORMATION CONTACT: Ms. Jacqueline Santiago, Chief, Loan Repayment Branch, Division of Health Professions Support, Indian Health Service, 12300 Twinbrook Parkway, Suite 100A, Rockville, MD 20852, Telephone 301–443–3396. (This is not a toll-free number.)

Correction

In the **Federal Register** of May 18, 2001, in FR Doc. 01–12529, on page 27665, in the third column, correct the **DATES** caption to read:

DATES: A. Application Receipt Date—An original and two (2) copies of the completed grant application must be submitted with all required documentation to the Grants Management Branch, Division of Acquisition and Grants Management, Twinbrook Building, Suite 100, 12300 Twinbrook Parkway, Rockville, Maryland 20852, by close of business June 18, 2001.

Dated: March 29, 2001.

Michel E. Lincoln,

Deputy Director.

[FR Doc. 01–13987 Filed 6–4–01; 8:45 am]

BILLING CODE 4160–16–M