CANCER STATISTICS REVIEW 1975-2011: INTRODUCTION

The annual SEER Cancer Statistics Review (**CSR**) contains incidence, mortality, prevalence, and survival statistics from 1975 through the most recent year for which data are available. This report is published by the Surveillance Research Program of the National Cancer Institute, which manages the Surveillance, Epidemiology, and End Results (SEER) Program. The scope and purpose of the **CSR** follow a report to the Senate Appropriations Committee (Breslow, 1988), which recommended that a broad profile of cancer be presented regularly to the American public.

The SEER program is an authoritative source of information on cancer incidence and survival in the United States. SEER collects and publishes these statistics from population-based registries covering 28% of the US population. The 18 SEER registries routinely collect data on patient demographics, primary tumor site, tumor morphology, extent of disease, first course of treatment, and active follow-up for vital status. Detailed information describing these fields can be found at http://seer.cancer.gov/resources/.

This report presents statistics on 29 primary sites and subsites, organized into site-specific chapters. Detailed statistics on cancer incidence, mortality, survival, and prevalence are reported by sex, race and ethnicity, age, stage at diagnosis, and geographic area. Information on tumor morphology is also presented. In addition, the *CSR* features a chapter on adolescent and young adult cancers and a chapter on childhood cancers. Information on some rare cancers can be found in the summary tables of section I. For a detailed list of primary sites, the summary tables provide incidence and death rates for the most recent 5-year period, trends from 1975 to the most recent year, median age at diagnosis, median age at death, and survival rates.

Delay-adjusted cancer incidence rates are a distinctive feature of the *CSR*. Delay-adjustment corrects the current case count to account for underreporting and corrections to the data. The final delay-adjusted rates are valuable in more precisely estimating trends.

New features added to the CSR include:

- Statistics for lung and bronchus cancer are shown by new histology groupings in Chapter 15 (Lewis DR, et al., 2014).
- Confidence intervals for state ranks in mortality were added (Zhang S, et al., 2014).

Changes in methodology to *CSR* include:

- New standard error calculation for delay-adjusted incidence rates was used. See http://surveillance.cancer.gov/reports/tech2014.01.pdf for more information.
- The default censoring age for survival calculations has changed from 199 to 99 years when using newly available expected survival tables. For most survival calculations there are no changes. Minimal changes may occur in survival for older age groups. See http://seer.cancer.gov/expsurvival/ for more information.

The *CSR* files are provided in both PDF and HTML formats. The HTML format is provided as an alternative and accessible version of the *SEER Cancer Statistics Review*. The current edition of the *CSR* is available on the web at <u>http://seer.cancer.gov/csr/</u>. Statistics from SEER may also be obtained via *FastStats* (<u>http://seer.cancer.gov/faststats/</u>) or *Cancer Query Systems* (<u>http://seer.cancer.gov/canques/</u>), which allow the user to access over 10,000,000 cancer statistics. The SEER Research Data file (<u>http://seer.cancer.gov/data/</u>) may be accessed by the public, either through *SEER*Stat* software or in an ASCII text format that can be analyzed with standard statistical software.

While most of the rates in this publication have been age-adjusted to the 2000 US standard population, some previous SEER publications have used the 1970 US standard million population. Therefore, rates given in this publication cannot be compared to rates given in those publications. This change conforms to a federal policy for reporting disease rates; it allows for the age-adjusted rate to more accurately reflect the current age distribution and burden of cancer.

INTERPRETATION OF CANCER STATISTICS

A number of factors may affect the interpretation of cancer incidence, mortality, and survival statistics provided in this report.

Survival rates for all cancers combined: The mix of cancers changes over time as the incidence of some cancers increases and the incidence of others decreases. The overall cancer survival rate can fluctuate even when the survival rates for site-specific cancers remain unchanged. (While it is possible to adjust the survival rate for all cancers combined on the basis of the relative frequencies of the component cancers, rates adjusted in this manner differ by only a small amount from unadjusted rates. In the future, such an adjustment may become more important if there are substantial changes in the incidence of various cancers.)

Early detection/screening: The improved earlier detection and diagnosis of cancers caused by new screening procedures may produce an *increase* in both incidence rates and survival rates. These increases can occur as a result of the introduction of a new procedure to screen subgroups of the population for a specific cancer; they need not be related to whether use of the screening test results in a decrease in mortality from that cancer. As the proportion of cancers detected at screening increases, presumably as a result of increased screening of the population, patient survival rates will *increase*, because they are based on survival time *after diagnosis*. The interval between the time a cancer is diagnosed by a screening is called **lead-time** (Zelen, 1976). (Screening for breast cancer has been demonstrated to result in increased survival over and above that resulting from lead-time alone and to reduce breast cancer mortality. The benefit of screening is being studied for some other cancers.)

If a new screening procedure consistently detects cancer in a *preinvasive* phase, it may result in a *decrease* in survival rates for *invasive* cancer. In this case, **length-biased sampling** (Zelen, 1976) may be operating. Length-biased sampling would result in the preferential detection—in a preinvasive phase—of those cancers that would have had a relatively good prognosis had they progressed to invasive disease; these potentially invasive cancers would be systematically eliminated. If this occurs, the mix of cancers that are not detected at screening and then progress to invasive behavior may become less prognostically favorable, resulting in a *decrease* in survival rates for patients with invasive cancers. (Length-biased sampling may at least partially explain survival trends for cervical cancer. Other cancers possibly affected include breast, colon, rectum, and prostate.)

Changes in diagnostic criteria: Early detection of cancer resulting from either screening or earlier response to symptoms may result in the increasing diagnosis of small tumors that are not yet life-threatening. This may have the effect of raising the incidence rates and survival estimates without changing the mortality rates. Breast, colon, prostate, cervix uteri, bladder, and skin (melanoma) are the cancer sites most likely to be affected.

Technological advances in diagnostic procedures: In this report, trends in survival by stage at diagnosis for specific cancers are not presented; trends in stage distributions are presented rarely. However, it is possible to compare survival by stage.

The assignment of a given stage to a particular cancer may change over time due to advances in diagnostic technology. Introduction of new technology can give rise to a phenomenon known as stage migration. Stage migration occurs when diagnostic procedures change over time, resulting in an increase in the probability that a given cancer will be diagnosed in a more advanced stage. For example, certain distant metastases that would have been undetectable a few years ago can now be diagnosed by a computer tomography (CT) scan or by magnetic resonance imaging (MRI). Therefore, some patients who would have been diagnosed previously as having cancer in a localized or regional stage are now diagnosed as having cancer in a *distant* stage. The likely result would be to remove the worst survivors, those with previously undetected distant metastases, from the localized and regional categories and put them into the distant category. As a result, the stage-at-diagnosis distribution for a cancer may become less favorable over time, but the survival for each stage may improve: The early stage will lose cases that will survive shorter than those remaining in that category, while the advanced stage will gain cases that will survive longer than those already in that category. However, overall survival would not change (Feinstein et al., 1985). Stage migration is an important concept to understand when examining temporal trends in survival by stage at diagnosis as well as temporal trends in stage distributions; it could affect the analysis of virtually all solid tumors.

Evolution of stage classifications: Every few years, the American Joint Committee on Cancer produces a new cancer-staging manual; the seventh edition is the most recent (Edge et al., 2010). The evolution of such classifications reflects the identification of new prognostic factors that may influence choice of treatment. Historically, the SEER Program has only collected data

on **extent of disease** (**EOD**), rather than stage. EOD is *more specific* than stage and usually determines stage, even when stage definitions change. Thus, SEER easily adapts to changes in stage definitions; moreover, trends in a newly redefined stage can usually be calculated. Recently the SEER Program has begun collecting **Collaborative Stage**. Collaborative Stage has the advantage of being a consolidated data collection system of three main staging systems (TNM, EOD, and Summary Stage) and allows combined pathological and clinical stage to be captured. New prognostic variables are introduced into staging for some cancers and so previously collected EOD data cannot determine new stage categories. There can be problems in assessing trends in stage of disease for these cancers. Only by reviewing the evolution of staging for a given cancer is it possible to determine what effects changes in stage definitions have had on stage-specific survival and on stage-at-diagnosis distributions. Stage migration (mentioned above) and EOD migration need also be taken into account. For some sites, the historic stage (*localized, regional,* or *distant*) is not shown, either because of inconsistencies in its definition over time or because stage is not appropriate (such as for leukemias, which are all considered to be distant at diagnosis).

Interpreting relative survival: The relative survival estimate is the ratio of observed survival to expected survival for a given patient cohort. Expected survival is based on mortality rates for the entire population, taking into account, as appropriate, the age, sex, race, and year of diagnosis of the patients. Assuming that the presence of cancer is the only factor that distinguishes the cancer patient cohort from the general population, relative survival estimates the probability that a patient will *not* die of the diagnosed cancer within the given time interval. This is the same as the probability that the patient will either survive the interval or die of a different cause.

A factor related to the risk of a cancer may also be related to the risk of dying from causes unrelated to the cancer. An example of such a factor is smoking. Smoking is a major risk factor for lung cancer; therefore, a cohort of lung cancer patients will contain a much higher proportion of smokers than the general population. However, smoking is also a risk factor for other diseases so smokers have a shorter life expectancy than nonsmokers. For this reason, expected survival estimates for lung cancer patients based on life tables for the general population will be unrealistically high; since relative survival = observed / expected, this will result in relative-survival estimates that are *lower* than they would be if the population consisted only of smokers. The problem cannot be easily corrected because separate life tables for smokers and nonsmokers are not available. Moreover, amount of smoking (usually measured in pack-years) is an important variable and cannot be easily quantified. In addition, expected survival may not be appropriate for patients with cancers of the cervix uteri or breast because the risk of these cancers has been associated with socioeconomic status (Baquet et al., 1991) which may be related to life expectancy. This should be considered when interpreting relative survival for these cancers.

Previous to the *CSR* for 1973–1996, the expected survival tables used were for 1970 and 1980; there were separate tables for whites, blacks, American Indians, Chinese, Japanese, Filipinos, white Hispanics, and Hawaiians. In updating the tables for 1990, several problems emerged.

The US life tables are based on age, race, and sex information from death certificates. The information on race on the death certificate may not be accurate (Rosenberg et al., 1999). One reason is that funeral directors may inaccurately report race on a death certificate. Also, reported age at death, especially for those older than 85, may not be accurate because birth certificates were not issued with as much regularity in the early 1900s as they are today. Although race misclassification and age-at-death misreporting exist across all races, they may be more problematic for races other than white or black because of those races' smaller population sizes. Therefore, life tables were generated for 1970, 1980, 1990, and 2000 only for white, black, and other; these life tables were used to produce the relative survival estimates in this review. There may be small variations among survival estimates calculated in this *CSR* and those in *CSR*s prior to 1973–1996.

Comparison with other databases: The SEER data are obtained from population-based cancer registries covering about 28 percent of the US population. It is sometimes of interest to compare cancer statistics for SEER areas with those from other registries both in the US and worldwide. In making such comparisons, one must carefully consider the factors mentioned above for both data sources. In addition, one should assess all of the following: (1) completeness of case ascertainment, (2) rules used to determine multiple primaries, (3) follow-up, (4) rules used in assigning and coding cause of death, and (5) the sources and procedures used in obtaining population estimates. Depending on the rates being compared, there could be other confounding factors which should be considered. The same standard or standard million population should be used for the age-adjustment of each group being compared; most statistics from outside the US are based on the 2000 world standard million population. Examples of other databases are US Cancer Statistics (<u>http://apps.nccd.cdc.gov/uscs</u>) and CINA+ Online (<u>http://www.cancer-rates.info/naaccr/</u>).

It is sometimes of interest to compare survival for cancer patients in SEER areas with data from clinical trials. *This must be done with great caution.* Survival data from clinical trials may have been obtained from a patient population that differs from that of SEER patients in prognostic factors for the given cancer; any survival comparisons would have to adjust for such differences. Also, it is necessary to verify that the methodology used in computing survival is the same for both data sources. Furthermore, patients on clinical trials may differ from SEER patients in characteristics that may be related to survival but are not recorded in either database. If this were true for a given cancer, it would not be possible to make valid comparisons of this type.

Errors in data collection: In the process of registering cancer patients, errors may be made in abstracting and coding the data, which include demographic information, cancer site, histology, extent of disease, treatment, and patient survival. Quality control studies are periodically carried out to detect and correct this type of error, but no attempt is made to incorporate this source of error into the variance estimates of cancer rates reported here.

Comparison of this report with previous reports: The cancer registries that participate in the SEER Program submit data on all cancers diagnosed in their coverage areas to the NCI each

year. Because of the dynamic nature of the registries' databases, *the reported number of new cancer cases in a particular race, sex, age, cancer category in a given calendar year may change from that which has been reported in a previous publication.* For a given diagnosis year, additional cancer cases that were previously overlooked may have been found and reported to the central registry. There may have been follow-back of cancers diagnosed by death certificate only; successful efforts to establish the dates of diagnosis for such patients will change the number of patients reported for a given diagnosis year. Code changes may occur when a patient dies; for example, information on race is generally available on the death certificate and may be used to update a previously unknown value. There may have been elimination of duplicate records for the same patient, often due to name changes or misspellings.

Thus, a recent report may have a different number of cases for a given diagnosis year than an earlier report, with resulting effects on incidence and possibly survival. Population estimates may also change from one report to another for some calendar years. This occurs because the NCI receives population estimates that are regularly revised and updated by the Bureau of the Census (**BOC**). Such changes may result in some differences between incidence and mortality rates for a given calendar period as published in different reports. See our website for the most current information about the population estimates (<u>http://seer.cancer.gov/popdata/</u>).

REFERENCES

Baquet CR, Horm JW, Gibbs T, Greenwald P. Socioeconomic factors and cancer incidence among blacks and whites. *J Natl Cancer Inst* 1991; 83:551-557.

Breslow L (Chairman, Extramural Committee to Assess Measures of Progress Against Cancer). Measurement of progress against cancer: Final report to the Senate Appropriations Committee. Bethesda: National Cancer Institute; 1988.

Edge SB, Byrd DR, Compton CC, Fritz AG, Greene FL, Trotti A, editors. *AJCC Cancer Staging Manual*, 7th ed. New York (NY): Springer; 2010.

Feinstein AR, Sosin DM, Wells CK. The Will Rogers phenomenon: Stage migration and new diagnostic techniques as a source of misleading statistics for survival of cancer. *New Engl J Med* 1985;312:1604-1608.

Lewis DR, Check DP, Caporaso NE, Travis WD, Devesa SS. U.S. Lung Cancer Trends by Histologic Type. *Cancer*, accepted March 13, 2014, in production.

Rosenberg HM, Maurer JD, Sorlie PD, Johnson NJ, MacDorman MF, Hoyert DL, Spitler JF, Scott C. Quality of death rates by race and Hispanic origin: A summary of current research. Hyattsville (MD): National Center for Health Statistics; Vital and Health Statistics, Series 2, No. 128, 1999.

Zelen M. Theory of early detection of breast cancer in the general population. In: Heuson J-C, Mattheiem WH, Rozencweig M, editors. *Breast Cancer: Trends in Research and Treatment*. New York (NY): Raven Press; 1976. p. 287-299.

Zhang S, Luo J, Zhu L, Stinchcomb D, Campbell D, Carter G, Gilkeson S, Feuer EJ. (2014), Confidence intervals for ranks of age-adjusted rates across states or counties. *Statistics in Medicine*. doi: 10.1002/sim.6071

TECHNICAL NOTES

There are four measures commonly used to assess the impact of a cancer in the general population and are reported in this review. The **incidence rate** is the number of new cases per year per 100,000 persons. The **death** (or **mortality**) **rate** is the number of deaths per year per 100,000 persons. The survival estimate is the proportion of patients alive at some point subsequent to the diagnosis of their cancer. The **prevalence count** is the number of people alive that have ever been diagnosed with a cancer. The Surveillance, Epidemiology, and End Results (**SEER**) Program (<u>http://seer.cancer.gov</u>) (based within the Surveillance Research Program (**SRP**) at the National Cancer Institute (**NCI**) collects incidence and survival data for all areas that participate in the Program. The National Center for Health Statistics (**NCHS**) provides mortality data for the entire United States (**US**). All incidence and mortality rates in this report are age-adjusted (see below) to the 2000 US standard population (see Appendix) unless otherwise specified. Age-adjustment minimizes the effect of a difference in age distributions when comparing rates.

