

reasonable basis for these claims. The complaint also alleges that respondents falsely claimed that scientific research proves that Native Essense Plus prevents breast cancer, and that scientific studies prove that Native Essense with Cat's Claw is effective in the treatment of cancer.

Regarding chaparral herb, the Commission's complaint alleges that respondents claimed that chaparral herb is effective in treating and curing cancer, is effective in causing people with cancer to go into complete remission without the need for any other form of treatment, and is effective in shrinking or eliminating cancerous tumors. The complaint alleges that respondents lacked a reasonable basis for these claims.

The complaint also alleges that respondents lacked a reasonable basis for the claims that Mai-T Mushroom Plus is effective in preventing, treating and curing cancer, including but not limited to lung cancer, stomach cancer, hepatocellular cancer, leukemia, and Kaposi's sarcoma; and that Mai-T Mushroom Plus is effective in inhibiting the growth of cancerous tumors. Finally, the complaint alleges that respondents falsely claimed that clinical studies prove that Maitake mushrooms and Mai-T Mushroom Plus prevent and treat lung cancer, stomach cancer, hepatocellular cancer, leukemia, and Kaposi's sarcoma, and inhibit tumor growth.

The proposed consent order contains provisions designed to prevent respondents from engaging in similar acts and practices in the future. Part I requires respondents to have competent and reliable scientific evidence substantiating any claim that Native Essense (Rene Caisse) Formula tea or extract, Native Essense Plus tea or extract, Native Essense with Cat's Claw tea or extract, chaparral herb (or any product containing chaparral herb), Maitake mushroom extract, or Mai-T Mushroom Plus Formula extract, or any other covered product or service, is effective in the treatment or cure of cancer; prevents or lowers the risk of cancer; is effective in reducing the size of, or eliminating, cancerous tumors; or is safe or non-toxic or has no side effects. A "covered product or service" is defined as any food, dietary supplement, or drug, including, but not limited to any of the above products, or any other health-related product, service, or program.

Part II requires that any future claim about the efficacy, performance, or health-related benefits of any covered product or service be truthful and supported by competent and reliable scientific evidence. Part III requires that

respondents, in connection with the advertising of any product, shall not misrepresent, in any manner, expressly or by implication, the existence, contents, validity, results, conclusions, or interpretations of any test, study, or research.

Part IV of the proposed order provides that the order does not prohibit respondents from making representations for any drug that are permitted in labeling for the drug under any tentative or final Food and Drug Administration ("FDA") standard or under any new drug application approved by the FDA, and representations for any product that are specifically permitted in labeling for that product by regulations issued by the FDA under the Nutrition Labeling and Education Act of 1990.

Part V of the proposed order requires respondents to compile a list of all consumers who purchased Native Essense (Rene Caisse) Formula tea or extract, Native Essense Plus tea or extract, Native Essense with Cat's Claw tea or extract, chaparral herb (or any product containing chaparral herb), Maitake mushroom extract, or Mai-T Mushroom Plus Formula extract from respondents since July 1, 2005, and to mail a letter (attached to the proposed order as Attachment A) to each such purchaser describing the scientific evidence related to these products. Part V also prohibits respondents from providing any identifying information about these purchasers to anyone other than the Commission, another law enforcement agency, or as required by law.

Part VI of the proposed order requires respondents to keep copies of relevant advertisements and materials that substantiate claims made in the advertisements. Part VII requires respondents to provide copies of the order to certain of their employees. Part VIII requires the corporate respondent to notify the Commission at least thirty days prior to any change in the corporation that may affect compliance obligations arising under this order. Part IX requires the individual respondents to notify the Commission of their affiliation with any new business or employment. Part X requires respondents to file compliance reports with the Commission. Part XI of the proposed order is a "sunset" provision, dictating that the order will terminate twenty years from the date it is issued or twenty years after a complaint is filed in federal court, by either the United States or the FTC, alleging any violation of the order.

The purpose of this analysis is to facilitate public comment on the

proposed order. It is not intended to constitute an official interpretation of the agreement and proposed order or to modify in any way their terms.

By direction of the Commission.

Richard C. Donohue,

Acting Secretary.

[FR Doc. E9-8140 Filed 4-8-09; 8:45 am]

BILLING CODE: 6750-01-S

DEPARTMENT OF HEALTH AND HUMAN SERVICES

Centers for Disease Control and Prevention

Advisory Council for the Elimination of Tuberculosis: Notice of Charter Renewal

This gives notice under the Federal Advisory Committee Act (Pub. L. 92-463) of October 6, 1972, that the Advisory Council for the Elimination of Tuberculosis, Department of Health and Human Services, has been renewed for a 2-year period through March 15, 2011.

For information, contact Hazel Dean, Sc.D., M.P.H., Executive Secretary, Advisory Council for the Elimination of Tuberculosis, Department of Health and Human Services, 1600 Clifton Road, NE., Mailstop E-10, Atlanta, Georgia 30333, telephone 404/639-8000 or fax 404/639-8600.

