

academic, and private research and commercial organizations in tracking changes in trends of vital events.

Respondents for the National Vital Statistics Report form (CDC 64.146) are registration officials in each State and Territory, the District of Columbia, and New York City. In addition, 33 local (county) officials in New Mexico who record marriages occurring in each county of New Mexico will use this form. The data are routinely available in each reporting office as a by-product of ongoing activities. 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 Occurrence Report form (CDC 64.147) 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 National Vital Statistics Report form in that they represent complete and final 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 Occurrence 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. The data are routinely available in each reporting office as a by-product of ongoing activities.

There are no costs to the respondents other than their time. The total estimated annualized burden hours are 208.

ESTIMATED ANNUALIZED BURDEN TABLE

Respondents	No. of respondents	No. of responses per respondent	Average burden per response (in hours)
CDC64.146: State and Territory registration officials	58	12	12/60
CDC64.146: New Mexico County marriage registrars	33	12	6/60
CDC64.147: State/Territory/City registration officials	58	1	30/60

Dated: January 11, 2006.

Betsey S. Dunaway,

Acting Reports Clearance Officer, Centers for Disease Control and Prevention.

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DEPARTMENT OF HEALTH AND HUMAN SERVICES

Centers for Disease Control and Prevention

Clinical Laboratory Improvement Advisory Committee

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) announces the following committee meeting.

Name: Clinical Laboratory Improvement Advisory Committee (CLIAC).

Times and Dates: 8:30 a.m.-5 p.m., February 8, 2006; 8:30 a.m.-3 p.m., February 9, 2006.

Place: Doubletree Hotel (Atlanta/Buckhead), 3342 Peachtree Road NE., Atlanta, Georgia 30326, Telephone: (404) 231-1234.

Status: Open to the public, limited only by the space available. The meeting room accommodates approximately 100 people.

Purpose: This Committee is charged with providing scientific and technical advice and guidance to the Secretary, Department of Health and Human Services; the Assistant Secretary for Health; and the Director, CDC, regarding the need for, and the nature of, revisions to the standards under which

clinical laboratories are regulated; the impact on medical and laboratory practice of proposed revisions to the standards; and the modification of the standards to accommodate technological advances.

Matters To Be Discussed: The agenda will include updates from the Food and Drug Administration, the Centers for Medicare & Medicaid Services, and CDC; reports on national cytology proficiency testing status and Coordinating Council on the Clinical Laboratory Workforce activities addressing laboratory personnel shortages; and the role of the public health laboratory, including scope of services, customers, connectivity, and preparedness.

Agenda items are subject to change as priorities dictate.

Providing Oral or Written Comments: It is the policy of CLIAC to accept written public comments and provide a brief period for oral public comments whenever possible. *Oral Comments:* In general, each individual or group requesting to make an oral presentation will be limited to a total time of five minutes (unless otherwise indicated). Speakers must also submit their comments in writing for inclusion in the meeting's Summary Report. To assure adequate time is scheduled for public comments, individuals or groups planning to make an oral presentation should, when possible, notify the contact person below at least one week prior to the meeting date. *Written Comments:* For individuals or groups unable to attend the meeting, CLIAC accepts written comments until the date of the meeting (unless otherwise stated). However, the comments should be received at least one week prior to the meeting date so that the comments may be made available to the Committee for their consideration and public distribution. Written comments, one hard copy with original signature, should be

provided to the contact person below. Written comments will be included in the meeting(s) Summary Report.

Contact Person for Additional Information: Devery Howerton, Acting Chief, Laboratory Practice Standards Branch, Division Public Health Partnerships—Laboratory Systems, National Center for Health Marketing, Coordinating Center for Health Information and Service, CDC, 4770 Buford Highway NE., Mailstop G-23, Atlanta, Georgia 30341-3717; telephone (770) 488-8155; fax (770) 488-8279; or via e-mail at DHowerton@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 CDC and the Agency for Toxic Substances and Disease Registry.

Dated: January 10, 2006.

Alvin Hall,

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

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DEPARTMENT OF HEALTH AND HUMAN SERVICES

Centers for Disease Control and Prevention

Healthcare Infection Control Practices Advisory Committee

In accordance with section 10(a)(2) of the Federal Advisory Committee Act (Pub. L. 92-463), the Centers for Disease