THE SEER PROGRAM

The National Cancer Act of 1971 mandated the collection, analysis, and dissemination of data useful in the prevention, diagnosis, and treatment of cancer. This mandate led to the establishment of the SEER Program. The population-based cancer registries participating in NCI's SEER Program routinely collect data on all cancers occurring in residents of the participating areas. Trends in cancer incidence and patient survival in the US are derived from this database. See the SEER Research Data (<u>http://seer.cancer.gov/data/</u>) for more information.

The SEER Program is a sequel to two earlier NCI programs—the End Results Program and the Third National Cancer Survey. The initial SEER reporting areas were the States of **Connecticut**, **Iowa**, **New Mexico**, **Utah**, and **Hawaii**; the metropolitan areas of **Detroit**, Michigan, and **San Francisco-Oakland**, California; and the Commonwealth of Puerto Rico. Case ascertainment began with January 1, 1973, diagnoses.

In 1974-1975, the program was expanded to include the metropolitan area of New Orleans, Louisiana, the thirteen-county **Seattle-Puget Sound** area in the State of Washington, and the metropolitan area of **Atlanta**, Georgia. New Orleans participated in the program only through the 1977 data collection year. In 1978, ten predominantly African-American counties in **rural Georgia** were added. **American Indian residents of Arizona** were added in 1980. In 1983, four counties in New Jersey were added with coverage retrospective to 1979. New Jersey and Puerto Rico participated in the program until the end of the 1989 reporting year. The National Cancer Institute also began funding a cancer registry that, with technical assistance from SEER, collects information on cancer cases among **Alaska Native** populations residing in Alaska. In 1992, the SEER Program was expanded to increase coverage of minority populations, especially Hispanics, by adding **Los Angeles County** and four counties in the **San Jose-Monterey** area south of San Francisco. In 2001, the SEER Program expanded coverage to include **Kentucky**, **Greater California** (the counties of California that were not already covered by SEER), **New Jersey**, and **Louisiana**. In 2012, **Greater Georgia** (the parts of Georgia not included in Atlanta and Rural Georgia) was added to the SEER Program, with data retroactive to 2000.

The long-term incidence trends and survival data for this report are from five states (Connecticut, Hawaii, Iowa, New Mexico, and Utah) and four metropolitan areas (Detroit, Atlanta, San Francisco-Oakland, and Seattle-Puget Sound) (Fig. I-1); this set of registries is called the **SEER 9**. Additional tables show more recent incidence trends for the **SEER 13** areas (the 9 areas above plus Los Angeles, San Jose-Monterey, Alaska Native Registry, and rural Georgia) since 1992 and additional information on race and ethnicity. Other tables give statistics for the **SEER 18** areas; these are the SEER 13 plus Kentucky, Greater California, New Jersey, Louisiana, and Greater Georgia.

The participating regions were selected principally for their ability to operate and maintain a population-based cancer reporting system and for their epidemiologically significant population subgroups. With respect to selected demographic and epidemiologic factors, they are when combined a reasonably representative subset of the US population. Data from the 9, 13, or 18 SEER geographic areas are used in this report; the given groups contain, respectively, approximately 9, 14, or 28 percent of the US population. By the end of the 2011 diagnosis year, the database of the 18 SEER registries (plus Arizona Indians) contained information on over 7 million cases diagnosed since 1973. New cases added in the most recent data year numbered over 449,000.

The goals of the SEER Program are:

- 1) to assemble and report, on a periodic basis, estimates of cancer incidence, mortality, survival, and prevalence in the US;
- to monitor annual cancer incidence trends to identify unusual changes in specific forms of cancer occurring in population subgroups defined by geographic and demographic characteristics;
- 3) to provide continuing information on trends over time in the extent of disease at diagnosis, trends in therapy, and associated changes in patient survival; and
- 4) to promote studies designed to identify factors amenable to cancer control interventions, such as: (a) environmental, occupational, socioeconomic, dietary, and health-related exposures; (b) screening practices, early detection and treatment; and (c) determinants of the length and quality of patient survival.

DATA SOURCES

INCIDENCE AND SURVIVAL DATA

The SEER Program contracts with nonprofit, medically-oriented organizations having statutory responsibility for registering diagnoses of cancer among residents of their respective geographic coverage areas. Each SEER contractor:

1) maintains a cancer information reporting system;

- 2) abstracts records for *resident* cancer patients seen in every hospital both inside and outside the coverage area;
- 3) abstracts all death certificates of *residents* (dying both inside and outside the coverage area) on which cancer is listed as a cause of death;
- strives for complete ascertainment of cases by searching records of private laboratories, radiotherapy units, nursing homes, and other health services units that provide diagnostic service;
- 5) registers all in situ and malignant neoplasms (with the exceptions of certain histologies for cancer of the skin and—beginning in 1996—in situ neoplasms of the cervix uteri);
- records data on all newly diagnosed cancers, including selected patient demographics, primary site, morphology, diagnostic confirmation, extent of disease, and first course of cancer-directed therapy;
- 7) provides active follow-up on all living patients (except for those with in situ cancer of the cervix uteri);
- 8) maintains confidentiality of patient records;
- 9) at least annually submits electronically to NCI data on all reportable diagnoses of cancer made in residents of the coverage area.

For 1992 to 2000 diagnoses, the SEER program codes site and histology by the *International Classification of Diseases for Oncology*, second edition (**ICD-O-2**) (Percy et al., 1990). All cases before 1992 were machine-converted to ICD-O-2. Cases diagnosed 2001-2009 have been coded according to the third edition (**ICD-O-3**) (Fritz et al., 2000). Starting with patients diagnosed in 2007, the new multiple primary and histology coding rules may impact their incidence data for some cancer sites (e.g., female breast). However, the impact of the new rule on observed incidence is negligible for a majority of the cancer sites. To learn more about the multiple primary rules, visit: <u>http://seer.cancer.gov/tools/mphrules/</u>. Beginning with 2010 diagnoses, cases are coded based on ICD-O-3 updated for hematopoetic codes based on *WHO Classification of Tumours of Haematopoietic and Lymphoid Tissues* (2008). The primary site groupings used for incidence are found in the Appendix. Changes were made to the site recode for ICD-O-2 for comparability with cases coded to ICD-O-3. Follow-up rates are also in the Appendix.

Underreporting Adjustment for Veterans Affairs Cases: A CSR section on Department of Veterans Affairs (VA) underreporting (Howlader et al., 2009) was included in recent versions of the CSR. As of the current CSR this section was removed since available evidence indicates that VA underreporting is resolved as of diagnosis year 2010. The section of the CSR introduction about the reporting delay describes measures to address any backlog of VA cases reported after the initial reporting year.

Excluded cancers: Some cancers were excluded from most of the analyses. Myelodysplastic syndrome (MDS), for example, was reclassified in ICD-O-3 (effective diagnosis year 2001) from nonmalignant to malignant; other cancers so reclassified include endometrial stromal sarcoma (low grade), papillary ependymoma, papillary meningioma, polycythemia vera, chronic myeloproliferative disease (NOS), myelosclerosis with myeloid metaplasia,

essential thrombocythemia, refractory anemia, refractory anemia with sideroblasts, refractory anemia with excess blasts, and refractory anemia with excess blasts in transformation. In contrast, borderline tumors of the ovary were reclassified from malignant to nonmalignant at the same time. In addition, benign brain/CNS tumors were collected beginning for 2004 diagnoses. All of these cancers were excluded from most of the analyses, especially time trends. Pilocytic astrocytoma, although reclassified in ICD-O-3, was not excluded. Separate tables for MDS and benign brain/CNS are shown.

MORTALITY DATA

The SEER Program annually obtains from the National Center for Health Statistics (NCHS) a file containing information on all deaths occurring in the US by calendar year. Information on each death includes age at death, sex, geographic area of residence, and underlying and contributing causes of death. For this publication, only the underlying cause of death is used in the calculation of death rates. Cause of death for 1969-1978 was coded according to ICD-8; for 1979-1998, ICD-9 was used; beginning with deaths in 1999, ICD-10 was used. Mortality rates for the SEER geographic areas, for each state, and for the entire US are obtained from these data. A list of the mortality site groupings used in this publication is in the Appendix and reflects updates made in 2004.

POPULATION DATA

The population estimates used in the SEER*Stat software to calculate cancer incidence and mortality rates for this report are a modified version of the intercensal and Vintage 2011 annual time series of July 1 county population estimates by age, sex, race, and Hispanic origin that are produced by the Population Estimates Program of the US Census Bureau (http://www.census.gov/popest/) with support from the NCI through an interagency agreement. Descriptions of the methodologies employed by the Census Bureau for various sets of estimates may be found on the same website. Vintage 2011 population estimates were used; these estimates were developed from the actual 2010 census results.

County population estimates for 2000 and later years must be bridged from 31 race categories used in Census 2000 to the four race categories specified under the 1997 OMB standards in order to report long-term cancer trends. The bridging methodology was developed by the National Center for Health Statistics and is described in a report (Ingram et al., 2003) and on their website http://www.cdc.gov/nchs/nvss/bridged_race.htm

Modifications made by the NCI to the population estimates are documented in "Population Estimates Used in NCI's SEER*Stat Software" (<u>http://seer.cancer.gov/popdata/methods.html</u>) and the population data files are available for download (see "Download US Population Data" from <u>http://seer.cancer.gov/popdata/download.html</u>). Several of the modifications pertaining to the grouping of specific counties needed to assure the compatibility of all incidence, mortality and population datasets. Another modification affects only population estimates for the State of Hawaii. The Epidemiology Program of the Hawaii Cancer Research Center has developed its own set of population estimates, based on sample survey data collected by the Hawaii Department of Health. This effort grew out of a concern that the native Hawaiian population has been vastly undercounted in previous censuses. The "Hawaii adjustment" to the Census Bureau's estimates has the net result of reducing the estimated white population and increasing the estimated Asian and Pacific Islander population for the state. The estimates for the total population, black population, and American Indian and Alaska Native populations in Hawaii are not modified.

The cancer incidence and mortality rates for American Indians and Alaska Natives (AI/AN) are based on the geographic areas (counties) included in the Indian Health Service's Contract Health Service Delivery Area (CHSDA). This reflects a concern that previously reported AI/AN rates were underestimated due to racial/ethnic misclassification of American Indian cases in geographic areas outside of CHSDA. This change has the net effect of higher, and more accurate, incidence and mortality rates for this population. Beginning in 2013, CSR reporting diagnoses 1975-2010, CHSDA counties were updated with 9 new counties designated as CHSDA. Four of these are in SEER areas. This addition was made to better reflect AI/AN populations that had been living in these counties.

Usually the use of a population estimate for July 1 of a particular year reflects the average population of that area for the year. Both Hurricane Katrina and Hurricane Rita struck the Gulf Coast area of the United States in 2005. This had the effect of displacing large populations. Since there weren't any population estimates by age, race, sex, and county for time periods just after the hurricanes, it is very difficult to estimate the actual population at risk for certain areas along the Gulf Coast for 2005. For Louisiana, only the first six months of incidence data for 2005 coupled with ½ of the population estimate for July 1, 2005, were used to calculate cancer incidence. For death rate calculations, no adjustments were made to the total US population, but for the Gulf area, an adjustment for displaced populations was made for 2005 state rates. For more details, see http://seer.cancer.gov/popdata/methods.html.

2000 US STANDARD POPULATION

Starting with the November 2004 SEER submission of data (diagnoses through 2002), the SEER Program age-adjusts using the 2000 US standard population based on single years of age from the Census P25-1130 series estimates of the 2000 US population (Day, 1996). For the *CSR*, 19 age groupings were used for age-adjustment: <1, 1–4, 5–9, ..., 80–84, 85+.

STATISTICAL METHODS

ESTIMATED CANCER CASES AND DEATHS IN 2014

The American Cancer Society (**ACS**) projects the numbers of new cancer cases and cancer deaths in the US in 2014 (American Cancer Society, 2012). The ACS projects incidence in 2014 based on incidence rates for 1995-2009 from 49 states and the District of Columbia, representing about 98% of the US population. These high-quality incidence data were submitted to the North American Association of Central Cancer Registries (NAACCR) by 49 states (and District of Columbia) belonging to the SEER Program and/or the National Program of Cancer

Registries (NPCR). For additional details please refer to http://www.cancer.org/docroot/STT/STT_0.asp

LONG-TERM TRENDS, 1950-2011

Trends in cancer mortality from 1950 to 2011 are summarized by age both for all cancers combined and for lung cancer (Table 1-2). These cancer mortality trends are based on the mortality experience in the entire US. Summaries of long-term trends back to 1950 in cancer survival are also shown for whites.

Use caution when interpreting these statistics. Evaluating trends over a long period of time may hide recent changes in the trends.

YEARS OF LIFE LOST DUE TO PREMATURE DEATH FROM VARIOUS CAUSES

Death rates alone give an incomplete picture of the burden that deaths impose on the population. Another measure is the years of life lost due to premature death. This shows the extent to which life is cut short by a particular cause or disease.

This measure is estimated by linking life table data to each death of a person of a given age and sex. The life table permits a determination of the number of additional years an average person of that age, race, and sex would be expected to live. In this report, the age groups used in the calculation were 1-year intervals. These remaining years of life left are summed over all deaths due to a particular cause, yielding the estimate of the number of person-years of life lost (**PYLL**). The average years of life lost (**AYLL**) is obtained by dividing the PYLL by the number of deaths. Both of these measures can be calculated for any cause of death.

RELATIVE SURVIVAL

Relative survival (Ederer, 1961) was developed to provide an objective measure of the probability of survival of cancer in the absence of other causes of death. It is a measure that is not influenced by changes in mortality from other causes and, therefore, provides a useful measure for both tracking survival across time and comparisons between racial/ethnic groups or between registries. For most cancer registries, cause-of-death information obtained from death certificates is either unavailable or unreliable due to misclassification error. Therefore, instead of calculating the probability of surviving cancer in the usual (cause-specific) way, considering deaths from other causes as censoring events, relative survival compares the observed survival proportion of a group of cancer patients with the survival of a "similar" theoretical cancer-free group. Relative survival is formally defined as the ratio of the observed survival (all causes of death) of a cohort of cancer patients to the expected survival of a comparable set of cancer-free individuals. Since a cohort of cancer-free individuals is difficult to obtain, life tables representing survival of the general population are used instead. The underlying assumption is that the cancer deaths are a negligible proportion of all deaths. To learn more on this topic, visit: http://surveillance.cancer.gov/survival/measures.html.

Expected survival can be calculated using different methods which vary with respect to the definition of the matching group. The three most common methods are: Ederer I (Ederer, et al., 1961), Ederer II (Ederer and Heise, 1959) and Hakulinen (Hakulinen, 1982). In previous versions of SEER*Stat, relative survival has been calculated using Ederer I and Hakulinen methods, Ederer I being the default for calculations in the Cancer Statistics Review. In the Ederer I and Hakulinen methods, theoretical individuals are matched to each patient and are considered to be at risk for the entire follow-up. Hakulinen adjusts for potential follow-up times. Relative survival using expected rates derived via these two methods are very similar. However, recent research on relative survival has resuscitated the initial method to estimate expected rate: the Ederer II method. Although none of the three methods can be considered a gold standard, the Ederer II method has be shown to be in better alignment with the concept of net cancer survival. For that reason, as of 2011, we have switched to Ederer II as our default choice for calculating expected rate in SEER*Stat and the CSR. For more detail regarding this topic, read Cho et al., 2011 at: http://surveillance.cancer.gov/reports/. As of 2013, Survival time was calculated using pre-calculated months based on the exact day information. See http://seer.cancer.gov/survivaltime/. As of 2014, the default censoring age for survival calculations has changed from 199 to 99 year when using newly available expected survival tables. Minimal changes may occur in survival for older age groups. See http://seer.cancer.gov/expsurvival/ for more information.