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: March 31, 2009.

Elaine L. Baker,

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

[FR Doc. E9-8072 Filed 4-8-09; 8:45 am]

BILLING CODE 4163-18-P

DEPARTMENT OF HEALTH AND HUMAN SERVICES

Centers for Disease Control and Prevention

CDC/Health Resources and Services Administration (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 CDC announces

the following meeting of the aforementioned committee:

Times and Dates:

8 a.m.–5 p.m., May 19, 2009

8 a.m.–3 p.m., May 20, 2009

Place: JW Marriott Buckhead, 3300 Lenox Road, Atlanta, Georgia 30326, *Telephone:* (404) 262–3344, *Fax:* (404) 262–8689.

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 issues pertaining to: (1) Updates on HIV Testing, Syphilis Elimination and Viral Hepatitis Prevention; (2) Preventing HIV, STD, Hepatitis and TB in Correctional Settings; (3) Update on HIV Surveillance; and (4) Challenges and Opportunities to STD Prevention. Agenda items are subject to change as priorities dictate.

Contact Person for More Information: Margie Scott-Cseh, Committee Management Specialist, CDC, Strategic Business Unit, 1600 Clifton Road, NE., Mailstop E-07, Atlanta, Georgia 30333, *Telephone:* (404)

639–8317, *Fax:* (404) 639–8600, *E-mail:* zkr7@cdc.gov.

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 CDC and the Agency for Toxic Substances and Disease Registry.

Dated: April 3, 2009.

Elaine L. Baker,

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

[FR Doc. E9–8071 Filed 4–8–09; 8:45 am]

BILLING CODE 4163–18–P

DEPARTMENT OF HEALTH AND HUMAN SERVICES

Administration for Children and Families

Submission for OMB Review; Comment Request

Title: Court Improvement Program.
OMB No.: 0970–0245.

Description: The Court Improvement Program provides grants to State court systems to conduct assessments of their

foster care and adoption laws and judicial processes and to develop and implement a plan for system improvement. ACF proposes to collect information from the States about this program (applications, program reports) by way of a Program Instruction, which (1) describes the requirements for States under the reauthorization of the Court Improvement Program; (2) outlines the programmatic and fiscal provisions and reporting requirements of the program; (3) specifies the application submittal and approval procedures for the program for Fiscal Years 2007 through 2011; and (4) identifies technical resources for use by State courts during the course of the program. This Program Instruction contains information collection requirements pursuant to receiving a grant award that are found in Public Law 103–66, as amended by Public Law 105–89, Public Law 107–133, Public Law 109–239, and Public Law 109–288. The agency will use the information received to ensure compliance with the statute and provide training and technical assistance to the grantees.

Respondents: State Courts.

ANNUAL BURDEN ESTIMATES

Instrument	Number of respondents	Number of responses per respondent	Average burden hours per response	Total burden hours
Application	52	1	40	2,080
Annual program report	52	1	36	1,872

Estimated Total Annual Burden Hours: 3,952.

Additional Information: Copies of the proposed collection may be obtained by writing to the Administration for Children and Families, Office of Information Services, 370 L'Enfant Promenade, SW., Washington, DC 20447, *Attn:* ACF Reports Clearance Officer. All requests should be identified by the title of the information collection. *E-mail address:* infocollection@acf.hhs.gov.

OMB Comment: OMB is required to make a decision concerning the collection of information between 30 and 60 days after publication of this document in the **Federal Register**. Therefore, a comment is best assured of having its full effect if OMB receives it within 30 days of publication. Written comments and recommendations for the proposed information collection should be sent directly to the following: Office of Management and Budget, Paperwork

Reduction Project, *Fax:* 202–395–6974, *Attn:* Desk Officer for the Administration for Children and Families.

Dated: April 2, 2009.

Janean Chambers,

Reports Clearance Officer.

[FR Doc. E9–8008 Filed 4–8–09; 8:45 am]

BILLING CODE 4184–01–M

DEPARTMENT OF HEALTH AND HUMAN SERVICES

Administration for Children and Families

Proposed Information Collection Activity; Comment Request Proposed Projects

Title: Territory TANF Financial Report (ACF–196TR).

OMB No.: New collection.

Description: Authority to collect and report this information is found in the

Personal Responsibility and Work Opportunity Reconciliation Act of 1996 (PRWORA), Public Law 104–193. Territories with approved plans for implementation of the TANF program are required by statute to report financial data. Form ACF–196TR provides for collection of Federal expenditures data. Failure to collect this data would seriously compromise the Administration for Children and Families (ACF) ability to monitor expenditures and maintain financial management of the Territories TANF program. The financial data collected is also used to estimate outlays and may be used to prepare ACF budget submissions to Congress. Federal policy requires the strictest controls on funding requirements, which necessitates review of documentation in support of Territories expenditures for reimbursement.

Respondents: All Territories TANF Agencies.