

The National Vital Statistics Report forms provide counts of monthly occurrences of births, deaths, infant deaths, marriages, and divorces. Similar data have been published since 1937 and are the sole source of these data at the National level. The data are used by the Department of Health and Human Services and by other government, academic, and private research and commercial organizations in tracking changes in trends of vital events.

Respondents for the Monthly Vital Statistics Report Form are registration officials in each State and Territory, the District of Columbia, and New York City; in addition, 60 local (county) officials in New Mexico who record marriages occurring and divorces and annulments granted in each county of New Mexico will use this form. This form is designed to collect counts of monthly occurrences of births, deaths,

infant deaths, marriages, and divorces immediately following the month of occurrence.

The Annual Marriage and Divorce Statistical Report Form collects final annual counts of marriages and divorces by month for the United States and for each State. The statistical counts requested on this form differ from provisional estimates obtained on the Monthly Vital Statistics Report Form in that they represent complete counts of marriages, divorces, and annulments occurring during the months of the prior year. These final counts are usually available from State or county officials about eight months after the end of the data year. The data are widely used by government, academic, private research, and commercial organizations in tracking changes in trends of family formation and dissolution.

Respondents for the Annual Marriage and Divorce Statistical Report Form are registration officials in each State, the District of Columbia, New York City, Guam, Puerto Rico, Virgin Islands, Northern Marianas, and American Samoa. In addition, counts of marriages will be collected from individual counties in New Mexico, and counts of divorces will be collected from individual counties in California, Colorado, Indiana, Louisiana, New Mexico, and the boroughs of New York City due to a lack of centralized complete collections in these registration areas.

This submission requests approval for three years. There are no costs to respondents other than their time; the data are routinely available in each reporting office as a by-product of ongoing activities.

ESTIMATED ANNUALIZED BURDEN TABLE

Type of respondents	Form name	Number of respondents	Number of responses per respondent	Average burden per response (in hours)	Total burden (in hours)
State, Territory and New Mexico County officials.	Monthly Vital Statistics Report	117	12	30/60	702
State, Territory and selected County, Borough and City officials.	Annual Marriage and Divorce Statistical Report.	404	1	30/60	202
Total					904

Dated: October 20, 2008.

Maryam I. Daneshvar,

Acting Reports Clearance Officer, Office of the Chief Science Officer, Centers for Disease Control and Prevention.

[FR Doc. E8-25779 Filed 10-28-08; 8:45 am]

BILLING CODE 4163-18-P?<

DEPARTMENT OF HEALTH AND HUMAN SERVICES

Administration for Children and Families

Proposed Information Collection Activity; Comment Request

Proposed Projects

Title: Help America Vote Act (HAVA) Voting Access Annual Report.
OMB No.: 0970-0327.

Description: This is a revision to include the application for the

previously cleared Help America Vote Act (HAVA) Annual Report, Payments to States and Units of Local Government, 42 U.S.C. 15421.

The Help America Vote Act (HAVA) application to States and Units of Local Government is required by federal statute and regulation. Each State or Unit of Local Government must prepare an application to receive funds under the Help America Vote Act (HAVA), Public Law 107-252, Title II, Subtitle D, Part 2, Sections 261 to 265, Payments to States and Units of Local Government to Assure Access for Individuals with Disabilities (42 U.S.C. 15421-25). The application is provided in writing to the Administration for Children and Families, Administration on Developmental Disabilities.

An annual report is required by Federal statute (the Help America Vote Act (HAVA) of 2002, Public Law 107-

252, Section 261, Payments to States and Units of Local Government, 42 U.S.C. 15421). Each State or Unit of Local Government must prepare and submit an annual report at the end of every fiscal year. The report addresses the activities conducted with the funds provided during the year. The information collected from the annual report will be aggregated into an annual profile of how States have utilized the funds and establish best practices for election officials. It will also provide an overview of the State election goals and accomplishments and permit the Administration on Developmental Disabilities to track voting progress to monitor grant activities.

Respondents: Secretaries of State, Directors, State Election Boards, State Chief Election officials.