CAUSE-SPECIFIC SURVIVAL

Cause-specific survival is a net-survival measure representing survival of a specified cause of death in the (theoretical) absence of other causes of death. Estimates are calculated by specifying the cause of death. Individuals who die of causes other than the specified cause are censored. This requires a cause-of-death variable that accurately captures all causes related to the specific cause. Cancer registries use algorithms to process causes of death from death certificates in order to identify a single, disease-specific, underlying cause of death. In some cases, attribution of a single cause of death may be difficult and misattribution may occur. For example, a death may be attributed to the site of metastasis instead of the primary site (Percy et al., 1981).

To capture deaths related to the specific cancer but not coded as such, the SEER causespecific death classification variable is defined by taking into account causes of deaths in conjunction with tumor sequence (i.e., only one tumor or the first of subsequent tumors), site of the original cancer diagnosis, and comorbidities (e.g., AIDS and/or site-related diseases). To learn more on this topic, please read the recent article published at the Journal of National Cancer Institute (Howlader et al., 2010) or visit: <u>http://seer.cancer.gov/causespecific/</u>.

CANCER PREVALENCE

Methods: In this report prevalence is calculated at 1/1/2011. Limited-duration prevalence is calculated using the counting method implemented in the SEER*Stat software. This method calculates the number or proportion of people alive at the prevalence date who had a diagnosis of the disease within the past *x* years (e.g., *x* = 5, 10, 20, or the full history of the registry).

Because SEER has available information for the various racial/ethnic groups for different numbers of years, different years and registries were used to estimate limited-duration prevalence. Prevalence estimates for all races combined, for whites, and for blacks use cases from 1975 through 2010 from the SEER 9 registries; prevalence estimates for Asian Pacific Islanders and Hispanics use cases diagnosed from 1990 through 2011 from the SEER 11 areas and rural Georgia.

The limited-duration prevalence method includes a correction for people lost to follow-up. For each individual lost to follow-up, a probability of being alive at the prevalence date is estimated from an appropriate survival function stratified by age at diagnosis (0–59, 60–69, 70+), sex, cancer site, year of diagnosis, and race, conditional on being alive at the time of loss to follow-up. Year of diagnosis is stratified into 5-year groups from the prevalence date, with the least recent interval being of varying length (4-8 years), depending on the length of years used to calculate prevalence. Race is stratified into white, black, other (American Indian/Alaska Native, Asian/Pacific Islander), and unknown/other-unspecified. When we use the SEER 11 registries, the same stratification as before is used, with American Indian/Alaska Native separated from Asian/Pacific Islander. Prevalence calculations for Hispanics use race stratified into: white, non-white, and unknown.

Different methods can be used to determine which tumors are to be included for people diagnosed with multiple tumors. Unless otherwise specified, prevalence calculations include only the *first malignant tumor per person*; that is, in situ cancers and second-or-later primary cancers were not included. Thus, if a woman had a melanoma prior to a breast cancer diagnosis, her melanoma would contribute to the prevalence of melanoma and to the prevalence of all sites, but the breast cancer would not contribute to the prevalence of breast cancer. Counting only one cancer per individual avoids some ambiguity in prevalence counts, and allows the counts for individual sites to sum to the all sites total. Prevalence using different selection criteria is compared in a table in the overview chapter. For more information on tumor selection criteria refer to http://surveillance.cancer.gov/prevalence/methods.html.

Complete prevalence is an estimate of the number of persons (or the proportion of population) alive on a specified date who had been diagnosed with the given cancer, no matter how long ago that diagnosis was. It was estimated for all races, whites, and blacks by applying the *completeness index method* (Capocaccia & De Angelis, 1997; Merrill et al., 2000; Mariotto et al., 2002) to limited-duration prevalence. The completeness index method is implemented in the COMPREV software, which can be found at http://surveillance.cancer.gov/comprev/. Validation of the completeness index for all races and for whites was made by using data from the Connecticut Tumor Registry (CTR) beginning with 1940. For blacks, SEER 9 data beginning with 1975 were used; identification of blacks is not possible in the CTR data prior to 1970. To validate the completeness index for blacks, we have compared the performance of the method to obtain 24-year prevalence from 10-year limited-duration prevalence. For all races combined and for whites, in cases where the validation indicated some lack of fit of the model, an approximation to the completeness index was derived from the CTR data. If there was a lack of

fit for blacks, no estimate of complete prevalence was reported. Complete prevalence for Asian/Pacific Islanders and Hispanics is not available at this time. Complete prevalence by age for all races combined was validated by comparing estimated 10-year complete prevalence with observed prevalence from the CTR data. Prevalence by age is reported for the sites that validated well.

The US cancer prevalence counts at 1/1/2011 were estimated by multiplying the SEER ageand race-specific prevalence proportions by the corresponding US population estimates based on the average of 2010 and 2011 population estimates from the US Census Bureau. US cancer prevalence counts for all races were estimated by summing the US estimated counts for whites/unknown, blacks, and other races. For Hispanics, the estimates for Hispanics of white or unknown race and for Hispanics of other races were summed.

Complete prevalence estimates of the number of individuals in the US diagnosed with cancer as children (ages 0-19), including those surviving for more than 36 years, is calculated using a statistical method that estimates the number of childhood survivors diagnosed before 1975 (Simonetti et al., 2008; Mariotto et al., 2009). Limited-duration prevalence proportions by age at prevalence are not shown for childhood cancers (age at diagnosis 0-19) since many of these estimates are not informative. For example, the number of people diagnosed with childhood cancers in the last 25 years and who are currently age 50-59 is zero by definition. For more details on available prevalence estimates, see http://surveillance.cancer.gov/prevalence/.

The overview chapter contains two prevalence tables. The first table reports US complete prevalence counts by age at prevalence and sex for some main cancer sites. The second table reports US prevalence counts for people diagnosed in the 5 years and 36 years prior to the prevalence date using different tumor inclusion criteria. Each site-specific chapter contains a prevalence table that reports limited-duration US prevalence counts by time since diagnosis for different racial/ethnic groups. US complete prevalence estimates are also reported when available. The second part of the site-specific tables displays the percent of the population in the SEER 11 areas diagnosed in the previous 19 years with the specific cancer by 10-year age groups for the different racial/ethnic groups.

PROBABILITY OF BEING DIAGNOSED WITH OR DYING FROM CANCER

Lifetime and interval risks of being diagnosed with cancer: The probability of being diagnosed with cancer is computed by applying cross-sectional age-specific 2008-2011 incidence rates from the SEER 17 areas and death rates from those same areas to a hypothetical cohort of 10,000,000 live births. This cohort is considered to be at risk for two mutually exclusive events: (1) developing the specified cancer, and (2) dying of other causes without developing the specified cancer. Using these two types of events, a standard **multiple decrement life table** (with 20 age groups from 0-4 to 90-94 and 95+) is derived. For each age interval, the number alive and free of the specified cancer at the beginning of the interval is decremented by the

risk of being diagnosed with the specified cancer is derived by summing all cancer cases from age 0-4 through age 95+ and dividing by 10,000,000. This calculation does not assume that an individual lives to any particular age; rather, it is the sum over all age intervals of the probability of living to the beginning of that interval without developing the given cancer times the probability of developing the cancer in that interval. The probability of developing cancer during any time period (e.g., between age 50 and age 60) is calculated by adding up all the cancers in the life table over the specified age range and dividing by the number of individuals alive and free of the specified cancer at the beginning of the period. The methodology is described in detail in (Fay et al., 2003) and (Fay, 2004). To improve the precision of the calculations, rates were calculated beyond the usual last open ended age interval (i.e. 85+) for the age groups 85-89, 90-94, and 95+.

Lifetime risk of dying from cancer: The lifetime risk of dying from a specified cancer is derived using a standard multiple decrement life table (Elandt-Johnson & Johnson, 1980). For each age, the risks of dying of the specified cancer and of all other causes are calculated, based on mortality data from the entire United States.

Detailed methodology and software: The estimates of developing and dying from cancer are implemented in DevCan (Probablity of DEVeloping or dying from CANcer software). More details on the software, various databases, and the methodology can be found at http://surveillance.cancer.gov/devcan/.

US CANCER DEATH RATES BY STATE

Each cancer-site-specific section presents the death rate for the given cancer for each state and the District of Columbia, specifying the five highest and the five lowest death rates by state for the most recent 5-year period for all persons, males only, and females only. The rates are per 100,000 persons; they are age-adjusted to the 2000 US standard population. (In some previous editions of the CSR, the 1970 US standard million population was used; *death rates standardized to the 2000 US standard million population cannot be compared to death rates standardized to the 1970 US standard million population.*)

The **percent difference (PD)** between a state rate and the rate for the total US is given by the formula:

PD = [(State Rate – Total US Rate)/Total US Rate] * 100

The **standard error** for each age-adjusted state death rate is calculated, based on the assumptions that (1) for each age-specific rate, the number of deaths is a Poisson random variable (Keyfitz, 1966) and (2) the variance of the age-adjusted rate is a linear combination of the variances of the age-specific rates (Snedecor & Cochran, 1980; pp. 188-9).

The standard error of the difference (SE_d) between a state rate and the total US rate is given

by the formula

$SE_d = Square Root of [SE_S^2 + SE_U^2 - 2 * Cov_{S,U}]$

where SE_S and SE_U are the standard errors of a state rate and of the total US rate, respectively, and Cov_{S,U} is the covariance between the two rates. The variance of each rate (i.e., the square of the standard error) and the covariance between the two rates are based on the Poisson assumption. The standard error does not represent the total error that may be present in the age-adjusted rate; it is merely the square root of the variance associated with the rates. In addition to this variance, there also exist potential biases and errors in the measurement of the rate that are difficult to assess accurately and probably impact differently on the error calculations for different states.

The difference between each age-adjusted state rate and the age-adjusted US rate is tested for statistical significance (see below) by calculating a Z (standard normal) statistic from the formula:

Z = (State rate – Total US rate) / SE_d

Although the rates being compared are not independent because each state is part of the US, the statistical test may not be substantially affected if the state represents a small proportion of the total US. There is also an adjustment for multiple comparisons; see below under *Statistical Significance*.

JOINPOINT REGRESSION ANALYSIS OF CANCER TRENDS

An advance in the presentation of cancer trends is the use of joinpoint models (Kim et al., 2000). In some past issues of the *Cancer Statistics Review*, certain time intervals (e.g., 1973–1996) were specified and the annual percent changes (APC) were computed over those intervals. The choices of where to start and where to end an interval were arbitrary and sometimes did not give an accurate picture of the trend for a given cancer site. For example, the rates might be increasing and decreasing in different parts of the same interval. For some sites, increases occurred in the earlier years, followed by declines in more recent years.

To achieve greater descriptive accuracy, a statistical algorithm finds the optimal number and location of places where a trend changes. The point (in time) when a trend changes is called a **joinpoint**. Trends may change in different ways at a joinpoint: from up to down, from down to up, from up to up at a different rate, or from down to down at a different rate. A **joinpoint regression model** describes the trends by a continuous, piecewise-exponential function. Adjacent segments are connected at a joinpoint. The segments are connected because we assume that rates generally change smoothly, rather than "jump" abruptly. In each segment, the rates are assumed to grow or decay exponentially ($y = e^{mx+b}$), i.e., to change by a constant percentage each year. Thus the "slope" *m* in each segment can be associated with a fixed annual percent change (**APC**) by $APC = 100(e^m - 1)$.

Joinpoint analysis first assumes no joinpoints are needed to describe the data accurately, i.e.,

the trend over the entire interval 1975-2011 does not change. Joinpoints are added in turn if they are statistically significant. Thus, in the final model, each joinpoint represents a significant change in trend. Smoother polynomial models may provide a good fit overall, but are less sensitive to what is occurring at the ends of the data.

In running the Joinpoint program, we set the program parameters as follows:

- (1) Joinpoints occur only at exact years; the joinpoint is not necessarily the same as the data point for that year;
- (2) The minimum time interval between consecutive joinpoints is three years;
- (3) The first joinpoint is not earlier than two years after the first year of data;
- (4) The last joinpoint is not later than two years before the last year of data;
- (5) The maximum number of joinpoints is five for 1975-2011 (SEER 9) data and three for 1992-2011 (SEER 13) data.

These restrictions provide some added stability to the resultant models. Different values for these parameters may yield a different joinpoint model. Since the test statistic to determine if additional joinpoints are necessary cannot be compared against any known standard distribution to determine significance (e.g., the normal, t, or f), a permutation test is used which simulates the distribution of the test statistic under the null hypothesis. Thus an element of randomness is introduced by the random number stream used. However, for greater consistency in the p-values obtained if one were to change the random seed for each run, we run the program for 4499 permutations.

A Windows-based program, Joinpoint, is freely available at

<u>http://surveillance.cancer.gov/joinpoint/</u>; it accepts data from the *SEER*Stat* program, as well as user-defined data. Further details on joinpoint regression may be found at the website. Starting with the 2011 edition of CSR, we have generated all our cancer trend statistics using a Linux-based *Joinpoint* program as opposed to the downloadable Windows-based program. As a result of using a different platform, in rare instances the results (e.g., # of joinpoints) may differ.

Average Annual Percent Change (AAPC) is a summary measure of a trend over a pre-specified fixed interval based on an underlying joinpoint model. It allows us to use a single number to describe the average trend over a period of multiple years. It can be estimated even if the joinpoint model indicates that there were changes in trends during those years, since it is estimated as a geometric weighted average of the joinpoint APCs, with the weights equal to the lengths of each segment over the pre-specified fixed interval. In this report, we have included AAPCs as an addendum to the underlying joinpoint trends, and as a summary measure to compare fixed interval trends by race/ethnicity. For more information on how the AAPC is calculated and the advantages of reporting an AAPC over APCs, see http://surveillance.cancer.gov/joinpoint/aapc.html.

REPORTING DELAY

Timely and accurate calculation of cancer incidence rates is hampered by reporting delay, the

time lapse before a diagnosed cancer case is reported to the NCI or the delay in receiving updated information for an existing case. Currently, the NCI allows a standard delay of 22 months between the end of the diagnosis year and the time the cancers are reported to the NCI in November, almost two years later. The data are released to the public in the spring of the following year. For example, cases diagnosed in 2011 were first reported to the NCI in November 2013 and released to the public in April 2014. However, in each subsequent release of the SEER data, *records from all prior diagnosis years* (e.g., diagnosis years 2010 and earlier in the 2013 submission to the NCI) *are updated* as either new cases are found or new information is received about previously submitted cases.

The submissions for the most recent diagnosis year are, in general, about two percent below the total number of cancers that will eventually be submitted for that year, although this varies by cancer site and other factors.

The idea behind modeling reporting delay is *to adjust the recent rates to anticipate future corrections (additions, changes, and deletions) to the data.* These adjusted rates and the associated delay model are valuable in more precisely determining current cancer trends, as well as in monitoring the timeliness of data collection—an important aspect of quality control (Clegg et al., 2002). Reporting delay models have been previously used in the reporting of AIDS cases (Brookmeyer & Damiano, 1989; Pagano et al., 1994; Harris, 1990).

In this report, we show SEER age-adjusted incidence rates and trends, along with their calculated delay adjustments for SEER 9 and SEER 13 areas. The adjusted rates, factors, and trends are available for all cancers combined (malignant only except for urinary bladder), for female breast in situ, for urinary bladder (in situ and malignant combined), and for 22 malignant cancer sites: melanoma (for all races combined and whites only), lung/bronchus, colon/rectum, prostate, female breast, liver and intrahepatic bile duct, pancreas, cervix uteri, corpus and uterus, ovary, testis, kidney and renal pelvis, brain and other nervous system, Hodgkin lymphoma, non-Hodgkin lymphoma, all leukemias, esophagus, larynx, myeloma, oral cavity and pharynx, thyroid, and stomach.

For more information on cancer incidence rates adjusted for reporting delay, see <u>http://surveillance.cancer.gov/delay/</u>. Estimates of observed incidence rates, delay-adjusted incidence rates, and delay-adjustments factors may be found in the Cancer Query Systems at <u>http://seer.cancer.gov/canques/</u>.

