and issues a periodic newsletter, and is the focal point for conference and meeting planning activities for the Bureau.

6. Division of State Systems (DSS) reviews, assesses, and inspects the planning, design and operation of State management information systems and approves advanced planning documents for automated data systems. The Division provides leadership for the provision of technical assistance to States on information systems projects and advances the use of computer technology in the administration of child welfare and social services programs by States. The Division reviews, analyzes, and approves/disapproves State requests for federal financial participation for automated systems development and activities which support child welfare programs, including foster care and adoption. It provides assistance to States in developing or modifying automation plans to conform to federal requirements. It monitors approved State system development activities and conducts periodic reviews to assure State compliance with regulatory requirements applicable to automated systems supported by Federal financial participation. It provides guidance to States on functional requirements for these automated information systems. It promotes interstate transfer of existing automated systems and provides assistance and guidance to improve ACYF's programs through the use of automated systems.

Dated: May 30, 2001.

James A. Harrell,

Acting Commissioner, Administration on Children, Youth and Families.

[FR Doc. 01–14058 Filed 6–4–01; 8:45 am]

BILLING CODE 4184–01–M

DEPARTMENT OF HEALTH AND HUMAN SERVICES

Agency for Toxic Substances and Disease Registry

Medical Testing Associated With Exposure to Asbestos; Meeting

The Agency for Toxic Substances and Disease Registry (ATSDR), Division of Health Studies (DHS) announces the following meeting:

Name: Medical Testing Associated with Exposure to Asbestos.

Times and Dates: 10:00 a.m.–6:00 p.m., June 18, 2001; 8:30 a.m.–5:00 p.m., June 19, 2001.

Place: Sheraton Buckhead Hotel; 3405 Lenox Road; Atlanta, GA 30326.

Status: Open to the public, limited only by the space available.

Purpose: This is a working group meeting to discuss issues related to initial and follow-up testing of persons with environmental and historic occupational exposure to asbestoform materials from vermiculite mined in Libby, MT.

Matters To Be Discussed: The agenda will include a discussion on the routes and duration of exposure to asbestoform materials through both historic environmental and occupational routes; commonly conducted screening tests; frequency and periodicity of follow-up testing; use of standard testing procedures; and testing of special and/or sensitive populations.

Agenda items are subject to change as priorities dictate.

Contact Person for More Information: Jeffrey A. Lybarger, M.D., director, Division of Health Studies, ATSDR, 1600 Clifton Road, NE, m/s E31, Atlanta, Georgia 30333. Telephone 404/639–6200.

The Director, Management Analysis and Services office has been delegated the authority to sign **Federal Register** notices pertaining to announcements of meetings and other committee management activities for both the Centers for Disease Control and Prevention and the Agency for Toxic Substances and Disease Registry.

Dated: May 24, 2001.

Carolyn J. Russell,

Director, Management Analysis and Services Office, Centers for Disease Control and Prevention.

[FR Doc. 01–14050 Filed 6–4–01; 8:45 am]

BILLING CODE 4163–70–P

DEPARTMENT OF HEALTH AND HUMAN SERVICES

Food and Drug Administration

[Docket No. 01N–0069]

Agency Information Collection Activities; Submission for OMB Review; Comment Request; Information From U.S. Processors That Export to the European Community

AGENCY: Food and Drug Administration, HHS.

ACTION: Notice.

SUMMARY: The Food and Drug Administration (FDA) is announcing that the proposed collection of information listed below has been submitted to the Office of Management and Budget (OMB) for review and clearance under the Paperwork Reduction Act of 1995.

DATES: Submit written comments on the collection of information by July 5, 2001.

ADDRESSES: Submit written comments on the collection of information to the Office of Information and Regulatory Affairs, OMB, New Executive Office Bldg., 725 17th St. NW., rm. 10235, Washington, DC 20503, Attn: Wendy Taylor, Desk Officer for FDA.

FOR FURTHER INFORMATION CONTACT: Peggy Schlosburg, Office of Information Resources Management (HFA–250), Food and Drug Administration, 5600 Fishers Lane, Rockville, MD 20857, 301–827–1223.

SUPPLEMENTARY INFORMATION: In compliance with 44 U.S.C. 3507, FDA has submitted the following proposed collection of information to OMB for review and clearance.

Request for Information From U.S. Processors That Export to the European Community (OMB Control Number 0910–0320)—Extension

The European Community (EC) is a group of 15 European countries that have agreed to harmonize their commodity requirements to facilitate commerce among member States. EC legislation for intra-EC trade has been extended to trade with non-EC countries, including the United States. For certain food products, including those listed below in this document, EC legislation requires assurances from the responsible authority of the country of origin that the processor of the food is in compliance with applicable regulatory requirements.

With the assistance of trade associations and State authorities, FDA requests information from processors that export certain animal-derived products (e.g., shell eggs, dairy products, game meat, game meat products, animal casings, and gelatin) to EC. FDA uses the information to maintain lists of processors that have demonstrated current compliance with U.S. requirements and provides the lists to EC quarterly. Inclusion on the list is voluntary. EC member countries refer to the lists at ports of entry to verify that products offered for importation to EC from the United States are from processors that meet U.S. regulatory requirements. Products processed by firms not on the list are subject to detention and possible refusal at the port. FDA requests the following information from each processor:

1. Business name and address;
2. Name and telephone number of person designated as business contact;
3. Lists of products presently being shipped to EC and those intended to be shipped in the next 6 months;
4. Name and address of manufacturing plants for each product;
5. Names and affiliations of any Federal, State, or local governmental agencies that inspect the plant, government-assigned plant identifier, such as plant number, and last date of inspection; and

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