

reports on the physical examination of herds, while § 1210.13 requires the reporting of tuberculin testing of the herds. In addition, the regulations in part 1210 require that dairy farmers and plants maintain pasteurization records (§ 1210.15) and that each container of

milk or cream imported into the United States bear a tag with the product type, permit number, and shipper's name and address (§ 1210.22). Section 1210.20 requires that an application for a permit to ship or transport milk or cream into the United States be made by the actual

shipper. Section 1210.23 allows permits to be granted based on certificates from accredited officials.

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

TABLE 1—ESTIMATED ANNUAL REPORTING BURDEN¹

21 CFR Section	Form No.	Number of respondents	Number of responses per respondent	Total annual responses	Average burden per response	Total hours
1210.11	FDA 1996/Sanitary inspection of dairy farms.	2	200	400	1.5	600
1210.12	FDA 1995/Physical examination of cows.	1	1	1	0.5	0.5
1210.13	FDA 1994/Tuberculin test	1	1	1	0.5	0.5
1210.14	FDA 1997/Sanitary inspections of plants.	2	1	2	2	4
1210.20	FDA 1993/Application for permit	2	1	2	0.5	1
1210.23	FDA 1815/Permits granted on certificates.	2	1	2	0.5	1
Total	607

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

TABLE 2—ESTIMATED ANNUAL RECORDKEEPING BURDEN¹

21 CFR Section	Number of recordkeepers	Number of records per recordkeeper	Total annual records	Average burden per recordkeeping	Total hours
1210.15	2	1	2	0.05	0.10

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

The estimated number of respondents and hours per response are based on FDA's experience with the import milk permit program and the average number of import milk permit holders over the past 3 years. FDA estimates that two respondents will submit approximately 200 Form FDA 1996 reports annually, for a total of 600 responses. FDA estimates the reporting burden to be 1.5 hours per response, for a total burden of 607 hours.

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. To date, Form FDA 1815 has been submitted in lieu of these forms. Because FDA has not received any Forms FDA 1994 and 1995 in the last 3 years, the Agency estimates no more than one will be submitted annually. FDA estimates the reporting burden for each to be 0.5 hours per response for a total burden reporting burden of 0.5 hours each.

FDA estimates that two respondents will submit one Form FDA 1997 report annually, for a total of two responses. FDA estimates the reporting burden to be 2.0 hours per response, for a total

burden of 4 hours. FDA estimates that two respondents will submit one Form FDA 1993 report annually, for a total of two responses. FDA estimates the reporting burden to be 0.5 hours per response, for a total burden of 1 hour. FDA estimates that two respondents will submit one Form FDA 1815 report annually, for a total of two responses. FDA estimates the reporting burden to be 0.5 hours per response, for a total burden of 1 hour.

With regard to records maintenance, FDA estimates that approximately two recordkeepers will spend 0.05 hours annually maintaining the additional pasteurization records required by § 1210.15, for a total of 0.10 hours annually.

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, 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 business activities.

Dated: April 16, 2012.

Leslie Kux,

Assistant Commissioner for Policy.

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DEPARTMENT OF HEALTH AND HUMAN SERVICES

Centers for Disease Control and Prevention

Health Resources and Services Administration

CDC/HRSA Advisory Committee on HIV and STD Prevention and Treatment

In accordance with section 10(a)(2) of the Federal Advisory Committee Act (Pub. L. 92–463), the Centers for Disease Control and Prevention (CDC) and the Health Resources and Services

Administration (HRSA) announce the following meeting of the aforementioned committee:

Times and Dates:

8 a.m.–5:30 p.m., May 8, 2012.

8 a.m.–3 p.m., May 9, 2012.

Place: The Westin Buckhead Atlanta, 3391 Peachtree Road, NE.,

Atlanta, Georgia, 30326, Telephone: (404) 365–0065.

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

The meeting room will accommodate approximately 100 people.

Purpose: This Committee is charged with advising the Director, CDC and the Administrator, HRSA, regarding activities related to prevention and control of HIV/AIDS and other STDs, the support of health care services to persons living with HIV/AIDS, and education of health professionals and the public about HIV/AIDS and other STDs.

Matters To Be Discussed: Agenda items include: (1) Enhancing Hepatitis Prevention Treatment and Care in the United States;

(2) Integrating HIV Prevention and Care Data Systems; (3) External Peer Review of CDC Youth HIV/STI Prevention and Sexual Health Activities; (4) Preparing for the Ryan White Reauthorization; and (5) CHAC Workgroups Update.