Adjustment for VA Case Backlog, Submission Year 2011

A policy change of the Department of Veterans Affairs (VA) regarding data sharing on VA cancer cases resulted in underreporting on VA hospital cases for submission years 2007-2011. Some special adjustments to case counts are necessary to fit the delay adjustment model. Beginning with the 2009 submission of SEER data, some SEER registries began accounting for the backlog of VA cases that would have been reported in 2006-2008. This upsurge in cases could cause perturbation in the delay model if fit in the usual manner.

As with the 2009 to 2011 submissions, to take account of the effect of the VA backlog in the 2012 submission on the delay adjustment model, the counts are adjusted by re-distributing VA cases to previous submission years according to the expected counts from the delay distribution conditional on the current submission. Specifically, for each of the diagnosis years 2004-2009, given the total cancer count in submission year 2012, the proportion of cumulative cancer count in each subsequent submission year is calculated based on the estimated parameters from previous year's reporting delay model. The VA cases in the 2012 submission are re-distributed to each of the prior submission years according to this proportion. The adjusted total cancer count in that submission year was then calculated by combining the non-VA cases and the re-distributed VA counts.

Delay-adjusted incidence rates and trends are reported for all cancers combined (malignant only except for urinary bladder), for female breast in situ, for urinary bladder (in situ and malignant combined), and for 22 malignant cancer sites: melanoma (for all races combined and whites only), lung/bronchus, colon/rectum, prostate, female breast, liver and intrahepatic bile duct, pancreas, cervix uteri, corpus and uterus, ovary, testis, kidney and renal pelvis, brain and other nervous system, Hodgkin lymphoma, non-Hodgkin lymphoma, all leukemias, esophagus, larynx, myeloma, oral cavity and pharynx, thyroid, and stomach.

STATISTICAL SIGNIFICANCE

Errors may be made in the estimation of a given statistic. In order to test whether two groups (such as the populations of a state and the entire US) have the same or different *actual* rates, the *observed* rates for the groups are compared. Statisticians consider that a difference in observed rates can be explained by one of two hypotheses: (H_0) The actual rates are really the same, but the observed rates are different because of some combination of error-causing factors, or (H_1) the actual rates of the groups are really different. H_0 is called the **null hypothesis** (because it says there is *no* real difference); H_1 is called the **alternate hypothesis**. Typically, H_0 is rejected only if there is strong evidence in favor of H_1 . (Thus, if the observed rates are equal, we cannot reject H_0 .)

Using statistical theory, one can determine the distribution of the rate difference under the assumption that H_0 is true. Then values of the rate difference that are very unlikely to occur if H_0 is true are identified. More specifically, a small positive number, called **alpha** (α), is chosen; usually, α is 0.05 or 0.01. (Alpha is called the **significance level** of the hypothesis test.) One can then identify limits for the difference in rates such that, if H_0 is true, the probability of the difference being outside of those limits is α . If the observed difference is *outside* of these limits, then the observed result is *very unlikely* to happen if H_0 is true, so H_0 is rejected.

Another way of looking at the same process is to calculate, assuming H_0 is true, the probability that the observed difference or any greater difference would occur; this number is called the *P*-**value** of the observed result. If the *P*-value of a comparison is less than α (that is, the observed difference is *very unlikely* to happen if the null hypothesis is true), H_0 will be rejected. If the *P*-value of a test is greater than the significance level α , H_0 will not be rejected. When a difference

in rates is sufficiently large to cause the null hypothesis to be rejected for a given value of α (usually 0.05), it is called a **statistically significant** difference.

When a null hypothesis is rejected, there remains a small chance that a wrong decision has been made. If many statistical comparisons are done, even with $\alpha = 0.01$, the chance of making at least one wrong decision becomes a concern. In testing the differences between the total US rate and the rate for each state (or for the District of Columbia) for a given cancer, 51 statistical comparisons of the type described above are performed. Based on one of Bonferroni's inequalities (if there are *n* events and *p_i* is the probability of success in event *i*, then *P*(at least 1 success) < *p*₁ + ... + *p_n*) (Snedecor & Cochran,1980; p. 115-117), the significance level α for each individual comparison was set equal to $0.01/51 \approx 0.0002$. Thus, only individual-state-to-total-US comparisons with an associated *P*-value less than 0.0002 are considered to be statistically significant. That is, a *very small* significance level α (0.0002) is used in order to minimize the total risk (0.01) of falsely deciding that some pair of equal rates are unequal.

Use caution in assessing statistically significant differences. Population size has an important role in any calculation of statistical significance. Some states may have estimated rates that are very close to the estimated total US rate, but because of their large population, the difference between their estimated rate and the estimated total US rate is found to be statistically significant. In this case, the true state rate and the true US rate are almost certainly different, because the observed difference, though small, is nearly impossible if the null hypothesis (equal rates) is true. A small difference in rates, however, may have no practical importance. On the other hand, some smaller states may have estimated rates that differ substantially from the estimated total US rate, but because of their relatively small population, the differences are found to be statistically nonsignificant. When this happens, if the true state rate and the true US rate were equal, the probability of obtaining a difference at least as large as what has been observed is greater than $\alpha \approx 0.0002$. Therefore, *because the evidence against it isn't strong enough, the null hypothesis (equal rates) is not rejected.*

If the percent difference (PD) between the two rates is small, there may be some question about the importance of the difference. It is difficult to specify a minimally significant absolute PD, below which the difference would always be unimportant, because the observed PD will depend on the populations of the areas involved. It may be of value to consider the size of the PD between a state rate and the US rate in assessing the importance of a statistically significant difference.

Comparing individual state rates with the US rate and assessing statistical significance is not an appropriate procedure for assessing geographic clustering of state rates. Identification of states which may represent regional clusters of high or low rates would require additional statistical and graphical analyses.

For a number of cancers, the District of Columbia has the highest death rates. *Use caution when comparing cancer rates for the District with those from the 50 states.* The District is an entirely urban area, whereas a state includes urban, suburban, and rural areas. Mortality rates

for many cancers are higher in urban areas. Also, the District has a higher percentage of blacks —51% of the total population in 2010 (US Census Bureau, 2013)—than any state. In addition, their higher mortality rates for several types of cancer elevate the overall rate for the District.

STANDARD ERRORS OF RATES

Survival rates: In the tables presenting survival estimates, the magnitude of the standard error is given as a measure of the reliability of a given rate: the greater the standard error, the more uncertainty associated with the estimated rate. In addition, if there were fewer than 25 diagnoses in the first interval of the life table constructed to calculate survival, or if all cases became lost to follow-up within an interval, a valid survival estimate could not be calculated, as is noted in the table footnotes.

The **standard error** (**SE**) of a relative survival estimate is obtained as follows (Ederer et al., 1961):

SE(CR_t) = CR_t * square root of $[q_1/(e_1-d_1) + q_2/(e_2-d_2) + ... + q_t/(e_t-d_t)]$

where CR_t is the *t*-year relative survival estimate, and for i = 1, ..., t, q_i is the probability of dying in year *i* after diagnosis, e_i is the effective number of patients at risk in year *i* after diagnosis, and d_i is the number of deaths in year *i* after diagnosis.

Incidence and mortality rates: The standard errors of age-adjusted incidence and mortality rates are often not specified. However, the reader can approximate the SE of a particular incidence or mortality rate by the SE of a crude incidence or mortality rate (Keyfitz, 1966), that is, the SE can be approximated by the rate divided by the square root of the number of cancer cases (or the number of deaths).

Appendix tables provide numbers of cancer diagnoses within SEER areas and numbers of deaths in the entire US, respectively, by race and sex for the most recent 5-year period. These can be used to obtain approximations of the standard errors for associated age-adjusted rates for the same time period using the above formula. To approximate the standard error of a rate for a single year, use the formula but replace the number of cancer cases or deaths with the number of cancer cases or deaths divided by 5.

DEFINITIONS

Several technical terms are used in presenting the data in this report. Their definitions are presented here to clarify them for the reader.

Incidence rate: The cancer incidence rate is the number of new cancers of a specific site/type occurring in a specified population during a year, usually expressed as the number of cancers

per 100,000 persons at risk. That is, Incidence rate = (New cancers / Population) * 100,000.

The *numerator* of the incidence rate is the number of new cancers; the *denominator* of the incidence rate is the size of the population. The number of new cancers may include multiple primary cancers occurring in one patient. The primary site reported is the site of origin and not the metastatic site. In general, the incidence rate would not include recurrences. *The population used depends on the rate to be calculated.* For cancer sites that occur in only one sex, the sexspecific population (e.g., females for cervical cancer) is used.

The incidence rate can be computed for a given type of cancer or for all cancers combined. Except for 5-year age-specific rates, all incidence rates in this report are *age-adjusted* (see below) to the 2000 US standard population (or, where appropriate, to the world standard million population). (In some previous editions of the *CSR*, the 1970 US standard million population was used; therefore, *incidence rates in this edition cannot be compared to rates published in those editions.*) Incidence rates are for *invasive cancer only*, unless otherwise specified. (Exceptions are the incidence rate for cancer of the urinary bladder (where both in situ and invasive cancers are counted) and breast cancer in situ, which is shown separately.)

Death rate: The cancer death (or mortality) rate is the number of deaths with cancer given as the underlying cause of death occurring in a specified population during a year, usually expressed as the number of deaths due to cancer per 100,000 persons. That is,

Death Rate = (Cancer Deaths / Population) * 100,000.

The *numerator* of the death rate is the number of deaths; the *denominator* of the death rate is the size of the population. As with the incidence rate, *the population used depends on the rate to be calculated.* The death rate can be computed for a given cancer site or for all cancers combined. Except for 5-year age-specific rates, all death rates in this report are *age-adjusted* (see below) to the 2000 US standard population (or, where appropriate, to the world standard million population). (In some previous editions of the *CSR*, the 1970 US standard million population was used; therefore, *death rates in this edition cannot be compared to rates published in those editions.*)

Age distribution: A table showing a partition of the entire lifespan into disjoint age intervals, along with the proportion of the population in each interval.

Median age: The age at which half of a population is younger and half is older.

Standard population: A standard population for a geographic area, such as the US or the world, is a table giving the proportions of the population falling into the age groups 0, 1-4, 5-9, ..., 80-84, and 85+. A standard million population for a geographic area is a table giving the number of persons in each age group 0, 1-4, ..., 85+ out of a theoretical cohort of 1,000,000 persons that is distributed by age in the same proportions as the standard population. Table A-7 shows the US 2000 standard population and the world standard million population. (Some

World Health Organization mortality publications use a different world standard million population.)

Age-adjusted rate: An age-adjusted incidence or mortality rate is a weighted average of the age-specific incidence or mortality rates, where the weights are the counts of persons in the corresponding age groups of a standard population. The potential confounding effect of age is reduced when comparing age-adjusted rates based on the same standard population. For this report, the 2000 US standard population (or, where appropriate, the world standard million population) is used in computing age-adjusted rates, unless otherwise noted.

Percent change: The percent change (**PC**) in a statistic over a given time interval is **Percent change = (Final value – Initial value) / Initial value * 100.**

A positive PC corresponds to an increasing trend, a negative PC to a decreasing trend.

Annual percent change: The annual percent change (APC) is calculated by first fitting a regression line to the natural logarithms of the rates (r) using calendar year (x) as a regressor variable. In this report the method of *weighted least squares* is used to calculate the regression equation. If ln(r) = mx + b is the resulting regression equation (with slope m), then APC = 100 * ($e^m - 1$). A positive APC corresponds to an increasing trend, a negative APC to a decreasing trend.

Because the methods used in their calculation are mathematically different, *the signs of the PC and the APC for a given statistic and time interval may differ*, as occurs in a few of the tables presented. That is, one of these statistics may show an increasing trend, the other a decreasing trend.

Testing the hypothesis that the actual mean annual percent change is 0 is equivalent to testing the hypothesis that the theoretical slope estimated by the slope *m* of the line representing the equation $\ln(\mathbf{r}) = \mathbf{mx} + \mathbf{b}$ is 0. The latter hypothesis is tested using the *t* distribution of m / SE_m with n - 2 degrees of freedom. The standard error of *m*, called SE_m , is obtained from the fit of the regression (Kleinbaum et al., 1988). (This calculation assumes that the rates increased or decreased at a constant rate over the entire calendar year interval; the validity of this assumption was not assessed.) In those few instances where at least one of the rates was 0, the linear regression was not calculated.

Average Annual Percent Change: The average annual percent change (AAPC) is a summary measure of a trend over a pre-specified fixed interval based on an underlying joinpoint model. It allows us to use a single number to describe the average trend over a period of multiple years. It can be estimated even if the joinpoint model indicates that there were changes in trends during those years, since it is estimated as a weighted average of the joinpoint APCs, with the weights equal to the lengths of each subinterval over the pre-specified fixed interval.

Life table: A table for a given population listing, for each sex and each age from 0 to 120, how many members die at that age and how many survive one more year.

Observed survival: The observed survival estimate represents the proportion of cancer patients surviving for a specified time interval after diagnosis. Note that some of those not surviving died of the given cancer and some died of other causes.

Relative survival: The relative survival estimate is calculated using a procedure (Ederer et al., 1961; Ederer and Heise, 1959) whereby the observed survival estimate is adjusted for expected mortality. The relative survival estimate approximates the likelihood that a patient will not die from causes associated specifically with the given cancer before some specified time after diagnosis. It is always larger than the observed survival estimate for the same group of patients.

Standard error: The standard error of a rate is a measure of the sampling variability of the rate.

Person-years of life lost: The person-years of life lost (**PYLL**) is calculated as follows: For each individual who dies of the cancer of interest, the number of years of expected additional life for an average person of that age, race, and sex is obtained from life tables for the US population (available from the NCHS). The PYLL in the general population associated with a particular cancer for a given year is simply the sum of this expectation over all those individuals who died of that cancer in that year.

Average years of life lost: The average years of life lost (**AYLL**) associated with a particular cancer for a given year is the PYLL associated with that cancer in the general population divided by the number of deaths from that cancer in the general population in that year.

Prevalence: Prevalence is defined as the number or percent of people alive on a certain date in a population who previously had a diagnosis of the disease. It includes new (incident) and preexisting cases and is a function of past incidence, past survival, and the size and age structure of the population. *Limited-duration prevalence* represents the proportion of people alive on a certain day who had a diagnosis of the disease within the past *x* years (e.g. x = 5, 10, or 20 years). *Complete prevalence* is an estimate of the number of persons (or the proportion of the population) alive on a specified date who had been diagnosed with the given disease, no matter how long ago that diagnosis was. For more details on cancer prevalence definitions and methods, refer to <u>http://surveillance.cancer.gov/prevalence/</u>.

Stage of disease at diagnosis: Extent-of-disease information determines stage of disease at diagnosis. The SEER summary stage presented has four levels. An invasive neoplasm confined entirely to the organ of origin is said to be localized. A neoplasm that has extended beyond the limits of the organ of origin, either directly into surrounding organs or tissues or into regional lymph nodes, is said to be **regional**. A neoplasm that has spread to parts of the body remote from the primary tumor, either by direct extension or by discontinuous metastasis, is said to be **distant**. When information is not sufficient to assign a stage, a neoplasm is said to be **unstaged**. In situ tumors (except those of the cervix uteri) are also collected by SEER but generally are not published in this series. For some cancers and diagnosis years, the extent of

disease information can also be converted to Stages 0-IV as defined by the American Joint Committee on Cancer (Greene et al, 2002; Edge et al., 2010).

SOFTWARE USED TO GENERATE THE SEER CANCER STATISTICS REVIEW

The SEER Cancer Statistics Review includes statistics generated by a variety of statistical software including:

- <u>SEER*Stat</u>, statistical software for the analysis of SEER and other cancer databases, was used to generate incidence, mortality, prevalence, and survival statistics presented in the CSR.
- Analysis generated by the <u>Joinpoint Regression Program</u> are presented to better describe trends that are not constant over time.
- The <u>DevCan</u> system generated the probability of developing cancer from twelve SEER areas and the probability of dying from cancer from the total United States.
- The <u>ComPrev</u> software was used to calculate complete prevalence estimates.

Additional statistics can be obtained via SEER's <u>Cancer Query Systems</u>. These data retrieval applications provide access to pre-calculated cancer statistics stored in online databases.