ANNUAL BURDEN ESTIMATES

Instrument	Number of respondents	Number of responses per respondent	Average burden hours per response	Total burden hours
Help America Vote Act (HAVA) Voting Access Annual Report	50	1	24	1,200
Help America Vote Act (HAVA) Voting Access Application	55	1	50	2,750
Estimated Total Annual Burden Hours				3,950

In compliance with the requirements of Section 506(c)(2)(A) of the Paperwork Reduction Act of 1995, the Administration for Children and Families is soliciting public comment on the specific aspects of the information collection described above. Copies of the proposed collection of information can be obtained and comments may be forwarded by writing to the Administration for Children and Families, Office of Administration, Office of Information Services, 370 L'Enfant Promenade, SW., Washington, DC 20447, Attn: ACF Reports Clearance Officer. E-mail address: infocollection@acf.hhs.gov. All requests should be identified by the title of the information collection.

The Department specifically requests comments on: (a) Whether the proposed collection of information is necessary for the proper performance of the functions of the agency, including whether the information shall have practical utility; (b) the accuracy of the agency's estimate of the burden of the proposed collection of information; (c) the quality, utility, and clarity of the information to be collected; and (d) ways to minimize the burden of the collection of information on respondents, including through the use of automated collection techniques or other forms of information technology. Consideration will be given to comments and suggestions submitted within 60 days of this publication.

Dated: October 24, 2008.

Janean Chambers,

Reports Clearance Officer.

[FR Doc. E8-25752 Filed 10-28-08; 8:45 am]

BILLING CODE 4184-01-P

DEPARTMENT OF HEALTH AND HUMAN SERVICES

Food and Drug Administration

[Docket No. FDA-2008-D-0030] (formerly Docket No. 2004D-0466)

Agency Information Collection Activities; Announcement of Office of Management and Budget Approval; Substantiation for Dietary Supplement Claims Made Under the Federal Food, Drug, and Cosmetic Act

AGENCY: Food and Drug Administration, HHS.

ACTION: Notice.

SUMMARY: The Food and Drug Administration (FDA) is announcing that a collection of information entitled "Substantiation for Dietary Supplement Claims Made Under the Federal Food, Drug, and Cosmetic Act" has been approved by the Office of Management and Budget (OMB) under the Paperwork Reduction Act of 1995.

FOR FURTHER INFORMATION CONTACT:

Jonna Capezzuto, Office of Information Management (HFA-710), Food and Drug Administration, 5600 Fishers Lane, Rockville, MD 20857, 301-796-3794.

SUPPLEMENTARY INFORMATION: In the **Federal Register** of April 25, 2008 (73 FR 22423), the agency announced that the proposed information collection had been submitted to OMB for review and clearance under 44 U.S.C. 3507. An agency may not conduct or sponsor, and a person is not required to respond to, a collection of information unless it displays a currently valid OMB control number. OMB has now approved the information collection and has assigned OMB control number 0910-0626. The approval expires on August 31, 2011. A copy of the supporting statement for this information collection is available on the Internet at <http://www.reginfo.gov/public/do/PRAMain>.

Dated: October 22, 2008.

Jeffrey Shuren,

Associate Commissioner for Policy and Planning.

[FR Doc. E8-25791 Filed 10-28-08; 8:45 am]

BILLING CODE 4160-01-S

DEPARTMENT OF HEALTH AND HUMAN SERVICES

Food and Drug Administration

[Docket No. FDA-2008-N-0543]

Agency Information Collection Activities; Proposed Collection; Comment Request; Waiver of In Vivo Demonstration of Bioequivalence of Animal Drugs in Soluble Powder Oral Dosage Form Products and Type A Medicated Articles

AGENCY: Food and Drug Administration, HHS.

ACTION: Notice.

SUMMARY: The Food and Drug Administration (FDA) is announcing an opportunity for public comment on the proposed collection of certain information by the agency. Under the Paperwork Reduction Act of 1995 (the PRA), Federal agencies are required to publish notice in the **Federal Register** concerning each proposed collection of information, including each proposed extension of an existing collection of information, and to allow 60 days for public comment in response to the notice. This notice solicits comments on the waiver requirement procedures that are recommended by the agency for in vivo demonstration of bioequivalence for generic soluble powder oral dosage form products and Type A medicated articles.

DATES: Submit written or electronic comments on the collection of information by December 29, 2008.

ADDRESSES: Submit electronic comments on the collection of information to <http://www.regulations.gov>. Submit written comments on the collection of information to the Division of Dockets Management (HFA-305), Food and Drug Administration, 5630 Fishers Lane, rm.1061, Rockville, MD 20852. All comments should be identified with the docket number found in brackets in the heading of this document.

FOR FURTHER INFORMATION CONTACT:

Denver Presley, Jr., Office of Information Management (HFA-710), Food and Drug