Agenda items are subject to change as priorities dictate.

Contact Person for More Information: Margie Scott-Cseh, CDC, National Center for HIV/AIDS, Viral Hepatitis, STD, and TB Prevention, 1600 Clifton Road, NE., Mailstop E–07, Atlanta, Georgia 30333, Telephone (404) 639–8317.

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: April 16, 2012.

Elaine L. Baker,

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

[FR Doc. 2012–9549 Filed 4–19–12; 8:45 am]

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DEPARTMENT OF HOMELAND SECURITY

U.S. Citizenship and Immigration Services

Agency Information Collection

Activities: Form I–361, Extension of a Currently Approved Information Collection; Comment Request

ACTION: 30-Day Notice of Information Collection Under Review: Form I–361, Affidavit of Financial Support and

Intent to Petition for Legal Custody for Public Law 97–359 Amerasian.

The Department of Homeland Security, U.S. Citizenship and Immigration Services (USCIS) will be submitting the following information collection request to the Office of Management and Budget (OMB) for review and clearance in accordance with the Paperwork Reduction Act of 1995. The information collection notice was previously published in the **Federal Register** on February 16, 2012, at 77 FR 9259, allowing for a 60-day public comment period. USCIS did not receive any comments for this information collection.

The purpose of this notice is to allow an additional 30 days for public comments. Comments are encouraged and will be accepted until May 21, 2012. This process is conducted in accordance with 5 CFR 1320.10.

Written comments and/or suggestions regarding the item(s) contained in this notice, especially regarding the estimated public burden and associated response time, should be directed to the Department of Homeland Security (DHS), and to the Office of Information and Regulatory Affairs, Office of Management and Budget (OMB), USCIS Desk Officer. Comments may be submitted to: USCIS, Chief Regulatory Coordinator, Regulatory Coordination Division, Office of Policy and Strategy, Clearance Office, 20 Massachusetts Avenue, Washington, DC 20529. Comments may also be submitted to DHS via facsimile at 202–272–0997 or via email at uscisfrcomment@dhs.gov, and to the OMB USCIS Desk Officer via facsimile at 202–395–6974 or via email at oira_submission@omb.eop.gov.

When submitting comments by email please make sure to add OMB Control Number 1615–0021 in the subject box. Written comments and suggestions from the public and affected agencies should address one or more of the following four points:

(1) Evaluate whether the collection of information is necessary for the proper performance of the functions of the agency, including whether the information will have practical utility;

(2) Evaluate the accuracy of the agencies estimate of the burden of the collection of information, including the validity of the methodology and assumptions used;

(3) Enhance the quality, utility, and clarity of the information to be collected; and

(4) 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 submission of responses.

Overview of This Information Collection

(1) *Type of Information Collection:* Extension of a currently approved information collection.

(2) *Title of the Form/Collection:* Affidavit of Financial Support and Intent to Petition for Legal Custody for Public Law 97–359 Amerasian.

(3) *Agency form number, if any, and the applicable component of the Department of Homeland Security sponsoring the collection:* Form I–361. U.S. Citizenship and Immigration Services.

(4) *Affected public who will be asked or required to respond, as well as a brief abstract:* *Primary:* Individuals and households. The information on this form is used in support of Form I–360 (Petition for Amerasian, Widow(er), or Special Immigrant) to ensure financial support for Public Law 97–359 Amerasian. The affidavit is used only to sponsor individuals eligible for immigration under Public Law 97–359.

(5) *An estimate of the total number of respondents and the amount of time estimated for an average respondent to respond:* 50 responses at 30 minutes (.50) per response.

(6) *An estimate of the total public burden (in hours) associated with the collection:* 25 annual burden hours.

If you have additional comments, suggestions, or need a copy of the information collection instrument, please visit the USCIS Web site at: <http://www.regulations.gov/>.

We may also be contacted at: USCIS, Regulatory Coordination Division, Office of Policy and Strategy, 20 Massachusetts Avenue NW., Washington, DC 20529, Telephone number 202–272–1470.

Dated: April 16, 2012.

Laura Dawkins,

Acting Chief Regulatory Coordinator, Regulatory Coordination Division, Office of Policy and Strategy, U.S. Citizenship and Immigration Services, Department of Homeland Security.

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