REFERENCES

American Cancer Society. *Cancer Facts & Figures 2012*. Atlanta: American Cancer Society; 2012.

Baquet CR, Horm JW, Gibbs T, Greenwald P. Socioeconomic factors and cancer incidence among blacks and whites. *J Natl Cancer Inst* 1991; 83:551-557.

Breslow L (Chairman, Extramural Committee to Assess Measures of Progress Against Cancer). Measurement of progress against cancer: Final report to the Senate Appropriations Committee. Bethesda: National Cancer Institute; 1988.

Brookmeyer R, Damiano A. Statistical methods for short-term projections of AIDS incidence. *Stat Med* 1989;8:23-34.

Byrne J, Kessler LG, Devesa SS. The prevalence of cancer among adults in the United States: 1987. *Cancer* 1992;68:2154-9.

Capocaccia R, De Angelis R. Estimating the completeness of prevalence based on cancer registry data. *Stat Med* 1997;16:425-40.

Cho H, Howlader N, Mariotto AB, Cronin KA. Estimating relative survival for cancer patients from the SEER Program using expected rates based on Ederer I versus Ederer II method. Surveillance Research Program, National Cancer Institute; 2011. Technical Report #2011-01.

Clegg LX, Feuer EJ, Midthune D, Fay MP, Hankey BF. Impact of reporting delay and reporting error on cancer incidence rates and trends. *J Natl Cancer Inst* 2002;94:1537-1545.

Clegg L, Gail M, Feuer EJ. Estimating the variance of disease prevalence estimates from population-based registries. *Biometrics* 2002;58(3):684-8.

Day JC. Population Projections of the United States by Age, Sex, Race, and Hispanic Origin: 1995 to 2050, US Census Bureau, Current Population Reports, P25-1130, US Government Printing Office, Washington, DC, 1996. Available from: <u>http://www.census.gov/prod/1/pop/p25-1130/p251130.pdf</u>

Ederer F, Axtell LM, Cutler SJ. The relative survival rate: A statistical methodology. *J Natl Cancer Inst Monogr* 1961;6:101-121.

Ederer F, Heise H. Instructions to IBM 650 Programmers in Processing Survival Computations, Technical, End Results Evaluation Section, National Cancer Institute, 1959.

Edge SB, Byrd DR, Compton CC, Fritz AG, Greene FL, Trotti A III. *AJCC Cancer Staging Manual*, 7th ed. New York (NY): Springer; 2010.

Elandt-Johnson RC, Johnson NL. *Survival Models and Data Analysis*. New York (NY): Wiley; 1980.

Fay MP. Estimating age conditional probability of developing disease from surveillance data. *Popul Health Metr.* 2004 Jul 27;2(1):6. Available from: http://www.pophealthmetrics.com/content/2/1/6

Fay MP, Pfeiffer R, Cronin KA, Le C, Feuer EJ. Age-conditional probabilities of developing cancer. *Stat Med.* 2003;22(11):1837-48.

Feinstein AR, Sosin DM, Wells CK. The Will Rogers phenomenon: Stage migration and new diagnostic techniques as a source of misleading statistics for survival of cancer. *New Engl J Med* 1985;312:1604-1608.

Feldman AR, Kessler L, Myers M, Naughton MD. The prevalence of cancer: Estimates based on the Connecticut Tumor Registry. *New Engl J Med* 1986; 315:1394-1397.

Feuer EJ, Wun L-M, Boring CC. Probability of developing cancer. In: Miller BA, Ries LAG, Hankey BF, Kosary CL, Edwards BK, editors. *Cancer Statistics Review: 1973-1989.* National Cancer Institute, NIH Pub. No. 92-2789, 1992. p. 1-8.

Feuer EJ, Wun L-M, Boring CC, Flanders WD, Timmel MJ, Tong T. The lifetime risk of developing breast cancer. *J Natl Cancer Inst* 1993;85:892-897.

Fritz A, Percy C, Jack A, Shanmugaratnam K, Sobin L, Parkin DM, Whelan S, editors. *International Classification of Diseases for Oncology*, 3rd ed. Geneva: World Health Organization; 2000.

Gail MH, Kessler L, Midthune D, Scoppa S. Two approaches for estimating disease prevalence from population-based registries of incidence and total mortality. *Biometrics* 1999;55:1137-44.

Greene FL, Page DL, Fleming ID, Fritz AG, Balch CM, Haller DG, Morrow M, editors. *AJCC Cancer Staging Manual*, 6th ed. New York (NY): Springer; 2002.

Hahn RA, Mulinare J, Teutsch SM. Inconsistencies in coding of race and ethnicity between birth and death in US infants. *JAMA* 1992;267:259-263.

Hakulinen T. Cancer survival corrected for heterogeneity in patient withdrawal. *Biometrics* 1982;38:933-942.

Harris JE. Reporting delays and the incidence of AIDS. J Am Stat Assoc 1990;85:915-924.

Howlader N, Ries LAG, Mariotto AB, Reichman ME, Ruhl J, Cronin KA. Improved estimates of cancer-specific survival rates from population-based data. *J Natl Cancer Inst* 2010;102:1-15.

Howlader N, Ries LAG, Stinchcomb DG, Edwards BK. The impact of underreported Veterans

Affairs data on national cancer statistics: analysis using population-based SEER registries. *J Natl Cancer Inst* 2009;101(7):533-536.

Ingram DD, Parker JD, Schenker N, Weed JA, Hamilton B, Arias E, Madans JH. United States Census 2000 population with bridged race categories. *Vital Health Stat 2*. 2003 Sep;(135):1-55.

Keyfitz N. Sampling variance of standardized mortality rates. Hum Biol 1966;38:309-317.

Kim H-J, Fay MP, Feuer EJ, Midthune DN. Permutation tests for joinpoint regression with applications to cancer rates. *Stat Med* 2000;19:335-351.

Kleinbaum DG, Kupper LL, Muller KE. *Applied Regression Analysis and Other Multivariable Methods*, 2nd ed. Boston: PWS-Kent, 1988.

Mariotto A, Gigli A, Capocaccia R, Clegg L, Scoppa S, Ries LA, Tesauro GS, Rowland JS, Feuer EJ. Complete and limited duration prevalence estimates. *SEER Cancer Statistics Review, 1973-1999.* 2002;19.

Merrill RM, Feuer EJ, Capocaccia R, Mariotto A. Cancer prevalence estimates based on tumor registry data in the SEER Program. *Int J Epidemiol* 2000;29:197-207.

Midthune DN, Fay MP, Clegg LX, Feuer EJ. Modeling reporting delays and reporting corrections in cancer registry data. *J Am Stat Assoc* 2005;100(469):61-70.

Pagano M, Tu XM, De Gruttola V, MaWhinney S. Regression analysis of censored and truncated data: estimating reporting-delay distributions and AIDS incidence from surveillance data. *Biometrics* 1994;50:1203-1214.

Percy C, Ries LAG, Van Holten VD. The accuracy of liver cancer as the underlying cause of death on death certificates. *Public Health Rep* 1990;105:361-368.

Percy C, Stanek E, Gloeckler L. Accuracy of cancer death certificates and its effect on cancer mortality statistics. *Am J Public Health* 1981;71: 3242-3250.

Percy C, Van Holten V, Muir C, editors. *International Classification of Diseases for Oncology*, 2nd ed. Geneva: World Health Organization;1990.

Ries LAG, Eisner MP, Kosary CL, Hankey BF, Miller BA, Clegg LX, Edwards BK (eds). *SEER Cancer Statistics Review, 1973-1997*. National Cancer Institute. NIH Pub. No. 00-2789. Bethesda, MD, 2000.

Robinson JG, West KK, Adlakha A. Coverage of the population in Census 2000: Results from demographic analysis. *Population Res Policy Rev* 2002;21:19-38.

Rosenberg HM, Maurer JD, Sorlie PD, Johnson NJ, MacDorman MF, Hoyert DL, Spitler JF, Scott C. Quality of death rates by race and Hispanic origin: A summary of current research. Hyattsville (MD): National Center for Health Statistics; Vital and Health Statistics, Series 2, No. 128, 1999.

Siegel R, Naishadham D, Jemal A. Cancer statistics, 2012. *CA Cancer J Clin* 2012 Jan-Feb;62(1):10-29.

Simonetti A, Gigli A, Capocaccia R, Mariotto A. Estimating complete prevalence of cancers diagnosed in childhood. *Stat Med* 2008 Mar 30;27(7):990-1007.

Snedecor GW, Cochran WG. *Statistical Methods*, 7th ed. Ames (IA): Iowa State University Press; 1980.

US Cancer Statistics Working Group. *United States Cancer Statistics: 1999-2002 Incidence and Mortality Web-based Report Version*. Atlanta: Department of Health and Human Services, Centers for Disease Control and Prevention, and National Cancer Institute; 2005. Available from: <u>http://www.cdc.gov/cancer/npcr/uscs/index.htm</u>

US Census Bureau. Current Population Reports; Series P-25 No. 985. Washington (DC): US Government Printing Office; 1986.

US Census Bureau: State and County QuickFacts. Data derived from Population Estimates, Census of Population and Housing, Small Area Income and Poverty Estimates, State and County Housing Unit Estimates, County Business Patterns, Nonemployer Statistics, Economic Census, Survey of Business Owners, Building Permits, Consolidated Federal Funds Report. Last Revised: Thursday, 04-Nov-2010 12:46:18 EDT.

Zelen M. Theory of early detection of breast cancer in the general population. In: Heuson J-C, Mattheiem WH, Rozencweig M, editors. *Breast Cancer: Trends in Research and Treatment*. New York (NY): Raven Press; 1976. p. 287-299.

Zou J, Huang L, Midthune D, Horner MJ, Krapcho M, Feuer EJ. Effect of reporting year on delay modeling. Statistical Research and Applications Branch, National Cancer Institute; 2009. Technical Report #2009-01. Available from: <u>http://surveillance.cancer.gov/reports/</u>.

Table 1.1

Estimated New Cancer Cases and Deaths for 2014 All Races, By Sex

	Esti	mated New (Cases	Estimated Deaths				
Primary Site	Total	Males	Females	Total	Males	Females		
All Sites	1,665,540	855,220	810,320	585,720	310,010	275,710		
Oral Cavity and Pharynx	42,440	30,220	12,220	8,390	5,730	2,660		
Tongue	13,590	9,720	3,870	2,150	1,450	700		
Mouth	11,920	7,150	4,770	2,070	1,130	940		
Pharynx	14,410	11,550	2,860	2,540	1,900	640		
Other Oral Cavity	2,520	1,800	720	1,630	1,250	380		
Digestive System	289,610	162,730	126,880	147,260	84,970	62,290		
Esophagus	18,170	14,660	3,510	15,450	12,450	3,000		
Stomach	22,220	13,730	8,490	10,990	6,720	4,270		
Small Intestine	9,160	4,880	4,280	1,210	640	570		
Colon ^a	96,830	48,450	48,380	50,310	26,270	24,040		
Rectum	40,000	23,380	16,620		,	,		
Anus, Anal Canal, and Anorectum	7,210	2,660	4,550	950	370	580		
Liver and Intrahepatic Bile Duct	33,190	24,600	8,590	23,000	15,870	7,130		
Gallbladder and Other Biliary	10,650	4,960	5,690	3,630	1,610	2,020		
Pancreas	46,420	23,530	22,890	39,590	20,170	19,420		
Other Digestive	5,760	1,880	3,880	2,130	870	1,260		
Respiratory System	242,550	130,000	112,550	163,660	90,280	73,380		
Larynx	12,630	10,000	2,630	3,610	2,870	740		
Lung and Bronchus	224,210	116,000	108,210	159,260	86,930	72,330		
Other Respiratory	5,710	4,000	1,710	790	480	310		
Bones and Joints	3,020	1,680	1,340	1,460	830	630		
Soft Tissue	12,020	6,550	5,470	4,740	2,550	2,190		
Skin (excl. basal & squamous)	81,220	46,630	34,590	12,980	8,840	4,140		
Melanoma of the Skin ^b	76,100	43,890	32,210	9,710	6,470	3,240		
Other non-epithelial skin	5,120	2,740	2,380	3,270	2,370	900		
Breast ^b	235,030	2,740	232,670	40,430	430	40,000		
Genital Organs	338,450	243,460	94,990	58,970	30,180	28,790		
Cervix (uterus)	12,360	243,400	12,360	4,020	30,100	4,020		
Endometrium (uterus)	52,630		52,630	8,590		8,590		
Ovary	21,980		21,980	14,270				
-						14,270		
Vulva	4,850		4,850	1,030		1,030		
Vagina and other genital organs, female	3,170		3,170	880	00.400	880		
Prostate	233,000	233,000		29,480	29,480			
Testis Penis and other genital	8,820 1,640	8,820 1,640		380 320	380 320			
organs, male	141 610	07 400	44 100	20.250	20 (10	0 740		
Urinary System	141,610	97,420	44,190	30,350	20,610	9,740		
Urinary Bladder	74,690	56,390	18,300	15,580	11,170	4,410		
Kidney and Renal Pelvis Ureter and other urinary	63,920 3,000	39,140 1,890	24,780 1,110	13,860 910	8,900 540	4,960 370		
organs	0 7 2 0	1 4 4 0	1 000	210	1 2 0	100		
Eye and Orbit Brain and Other Nervous	2,730 23,380	1,440	1,290	310	130	180		
System		12,820	10,560	14,320	8,090	6,230		
Endocrine System	65,630	16,600	49,030	2,820	1,300	1,520		
Thyroid	62,980	15,190	47,790	1,890	830	1,060		
Other Endocrine	2,650	1,410	1,240	930	470	460		
Lymphoma	79,990	43,340	36,650	20,170	11,140	9,030		
Hodgkin Lymphoma	9,190	5,070	4,120	1,180	670	510		
Non-Hodgkin Lymphoma	70,800	38,270	32,530	18,990	10,470	8,520		
Myeloma	24,050	13,500	10,550	11,090	6,110	4,980		
Leukemia	52,380	30,100	22,280	24,090	14,040	10,050		
Lymphocytic Leukemias	21,740	12,240	9,500	6,040	3,610	2,430		
Myeloid Leukemias	11,780	6,330	5,450	7,590	4,420	3,170		
Other leukemia	5,800	3,200	2,600	6,780	3,870	2,910		
All Other Sites ^c	31,430	16,370	15,060	44,680	24,780	19,900		

Cancer Facts & Figures - 2014, American Cancer Society (ACS), Atlanta, Georgia, 2014. Excludes basal and squamous cell skin and in situ carcinomas except urinary bladder.

Incidence projections are based on rates from the North American Association of Central Cancer Registries(NAACCR) from 1995-2010, representing about 89% of the US population. Estimated deaths are based on data from US Mortality Data, 1995-2010, National Center for Health Statistics, Centers for Disease Control and Prevention.

Estimated deaths for colon & rectum cancers are combined. Carcinoma *in situ* of the breast accounts for about 62,570 new cases annually, and melanoma *in situ* accounts for about 63,770 new cases annually. b

С

More deaths than cases suggests lack of specificity in recording underlying causes of death on death certificate.

а

Table 1.3

62-Year Trends in U.S. Cancer Death ${\tt Rates}^{\tt a}$

All Races, Males and Females

All Primary Cancer Sites Combined

Age Group	1950	1981	2011	Ann Percent 1950-1981		Total Percent Change 1950-2011
Ages 0-4	11.1	4.4	2.1	-3.2*	-2.6*	-81.1
Ages 5-14	6.7	4.1	2.1	-1.6*	-2.1*	-68.5
Ages 15-24	8.6	5.6	3.6	-1.2*	-1.5*	-57.5
Ages 25-34	20.4	13.3	8.5	-1.4*	-1.6*	-58.3
Ages 35-44	63.6	48.9	28.6	-0.8*	-1.8*	-55.1
Ages 45-54	174.2	172.9	106.6	0.1*	-1.7*	-38.8
Ages 55-64	391.3	430.7	292.7	0.4*	-1.5*	-25.2
Ages 65-74	710.0	823.9	660.0	0.5*	-0.8*	-7.0
Ages 75-84	1,167.2	1,229.5	1,167.2	0.2*	-0.2*	0.0
Ages 85+	1,450.7	1,580.4	1,682.7	0.3*	0.2*	16.0
All Ages	195.4	206.4	168.7	0.2*	-0.8*	-13.7

Lung and Bronchus Cancer^b

						Total
				Ann	Percent	
				Percent	Change	
Age Group	1950	1981	2011	1950-1981 1981-2011		1950-2011
Ages 0-4	-	-	-	-	-	-
Ages 5-14	-	-	-	-	-	-
Ages 15-24	0.2	0.1	0.1	-2.8*	-0.1	-55.3
Ages 25-34	0.8	0.6	0.3	-0.5	-2.4*	-62.8
Ages 35-44	4.6	9.5	3.1	2.5*	-2.8*	-33.2
Ages 45-54	20.2	52.5	24.9	3.2*	-2.7*	23.1
Ages 55-64	48.9	138.8	82.6	3.3*	-2.0*	69.0
Ages 65-74	59.4	238.1	218.8	4.1*	-0.4*	268.3
Ages 75-84	55.4	242.9	345.0	4.9*	1.0*	522.9
Ages 85+	42.3	178.9	325.5	5.1*	1.9*	669.6
All Ages	14.9	50.2	46.1	3.8*	-0.4*	208.4

Source: US Mortality Files, National Center for Health Statistics, Centers for Disease Control and Prevention. Rates are per 100,000 and age-adjusted to the 2000 US Std Population (18 age groups - Census P25-1130). a

b Due to coding changes throughout the years, Lung and Bronchus includes trachea and pleura.

*

The APC is significantly different from zero (p<.05). Statistic not shown. Rate based on less than 16 cases for the time interval. _

Trend based on less than 10 cases for at least one year within the time interval.

Table 1.4

Summary of Changes in Cancer Mortality, 1950-2011 and 5-Year Relative Survival (Percent), 1950-2010 Males and Females, By Primary Cancer Site

	Whites							
	Percent	ortality Change 2011 ^a	Surv	Relative ival cent) ^b				
Primary Site	Total	APC	1950-1954	2004-2010				
Oral cavity and pharynx	-51.9	-1.3*	46	67.4				
Esophagus	28.3	0.8*	4	21.0				
Stomach	-88.1	-3.4*	12	28.1				
Colon and rectum	-55.3	-1.3*	37	66.9				
Colon	-48.6	-1.0*	41	66.5				
Rectum	-70.3	-2.3*	40	68.0				
Liver and intrahepatic bile duct	53.0	0.8*	1	17.1				
Pancreas	26.1	0.1*	1	7.2				
Larynx	-39.2	-0.7*	52	64.0				
Lung and bronchus	207.5	1.4*	б	18.2				
Males	131.9	0.7*	5	15.8				
Females	543.9	2.9*	9	20.7				
Melanoma of the skin	173.7	1.3*	49	92.9				
Breast(females)	-35.8	-0.6*	60	91.8				
Cervix uteri	-81.9	-3.2*	59	71.0				
Corpus and uterus, NOS	-66.2	-1.6*	72	85.3				
Ovary	-12.1	-0.3*	30	44.3				
Prostate	-33.9	-0.4*	43	99.8				
Testis	-70.5	-2.8*	57	97.0				
Urinary bladder	-28.7	-0.7*	53	79.8				
Kidney and renal pelvis	33.7	0.5*	34	73.8				
Brain and nervous system	50.5	0.5*	21	33.4				
Thyroid	-40.4	-1.0*	80	98.3				
Hodgkin lymphoma	-80.9	-3.3*	30	88.3				
Non-Hodgkin lymphoma	82.6	0.9*	33	72.5				
Myeloma	217.3	1.2*	6	46.8				
Leukemia	-0.8	-0.3*	10	61.0				
Childhood (Ages 0-14)	-74.3	-2.7*	20	84.4				
All Sites	-14.1	-0.1*	35	69.4				

The APC is the Annual Percent Change over the time interval. Rates used in the calculation of the APC are age-adjusted to the 2000 U.S. standard population (18 age groups - Census P25-1130). U.S. Mortality Files, National Center for Health Statistics, Centers for Disease Control and Prevention. Due to coding changes throughout the years: Colon excludes other digestive tract; Rectum includes anal canal; Liver & intrahepatic bile duct includes gallbladder & biliary tract, NOS; Lung & bronchus includes trachea & pleura; Ovary includes fallopian tube; Urinary bladder includes other urinary organs; Kidney & Renal pelvis includes ureter; NHL and myeloma each include a small number of leukemias; NHL includes a small number of ill-defined sites. b Survival estimates for 1950-54 are from NCI Survival Report 5 with the exception of All Sites, Oral cavity & pharynx, Colon & rectum, Non-Hodgkin lymphoma and Childhood cancers which come from historical Connecticut data. Survival estimates for 2004-2010 are from the SEER 9 areas (San Francisco, Connecticut, Detroit, Hawaii, Iowa, New Mexico, Seattle, Utah, and Atlanta). Rates are based on follow-up of patients into 2011.

The APC is significantly different from zero (p<.05).

Table 1.5 Age-Adjusted SEER Incidence and U.S. Death Rates and 5-Year Relative Survival (Percent) By Primary Cancer Site, Sex and Time Period

	Incidence ^a (2007-2011)		US Mortality ^b (2007-2011)			Survival [°] (%) (2004-2010)			
Site	Total	Males	Females	Total	Males	Females	Total	Males	Females
All Sites	460.4	529.4	411.3	173.8	211.6	147.4	66.1	66.5	65.6
Oral Cavity & Pharynx:	11.0	16.5	6.2	2.5	3.8	1.4	62.7	61.9	64.6
Lip	0.7	1.1	0.3	0.0	0.0	0.0	89.5	90.5	86.5
Tongue	3.2	4.8	1.8	0.6	0.9	0.4	62.7	62.9	62.2
Salivary gland	1.3	1.7	1.0	0.2	0.4	0.1	72.4	65.4	81.5
Floor of mouth	0.6	0.8	0.3	0.0	0.0	0.0	51.4	49.1	57.4
Gum & other oral cavity	1.5	1.8	1.3	0.4	0.4	0.3	59.7	57.9	61.8
Nasopharynx	0.7	1.0	0.4	0.2	0.3	0.1	59.2	57.9	62.4
Tonsil	1.8	3.1	0.6	0.2	0.4	0.1	70.8	71.6	67.0
Oropharynx	0.4	0.7	0.2	0.2	0.4	0.1	41.7	43.1	36.5
Hypopharynx	0.6	1.1	0.2	0.1	0.2	0.0	31.9	31.5	33.6
Other oral cavity & pharynx	0.2	0.3	0.1	0.5	0.8	0.2	35.4	37.3	30.6
Digestive System:	84.3	103.2	68.9	42.5	54.6	32.8	44.3	42.6	46.4
Esophagus	4.4	7.7	1.8	4.2	7.5	1.6	17.5	17.6	17.2
Stomach	7.5	10.3	5.3	3.5	4.7	2.5	28.3	26.6	30.8
Small intestine	2.1	2.5	1.8	0.4	0.4	0.3	65.2	65.1	65.3
Colon & Rectum:	43.7	50.6	38.2	15.9	19.1	13.5	64.7	65.0	64.5
Colon	31.1	34.8	28.2	-	-	-	64.0	64.5	63.6
Rectum	12.6	15.7	10.0	-	-	-	66.5	66.0	67.2
Anus, anal canal & anorectum	1.8	1.5	2.0	0.2	0.2	0.3	65.5	60.8	68.6
Liver & intrahepatic	7.9	12.4	4.1	5.8	8.5	3.4	16.6	16.4	17.2
bile duct									
Gallbladder	1.2	0.8	1.4	0.6	0.5	0.7	17.4	15.5	18.2
Other biliary	1.9	2.3	1.5	0.4	0.5	0.4	16.3	17.6	14.8
Pancreas	12.3	14.0	10.9	10.9	12.5	9.6	6.7	6.3	7.0
Retroperitoneum	0.4	0.4	0.4	0.1	0.1	0.1	54.4	55.1	53.6
Peritoneum, omentum &	0.6	0.1	1.1	0.3	0.1	0.4	31.6	44.5	30.7
mesentery									
Other digestive system	0.6	0.7	0.5	0.3	0.4	0.2	11.3	10.4	12.1
Respiratory System:	64.3	79.3	53.0	49.8	64.0	39.1	19.8	18.9	20.9
Nose, nasal cavity &	0.7	0.9	0.5	0.1	0.2	0.1	54.7	55.7	53.4
middle ear									
Larynx	3.3	5.9	1.2	1.1	2.0	0.4	60.0	61.0	56.1
Lung & bronchus	60.1	72.2	51.1	48.4	61.6	38.5	16.8	14.4	19.6
Pleura ^d	0.0	0.0	0.0	0.1	0.1	0.0	19.3	16.4	23.5
Trachea & other	0.2	0.3	0.1	0.1	0.1	0.0	48.8	50.2	45.0
respiratory organs									
Bones & joints	0.9	1.1	0.8	0.4	0.5	0.3	66.6	64.9	68.7
Soft tissue (including heart)	3.3	4.0	2.8	1.3	1.5	1.1	65.3	64.5	66.4
Skin (excl. basal & squamous):	23.3	30.5	18.1	3.6	5.6	2.1	90.7	88.7	93.2
Melanoma of the skin	23.3	27.7	16.7	2.7	4.1	1.7	91.3	89.3	93.7
Other non-epithelial skin	21.3	2.8	1.4	0.9	1.5	0.4	84.3	81.7	87.7
conce non openicitat bain	2.0	2.0	±•±	0.9	1.5	0.1	01.0	··· /	0,.,
Breast	67.1	1.2	124.6	12.4	0.3	22.2	89.2	83.2	89.2
Breast (<i>in situ</i>)	16.8	0.2	31.7	-	-	-	100.0	100.0	100.0

Note: Incidence and death rates are per 100,000 and are age-adjusted to the 2000 US Std Population (19 age groups - Census P25-1130). а SEER 18 areas (San Francisco, Connecticut, Detroit, Hawaii, Iowa, New Mexico, Seattle, Utah, Atlanta, San Jose-Monterey, Los Angeles, Alaska Native Registry, Rural Georgia, California excluding SF/SJM/LA, Kentucky, Louisiana, New Jersey and

Georgia excluding ATL/RG).

b US Mortality Files, National Center for Health Statistics, Centers for Disease Control and Prevention.

С

SEER 18 areas. Based on follow-up of patients into 2011. Mesotheliomas of the Pleura are included in the separate group Mesothelioma for incidence but are included in the Pleura grouping for mortality. d

Statistic could not be calculated due to less than 16 cases in the time interval.

Age-Adjusted SEER Incidence and U.S. Death Rates and 5-Year Relative Survival (Percent) By Primary Cancer Site, Sex and Time Period

	Incidence ^a (2007-2011)			Mortalit 007-2011		Survival ^c (%) (2004-2010)			
Site	Total	Males	Females	Total	Males	Females	Total	Males	Females
Female Genital System: Cervix uteri Corpus uteri Uterus, NOS Ovary ^d Vagina Vulva Other female genital system	26.1 4.0 12.8 0.4 6.6 0.4 1.3 0.5	- - - - - - -	48.8 7.8 23.9 0.7 12.3 0.7 2.4 1.0	8.6 1.2 1.1 1.4 4.4 0.1 0.3 0.1	- - - - - - -	15.5 2.3 1.9 2.4 7.9 0.2 0.5 0.2	68.4 67.9 82.8 26.4 44.6 51.8 70.5 59.1	- - - - - -	68.4 67.9 82.8 26.4 44.6 51.8 70.5 59.1
Male Genital System: Prostate Testis Penis Other male genital system	70.3 67.0 2.8 0.4 0.1	154.5 147.8 5.6 0.9 0.3	- - - -	9.0 8.8 0.1 0.1 0.0	22.8 22.3 0.2 0.2 0.0	- - - -	98.6 98.9 95.3 67.9 86.9	98.6 98.9 95.3 67.9 86.9	- - - -
Urinary System: Urinary bladder Kidney & renal pelvis Ureter Other urinary system	36.9 20.5 15.5 0.6 0.3	58.8 36.2 21.2 0.8 0.5	20.1 8.8 10.7 0.4 0.2	8.6 4.4 4.0 0.1 0.1	13.8 7.7 5.8 0.1 0.2	4.9 2.2 2.6 0.1 0.1	74.7 77.4 72.4 47.6 50.7	75.9 78.9 72.1 47.6 54.6	72.2 72.7 72.9 47.9 44.3
Eye & Orbit	0.8	0.9	0.7	0.1	0.1	0.1	81.8	82.0	81.7
Brain & Nervous System: ^e Brain Cranial nerves & other nervous system	6.4 6.0 0.4	7.6 7.3 0.4	5.4 5.0 0.4	4.3 - -	5.2 - -	3.5 - -	33.4 30.3 77.2	32.5 29.8 75.6	34.5 30.9 78.7
Endocrine System: Thyroid Other endocrine & thymus	13.6 12.9 0.8	7.3 6.4 0.9	19.8 19.1 0.7	0.8 0.5 0.3	0.8 0.5 0.3	0.8 0.5 0.3	95.8 97.8 64.1	91.4 95.6 64.0	97.2 98.5 64.3
Lymphoma: Hodgkin lymphoma Non-Hodgkin lymphoma	22.4 2.7 19.7	27.0 3.1 23.9	18.8 2.4 16.3	6.7 0.4 6.3	8.5 0.5 8.1	5.3 0.3 5.0	71.6 85.3 69.3	70.4 84.4 68.0	73.0 86.4 70.9
Myeloma	6.1	7.7	4.9	3.4	4.3	2.7	44.9	46.0	43.5
Leukemia: Lymphocytic: Acute lymphocytic Chronic lymphocytic Other lymphocytic Myeloid & Monocytic: Acute myeloid Chronic myeloid Acute monocytic Other myeloid & monocytic Other leukemia: Other acute leukemia Aleukemic, subleukemic & NOS	$13.0 \\ 6.5 \\ 1.7 \\ 4.4 \\ 0.4 \\ 5.9 \\ 3.8 \\ 1.7 \\ 0.3 \\ 0.2 \\ 0.6 \\ 0.2 \\ 0.4$	16.7 8.6 1.9 6.0 0.6 7.3 4.6 2.2 0.3 0.2 0.8 0.3 0.5	$10.2 \\ 4.9 \\ 1.5 \\ 3.1 \\ 0.2 \\ 4.8 \\ 3.2 \\ 1.2 \\ 0.2 \\ 0.1 \\ 0.5 \\ 0.2 \\ 0.3$	$\begin{array}{c} 7.0\\ 2.0\\ 0.5\\ 1.4\\ 0.1\\ 3.4\\ 2.8\\ 0.3\\ 0.0\\ 0.2\\ 1.7\\ 0.6\\ 1.0 \end{array}$	$\begin{array}{c} 9.4\\ 2.8\\ 0.5\\ 2.0\\ 0.2\\ 4.5\\ 3.7\\ 0.4\\ 0.0\\ 0.3\\ 2.2\\ 0.8\\ 1.4\end{array}$	5.3 1.4 0.9 0.1 2.6 2.2 0.0 0.1 1.3 0.5 0.8	57.2 76.4 66.7 80.2 81.1 36.3 24.9 62.0 22.5 33.4 30.8 18.4 39.0	58.0 76.5 67.1 79.5 85.4 36.0 23.9 61.2 21.5 32.7 30.9 18.7 40.0	56.2 76.1 66.3 81.1 69.1 36.7 26.1 63.0 23.6 34.2 30.5 18.1 38.0
Kaposi Sarcoma ^f Mesothelioma ^f	0.5 1.0	1.0 1.9	0.1 0.4	-	-	-	72.1 8.3	71.5 6.6	76.5 13.5
Ill-defined & unspecified	8.9	10.3	7.8	12.9	16.3	10.3	17.2	21.3	13.2

Note: Incidence and death rates are per 100,000 and are age-adjusted to the 2000 US Std Population (19 age groups - Census P25-1130). a SEER 18 areas (San Francisco, Connecticut, Detroit, Hawaii, Iowa, New Mexico, Seattle, Utah, Atlanta, San Jose-Monterey, Los Angeles, Alaska Native Registry, Rural Georgia,

California excluding SF/SJM/LA, Kentucky, Louisiana, New Jersey and Georgia excluding ATL/RG).

b US Mortality Files, National Center for Health Statistics, Centers for Disease Control and Prevention. С

- SEER 18 areas. Based on follow-up of patients into 2011. d
- е
- Ovary excludes borderline cases or histologies 8442, 8451, 8462, 8472, and 8473. Due to coding changes, Brain & Nervous System mortality are no longer shown separately. Rate not shown for mortality. Category did not exist in mortality coding until 1999. Statistic could not be calculated due to less than 16 cases in the time interval. f

Table 1.6 Age-Adjusted SEER Incidence and U.S. Death Rates and 5-Year Relative Survival (Percent) By Primary Cancer Site, Sex and Time Period

Whites

			Whites							
	I	ncidence	ea	US	Mortali	t.v ^b	Survival ^c (%)			
		2007-201			2007-201			(2004-20	. ,	
Site	Total	Males	Females	Total	Males	Females	Total	Males	Females	
All Sites	468.9	532.1	424.4	173.3	209.8	147.5	66.7	66.9	66.4	
Oral Cavity & Pharynx:	11.4	17.0	6.4	2.4	3.7	1.4	64.3	64.0	65.1	
Lip	0.8	1.3	0.4	0.0	0.0	0.0	89.3	90.4	86.1	
Tongue	3.5	5.2	1.9	0.6	0.9	0.4	64.7	65.2	63.4	
Salivary gland	1.3	1.8	1.0	0.2	0.4	0.1	71.3	64.0	81.1	
Floor of mouth	0.6	0.9	0.4	0.0	0.0	0.0	52.5	50.5	57.6	
Gum & other oral cavity	1.5	1.8	1.3	0.4	0.4	0.3	59.8	58.1	61.9	
Nasopharynx	0.4	0.6	0.3	0.2	0.2	0.1	55.3	55.5	54.7	
Tonsil	2.0	3.4	0.7	0.2	0.4	0.1	72.8	73.5	69.1	
Oropharynx	0.4	0.6	0.2	0.2	0.3	0.1	45.4	47.5	37.8	
Hypopharynx	0.6	1.1	0.3	0.1	0.1	0.0	34.7	34.8	34.1	
Other oral cavity & pharynx	0.2	0.4	0.1	0.4	0.8	0.2	38.1	40.7	32.0	
Digestive System:	81.7	99.9	66.6	41.2	52.8	31.6	45.0	43.4	46.9	
Esophagus	4.6	8.0	1.7	4.3	7.8	1.5	18.3	18.4	17.8	
Stomach	6.6	9.2	4.5	3.0	4.1	2.1	26.9	25.3	29.5	
Small intestine	2.1	2.5	1.8	0.3	0.4	0.3	66.0	66.6	65.2	
Colon & Rectum:	42.9	49.6	37.3	15.5	18.5	13.0	65.4	65.7	65.0	
Colon	30.6	34.3	27.5	-	-	-	65.0	65.5	64.4	
Rectum	12.3	15.4	9.7	-	-	-	66.4	66.2	66.7	
Anus, anal canal & anorectum	1.9	1.5	2.2	0.2	0.2	0.3	67.2	63.0	69.8	
Liver & intrahepatic bile duct	7.0	10.8	3.6	5.3	7.8	3.2	16.1	15.9	16.5	
Gallbladder	1.1	0.8	1.4	0.6	0.4	0.7	17.5	15.7	18.2	
Other biliary	1.1	2.3	1.4	0.0	0.4	0.4	16.2	17.9	14.3	
-	12.2	14.0	10.7	10.4	12.5	0.4 9.4	6.6	6.3	6.9	
Pancreas Retroperitoneum	0.4	0.4	0.4	0.1	0.1	0.1	54.9	54.6	54.9	
Peritoneum, omentum &	0.4	0.4	1.2	0.1	0.1	0.1	31.3	44.4	30.4	
mesentery	0.7	0.1	1.2	0.3	0.1	0.4	51.5	11.1	30.4	
Other digestive system	0.5	0.6	0.5	0.3	0.4	0.2	11.4	11.3	11.3	
Respiratory System:	66.0	79.6	55.7	50.4	63.7	40.4	20.1	19.2	21.1	
Nose, nasal cavity & middle ear	0.7	0.9	0.5	0.1	0.2	0.1	55.9	58.0	53.0	
Larynx	3.4	5.9	1.3	1.0	1.8	0.4	60.8	61.8	56.7	
Lung & bronchus	61.7	72.4	53.8	49.1	61.4	39.8	17.1	14.7	19.8	
Pleura ^d	0.0	0.0	0.0	0.1	0.1	0.0	16.7	16.7	16.9	
Trachea & other	0.2	0.3	0.1	0.1	0.1	0.0	49.2	51.6	42.1	
respiratory organs										
Bones & joints	1.0	1.2	0.8	0.4	0.6	0.4	66.9	64.7	69.7	
Soft tissue (including heart)	3.4	4.2	2.8	1.3	1.5	1.1	66.5	65.6	67.6	
Skin (excl. basal & squamous):	27.2	35.3	21.4	4.1	6.3	2.4	90.4	88.3	93.0	
Melanoma of the skin	25.2	32.3	20.0	3.1	4.6	2.4	91.0	89.0	93.5	
Other non-epithelial skin	2.1	2.9	1.4	1.0	1.6	0.5	82.1	79.3	86.2	
time optimities shift	2.1	2.9		1.0	1.0	0.0	02.1			
Breast	68.3	1.2	128.0	12.1	0.3	21.7	90.4	85.7	90.4	
Breast (<i>in situ</i>)	16.8	0.1	32.1	-	-	-	100.0	100.0	100.0	

Note: Incidence and death rates are per 100,000 and are age-adjusted to the 2000 US Std Population (19 age groups - Census P25-1130). а SEER 18 areas (San Francisco, Connecticut, Detroit, Hawaii, Iowa, New Mexico, Seattle,

Utah, Atlanta, San Jose-Monterey, Los Angeles, Alaska Native Registry, Rural Georgia, California excluding SF/SJM/LA, Kentucky, Louisiana, New Jersey and Georgia excluding ATL/RG).

С

SEER 18 areas. Based on follow-up of patients into 2011. Mesotheliomas of the Pleura are included in the separate group Mesothelioma for incidence but are included in the Pleura grouping for mortality. d

Statistic could not be calculated due to less than 16 cases in the time interval.

b US Mortality Files, National Center for Health Statistics, Centers for Disease Control and Prevention.

Whites	

		2007-2011			Mortalit 007-2011			Survival ^c (2004-201	
Site	Total	Males	Females	Total	Males	Females	Total	Males	Females
Female Genital System: Cervix uteri Corpus uteri Uterus, NOS Ovary ^d Vagina Vulva	26.7 4.0 13.1 0.3 6.9 0.4 1.4		50.5 7.8 24.8 0.6 13.0 0.7 2.6	8.4 1.1 1.0 1.2 4.5 0.1 0.3		15.3 2.1 1.8 2.2 8.2 0.2 0.5	69.6 69.1 84.8 27.3 44.4 52.9 70.0		69.6 69.1 84.8 27.3 44.4 52.9 70.0
Other female genital system	0.5	-	1.0	0.1	-	0.2	59.9	-	59.9
Male Genital System: Prostate Testis Penis Other male genital system	68.1 64.2 3.4 0.4 0.1	147.7 139.9 6.6 0.9 0.3	- - - -	8.4 8.1 0.1 0.1 0.0	21.1 20.6 0.3 0.2 0.0	- - - -	98.9 99.3 95.4 66.9 88.4	98.9 99.3 95.4 66.9 88.4	- - - -
Urinary System: Urinary bladder Kidney & renal pelvis Ureter Other urinary system	39.2 22.4 15.9 0.6 0.3	62.5 39.4 21.7 0.9 0.5	21.0 9.5 11.0 0.4 0.2	8.9 4.6 4.0 0.1 0.1	14.3 8.1 5.9 0.2 0.2	5.0 2.2 2.6 0.1 0.1	75.2 77.9 72.6 48.1 51.9	76.3 79.2 72.4 49.0 53.7	72.8 73.7 73.1 47.1 48.2
Eye & Orbit	0.9	1.1	0.8	0.1	0.1	0.1	81.2	81.4	80.9
Brain & Nervous System: ^e Brain Cranial nerves & other nervous system	7.1 6.7 0.4	8.4 8.0 0.4	6.0 5.5 0.4	4.6 - -	5.6 - -	3.8 - -	32.0 29.1 78.5	31.3 28.8 76.9	32.8 29.4 80.1
Endocrine System: Thyroid Other endocrine & thymus	14.4 13.7 0.7	7.7 6.9 0.8	21.1 20.4 0.6	0.8 0.5 0.3	0.8 0.5 0.3	0.8 0.5 0.3	96.3 98.1 64.1	92.2 95.9 64.8	97.6 98.7 63.3
Lymphoma: Hodgkin lymphoma Non-Hodgkin lymphoma	23.6 2.9 20.6	28.2 3.3 24.9	19.8 2.6 17.2	7.0 0.4 6.6	8.9 0.5 8.4	5.5 0.3 5.2	72.1 85.7 69.9	71.1 85.0 68.9	73.3 86.6 71.2
Myeloma	5.6	7.2	4.3	3.1	4.0	2.5	44.8	46.7	42.4
Leukemia: Lymphocytic: Acute lymphocytic Chronic lymphocytic Other lymphocytic Myeloid & Monocytic: Acute myeloid Chronic myeloid Acute monocytic Other myeloid & monocytic Other leukemia: Other acute leukemia Aleukemic, subleukemic & NOS	13.77.01.94.70.46.13.91.70.30.20.60.20.4	17.59.22.16.40.77.64.82.20.40.20.20.80.30.5	10.7 5.3 1.7 3.4 0.2 4.9 3.3 1.2 0.2 0.2 0.1 0.5 0.2 0.3	$\begin{array}{c} 7.3\\ 2.0\\ 0.5\\ 1.4\\ 0.1\\ 3.5\\ 2.9\\ 0.3\\ 0.0\\ 0.2\\ 1.7\\ 0.6\\ 1.0 \end{array}$	9.7 2.9 0.6 2.1 0.2 4.6 3.8 0.4 0.0 0.3 2.2 0.8 1.4	5.4 1.4 0.4 1.0 0.1 2.7 2.3 0.2 0.0 0.2 1.3 0.5 0.8	57.4 76.6 80.2 82.2 35.2 24.2 60.5 22.8 32.7 29.8 16.1 39.2	58.1 76.7 66.9 79.6 86.1 34.9 23.2 60.1 22.2 31.9 28.9 15.7 39.1	56.5 76.3 66.2 81.1 71.2 35.6 25.3 61.1 23.4 33.5 30.5 16.6 38.9
Kaposi Sarcoma ^f Mesothelioma ^f	0.5 1.1	0.9 2.1	0.1 0.5	- -	-	-	78.0 8.2	77.2 6.5	84.8 13.7
Ill-defined & unspecified	9.0	10.5	7.9	12.9	16.4	10.3	17.8	22.6	13.1

Note: Incidence and death rates are per 100,000 and are age-adjusted to the 2000 US Std Population (19 age groups - Census P25-1130). a SEER 18 areas (San Francisco, Connecticut, Detroit, Hawaii, Iowa, New Mexico, Seattle, Utah, Atlanta, San Jose-Monterey, Los Angeles, Alaska Native Registry, Rural Georgia,

California excluding SF/SJM/LA, Kentucky, Louisiana, New Jersey and Georgia excluding ATL/RG).

b US Mortality Files, National Center for Health Statistics, Centers for Disease Control and Prevention.

- С d
- е

SEER 18 areas. Based on follow-up of patients into 2011. Ovary excludes borderline cases or histologies 8442, 8451, 8462, 8472, and 8473. Due to coding changes, Brain & Nervous System mortality are no longer shown separately. Rate not shown for mortality. Category did not exist in mortality coding until 1999. Statistic could not be calculated due to less than 16 cases in the time interval. f

Table 1.7 Age-Adjusted SEER Incidence and U.S. Death Rates and 5-Year Relative Survival (Percent) By Primary Cancer Site, Sex and Time Period

Blacks

			BIACKS							
	Incidence ^a US Mortality ^b						Survival ^c (%)			
		2007-201			2007-201			(2004-20		
Site	Total		Females	Total		Females	Total		Females	
All Sites	480.8	600.9	398.8	206.4	269.3	169.0	59.7	62.7	56.4	
Oral Cavity & Pharynx:	9.4	14.7	5.3	3.0	5.1	1.4	43.7	39.9	52.0	
Lip	0.1	0.1	0.1	-	_	-	79.4	72.3	89.6	
Tongue	2.2	3.6	1.1	0.6	0.9	0.3	35.9	34.5	38.2	
Salivary gland	1.0	1.1	1.0	0.2	0.3	0.1	73.7	66.4	79.5	
Floor of mouth	0.5	0.9	0.3	0.0	0.1	-	38.3	35.4	46.4	
Gum & other oral cavity	1.4	1.7	1.2	0.3	0.5	0.2	52.7	47.7	57.8	
Nasopharynx	0.7 1.6	1.1	0.3	0.3	0.4	0.1	53.4	52.3	56.4 48.2	
Tonsil		2.9	0.6	0.3	0.5	0.1	49.5	49.7		
Oropharynx	0.6 1.0	1.1 1.9	0.3	0.4	0.6	0.2 0.1	22.0	21.6	23.3 27.1	
Hypopharynx Othor aral gouitu (pharunu	0.3	1.9	0.3 0.2	0.2 0.8	0.3 1.5	0.1	18.3 20.4	16.6 20.5	27.1 21.9	
Other oral cavity & pharynx	0.5	0.4	0.2	0.0	1.5	0.5	20.4	20.5	21.9	
Digestive System:	104.7	129.4	86.9	56.8	74.9	44.3	39.1	35.8	42.6	
Esophagus	4.9	7.9	2.6	4.2	7.4	2.0	11.6	10.7	13.7	
Stomach	11.2	15.3	8.5	6.5	9.6	4.5	27.2	24.0	31.2	
Small intestine	3.5	4.2	3.0	0.6	0.7	0.5	63.0	59.0	66.6	
Colon & Rectum:	53.6	62.3	47.5	22.1	27.7	18.5	57.7	56.6	58.8	
Colon	40.1	45.4	36.4	-	-	-	56.2	55.8	56.7	
Rectum	13.5	16.9	11.1	-	-	-	62.0	58.5	65.6	
Anus, anal canal & anorectum	1.8	2.0	1.6	0.2	0.3	0.2	55.7	49.1	62.2	
Liver & intrahepatic	9.4	15.6	4.6	7.6	12.1	4.2	11.5	10.1	15.3	
bile duct	1 5	1 0	1 0	0 0	0 7	1 0	14.4	14 0	14 5	
Gallbladder	1.5	1.2	1.8	0.9	0.7	1.0	14.4	14.0	14.5	
Other biliary	1.8	2.1	1.6	0.4	0.4	0.4	12.4	10.6	13.9	
Pancreas	15.6 0.4	17.2 0.3	14.2 0.4	13.6 0.0	15.3 0.0	12.4 0.0	6.1 45.2	5.8 48.4	6.3 43.5	
Retroperitoneum Peritoneum, omentum &	0.4	0.3	0.4	0.0	0.0	0.0	45.2 31.3	40.4	28.2	
mesentery	0.4	0.1	0.5	0.2	0.1	0.2	31.3	45.2	20.2	
Other digestive system	0.7	1.0	0.6	0.4	0.5	0.3	10.8	9.5	12.1	
Other digestive system	0.7	1.0	0.0	0.4	0.5	0.5	10.0	9.5	12.1	
Respiratory System:	73.6	103.2	53.5	54.2	79.9	37.3	17.3	16.7	18.2	
Nose, nasal cavity &	0.6	0.9	0.4	0.2	0.2	0.1	44.9	42.3	49.2	
middle ear										
Larynx	4.8	9.0	1.7	1.9	3.8	0.6	53.6	54.1	51.6	
Lung & bronchus	68.0	93.0	51.2	52.0	75.7	36.5	13.9	11.8	16.5	
Pleura ^d	-	-	-	0.0	0.1	-	-	-	-	
Trachea & other	0.2	0.3	0.1	0.1	0.1	0.0	46.6	44.6	49.5	
respiratory organs										
Bones & joints	0.8	0.9	0.7	0.4	0.6	0.3	64.4	63.3	65.4	
Soft tissue (including heart)	3.3	3.4	3.2	1.4	1.5	1.4	58.7	57.6	59.7	
Skin (excl. basal & squamous):	2.1	2.2	2.0	0.9	1.3	0.6	84.7	83.5	85.0	
Melanoma of the skin	1.0	1.0	1.0	0.4	0.5	0.4	71.1	60.8	76.9	
Other non-epithelial skin	1.1	1.2	1.1	0.5	0.8	0.2	94.6	98.3	90.8	
Propet	70 0	1 0	100 0	10 0	0 5	20 E	70 1	67 0	70 1	
Breast	70.0	1.8	122.8	18.0	0.5	30.6	79.1	67.8	79.1	
Breast (<i>in situ</i>)	16.8	0.3	29.9	-	-	-	100.0	98.2	100.0	

Note: Incidence and death rates are per 100,000 and are age-adjusted to the 2000 US Std Population (19 age groups - Census P25-1130). а SEER 18 areas (San Francisco, Connecticut, Detroit, Hawaii, Iowa, New Mexico, Seattle, Utah, Atlanta, San Jose-Monterey, Los Angeles, Alaska Native Registry, Rural Georgia, California excluding SF/SJM/LA, Kentucky, Louisiana, New Jersey and

С d

SEER 18 areas. Based on follow-up of patients into 2011. Mesotheliomas of the Pleura are included in the separate group Mesothelioma for incidence but are included in the Pleura grouping for mortality.

Statistic could not be calculated due to less than 16 cases in the time interval.

Georgia excluding ATL/RG).

b US Mortality Files, National Center for Health Statistics, Centers for Disease Control and Prevention.

Age-Adjusted SEER Incidence and U.S. Death Rates and 5-Year Relative Survival (Percent) By Primary Cancer Site, Sex and Time Period

	(:	ncidence 2007-2011)	(2	Mortalit 007-2011)		Survival ^c (2004-201	.0)
Site	Total	Males	Females	Total	Males	Females	Total	Males	Females
Female Genital System: Cervix uteri Corpus uteri Uterus, NOS Ovary ^d Vagina Vulva Other female genital system	26.0 5.2 12.5 0.8 5.6 0.5 1.0 0.4		45.9 9.4 21.9 1.3 9.8 0.9 1.7 0.8	11.3 2.3 1.8 2.7 3.9 0.2 0.2 0.1		19.1 4.1 3.0 4.5 6.6 0.3 0.3 0.3	55.0 58.8 63.5 22.1 35.3 45.8 70.9 50.3		55.0 58.8 63.5 22.1 35.3 45.8 70.9 50.3
Male Genital System: Prostate Testis Penis Other male genital system	95.9 94.8 0.7 0.4 0.1	226.4 223.9 1.4 0.9 0.2	- - - -	17.5 17.4 0.1 0.1 0.0	49.4 48.9 0.1 0.3 0.1	- - - -	96.6 96.7 90.9 61.1 75.4	96.6 96.7 90.9 61.1 75.4	- - - -
Urinary System: Urinary bladder Kidney & renal pelvis Ureter Other urinary system	31.1 12.6 17.9 0.3 0.4	47.0 21.3 24.7 0.4 0.6	20.2 6.9 12.7 0.2 0.3	7.7 3.6 3.9 0.1 0.1	11.2 5.4 5.6 0.1 0.1	5.4 2.6 2.6 0.0 0.1	67.5 63.8 70.6 41.0 37.6	69.3 68.6 70.2 34.8 48.5	64.5 54.7 71.0 45.2 29.2
Eye & Orbit	0.2	0.3	0.2	0.0	0.0	0.0	81.9	79.8	84.8
Brain & Nervous System: ^e Brain Cranial nerves & other nervous system	4.1 3.7 0.4	4.7 4.4 0.3	3.6 3.3 0.4	2.5 - -	3.0 - -	2.1 - -	40.4 36.8 71.9	37.8 34.9 68.1	43.0 38.7 73.3
Endocrine System: Thyroid Other endocrine & thymus	8.5 7.6 0.9	4.3 3.3 1.0	12.1 11.3 0.9	0.9 0.5 0.4	0.8 0.4 0.4	0.9 0.6 0.3	92.8 96.7 63.2	83.1 91.4 59.9	95.4 97.6 66.1
Lymphoma: Hodgkin lymphoma Non-Hodgkin lymphoma	17.0 2.7 14.3	20.5 3.1 17.4	14.2 2.3 11.9	4.8 0.3 4.5	6.2 0.4 5.8	3.8 0.3 3.5	66.0 81.6 62.2	62.5 79.2 58.4	70.1 84.4 66.6
Myeloma	12.2	14.8	10.5	6.3	7.7	5.3	44.9	44.1	45.8
Leukemia: Lymphocytic: Acute lymphocytic Chronic lymphocytic Other lymphocytic Myeloid & Monocytic: Acute myeloid Chronic myeloid Acute monocytic Other myeloid & monocytic Other leukemia: Other acute leukemia Aleukemic, subleukemic & NOS	10.0 4.2 1.0 3.0 0.2 5.1 3.3 1.6 0.1 0.1 0.7 0.2 0.5	12.9 5.8 1.2 4.2 0.4 6.1 3.8 1.9 0.2 0.2 1.0 0.3 0.6	8.0 3.0 0.8 2.1 0.1 4.5 2.9 1.3 0.1 0.1 0.1 0.6 0.2 0.4	$\begin{array}{c} 6.0\\ 1.7\\ 0.3\\ 1.3\\ 0.1\\ 2.7\\ 2.2\\ 0.3\\ 0.0\\ 0.2\\ 1.7\\ 0.5\\ 1.2 \end{array}$	8.0 2.4 0.4 1.9 0.1 3.4 2.7 0.5 - 0.2 2.2 0.7 1.5	4.8 1.2 0.9 0.1 2.2 1.8 0.3 - 0.1 1.4 0.4 1.0	50.7 66.1 64.1 68.2 51.4 39.8 26.3 64.6 24.1 38.3 29.6 26.8 30.8	51.8 65.8 66.1 66.3 57.7 39.7 25.9 62.8 13.6 38.1 32.1 28.1 35.0	49.3 66.3 61.0 70.7 32.8 39.8 26.4 66.4 34.7 39.0 26.4 24.2 26.8
Kaposi Sarcoma ^f Mesothelioma ^f	1.1 0.5	2.1 1.0	0.2	- -	-	- -	53.8 10.2	53.4 8.2	56.5 15.0
Ill-defined & unspecified	10.1	11.4	9.1	14.7	19.2	11.7	12.7	12.9	12.5

Note: Incidence and death rates are per 100,000 and are age-adjusted to the 2000 US Std Population (19 age groups - Census P25-1130). а SEER 18 areas (San Francisco, Connecticut, Detroit, Hawaii, Iowa, New Mexico, Seattle, Utah, Atlanta, San Jose-Monterey, Los Angeles, Alaska Native Registry, Rural Georgia,

California excluding SF/SJM/LA, Kentucky, Louisiana, New Jersey and Georgia excluding ATL/RG).

b US Mortality Files, National Center for Health Statistics, Centers for Disease Control and Prevention.

С d

е

SEER 18 areas. Based on follow-up of patients into 2011. Ovary excludes borderline cases or histologies 8442, 8451, 8462, 8472, and 8473. Due to coding changes, Brain & Nervous System mortality are no longer shown separately. Rate not shown for mortality. Category did not exist in mortality coding until 1999. Statistic could not be calculated due to less than 16 cases in the time interval. f

				Tal	ole 1.8						
SEER	Incidence	and	U.S.	Mortality	Trends	by	Primary	Cancer	Site	and	Sex
				All Race	s, 2002	-20	11				

		Incidence	a	Ū	JS Mortalit	y ^b
	Total	Males	Females	Total	Males	Females
Site	APC	APC	APC	APC	APC	APC
All Sites	-0.6*	-1.2*	-0.2	-1.5*	-1.8*	-1.4*
Oral Cavity & Pharynx:	0.4*	0.4*	0.1	-1.0*	-0.9*	-1.5*
Lip	-3.5*	-4.1*	-2.1*	-1.8	-1.8	-2.0
Tongue	1.8*	2.2*	0.9	-0.8*	-0.6*	-1.4*
Salivary gland	0.5	0.4	0.3	-0.1	0.0	-0.6
Floor of mouth	-3.2*	-3.5*	-2.2	-8.2*	-8.9*	-5.8*
Gum & other oral cavity	-0.5	-0.8	0.0	-1.6*	-1.7*	-1.4*
Nasopharynx	-0.9	-0.8	-1.3	-1.1*	-0.6	-2.5*
Tonsil	2.9*	3.1*	1.9*	1.2*	1.5*	-0.3
Oropharynx	2.6*	3.0*	1.3	1.4*	1.0	1.8
Hypopharynx	-2.5*	-2.7*	-2.2*	-2.8*	-2.9*	-2.2
Other oral cavity & pharynx	-0.8	-0.5	-2.5	-2.4*	-2.1*	-3.7*
Digestive System:	-1.2*	-1.2*	-1.3*	-1.1*	-1.0*	-1.3*
Esophagus	-0.9*	-0.7	-2.2*	-0.7*	-0.6*	-1.6*
Stomach	-1.3*	-1.6*	-1.0*	-2.9*	-3.2*	-2.7*
Small intestine	1.8*	1.4*	2.3*	0.1	0.2	-0.2
Colon & Rectum:	-2.8*	-3.0*	-2.6*	-2.9*	-3.0*	-2.9*
Colon	-3.1*	-3.4*	-2.9*		_	_
Rectum	-2.0*	-2.2*	-1.9*	-	-	-
Anus, anal canal & anorectum	1.9*	1.3	2.4*	3.6*	4.2*	3.3*
Liver & intrahepatic	3.4*	3.7*	2.5*	2.5*	2.6*	1.8*
bile duct						
Gallbladder	-0.2	-0.4	-0.1	-0.9*	-0.2	-1.2*
Other biliary	1.0*	1.5*	0.2	-2.1*	-2.4*	-1.9*
Pancreas	0.7*	0.9*	0.6*	0.4*	0.3*	0.4*
Retroperitoneum	-1.0	-0.6	-1.6	-2.3*	-2.3	-2.4*
Peritoneum, omentum &	-1.2	0.1	-1.2	0.6	1.6	0.4
mesentery Other digestive system	1.8*	2.5*	1.2	0.2	0.5	0.0
Respiratory System:	-1.7*	-2.3*	-1.0*	-2.0*	-2.6*	-1.3*
Nose, nasal cavity &	-0.4	-0.4	-0.5	-1.1	-1.6	-0.3
middle ear	0.1	0.1	0.0		2.00	0.5
Larynx	-1.9*	-2.1*	-1.8*	-2.3*	-2.6*	-2.1*
Lung & bronchus	-1.7*	-2.4*	-1.0*	-1.9*	-2.6*	-1.2*
Pleura	-3.1	-2.1	-	-5.5*	-5.6*	-5.9*
Trachea & other	-1.1	-0.6	-1.9	-3.8*	-4.1*	-3.5
respiratory organs	1.1	-0.0	-1.9	-5.0	-1.1	-5.5
Bones & joints	-0.1	-0.1	-0.2	0.0	0.2	-0.2
Soft tissue (including heart)	0.8*	0.8*	0.8	0.7*	1.1*	0.3
Skin (excl. basal & squamous):	1.3*	1.7*	0.8	0.5*	0.8*	-0.3
Melanoma of the skin	1.4*	1.6*	0.9	0.4	0.7*	-0.3
Other non-epithelial skin	1.2	1.9*	0.0	1.0*	1.2*	-0.1
Breast	-0.3	1.8*	-0.2	-2.1*	-0.3	-2.0*
Breast (in situ)	0.9*	2.6	1.1*	-	-	-

- Trends are based on rates age-adjusted to the 2000 US Std Population
- (19 age groups Census P25-1130).

SEER 18 areas (San Francisco, Connecticut, Detroit, Hawaii, Iowa, New Mexico, Seattle, Utah, Atlanta, San Jose-Monterey, Los Angeles, Alaska Native Registry, Rural Georgia, California excluding SF/SJM/LA, Kentucky, Louisiana, New Jersey and a Georgia excluding ATL/RG).

b US Mortality Files, National Center for Health Statistics, Centers for Disease Control and

Prevention.

⁻

The APC is significantly different from zero (p<.05). Statistic could not be calculated. Trend based on less than 10 cases for at least one year within the time interval.

				Table 1.8	3 – cont	cinι	led				
SEER I	Incidence	and	U.S.	Mortality	Trends	by	Primary	Cancer	Site	and	Sex
				All Race	s, 2002	-20	11 -				

		Incidence	a	U	S Mortalit	yb
Site	Total APC	Males APC	Females APC	Total APC	Males APC	Females APC
Female Genital System:	-0.1	-	0.0	-1.2*	-	-1.0*
Cervix uteri	-1.7*	-	-1.6*	-1.3*	-	-1.1*
Corpus uteri	1.0*	-	1.2*	-0.7*	-	-0.4
Uterus, NOS	2.8*	-	3.3*	1.4*	-	1.8*
Ovary ^c	-1.9*	-	-1.7*	-2.2*	-	-2.0*
Vagina	0.2	-	0.5	-1.0*	-	-0.7
Vulva	0.9*	-	1.1*	0.9*	-	1.4*
Other female genital system	5.6*	-	5.6*	1.7	-	1.9
Male Genital System:	-1.7*	-2.2*	-	-2.6*	-3.3*	_
Prostate	-1.9*	-2.3*	-	-2.6*	-3.3*	-
Testis	0.6*	0.6*	-	-0.5	-0.6	-
Penis	0.8	0.4	-	-0.9	-1.1	-
Other male genital system	0.7	0.6	-	3.2	2.7	-
Urinary System:	0.3	0.1	0.4	-0.3*	-0.3*	-0.9*
Urinary bladder	-0.8*	-0.8*	-1.2*	0.1	0.1	-0.6*
Kidney & renal pelvis	1.9*	1.8*	1.8*	-0.9*	-0.8*	-1.2*
	-0.2	-0.3	-0.4	-0.5	-0.4	-0.9
Ureter Other urinary system	2.2*	2.0	2.5	1.5	2.6	-0.4
Other utiliary system	2.2	2.0	2.5	1.5	2.0	-0.4
Eye & Orbit	-0.9*	-1.3*	-0.5	0.8	0.1	1.2
Brain & Nervous System: ^d	-0.6*	-0.5*	-0.7*	-0.4	-0.4	-0.5*
Brain	-0.5	-0.5	-0.6	-	-	-
Cranial nerves & other nervous system	-1.6	-1.2	-1.8	-	-	-
Endocrine System:	5.5*	4.7*	5.8*	0.7*	1.0*	0.4
Thyroid	5.8*	5.3*	6.0*	1.3*	1.8*	0.9
Other endocrine & thymus	0.7	0.2	0.9	-0.3	-0.2	-0.5
Lymphoma:	-0.2	-0.1	-0.3	-2.6*	-2.3*	-3.0*
Hodgkin lymphoma	-0.4	-0.3	-0.5	-2.8*	-2.4*	-3.5*
Non-Hodgkin lymphoma	-0.1	0.0	-0.3	-2.6*	-2.3*	-3.0*
Myeloma	0.7*	0.7*	0.6	-1.5*	-1.3*	-1.9*
Leukemia:	0.0	-0.3	0.1	-0.9*	-1.0*	-1.0*
Lymphocytic:	-0.4	-0.7*	-0.1	-1.5*	-1.6*	-1.5*
Acute lymphocytic	1.1*	0.6	1.8*	-1.2*	-1.7*	-0.5
Chronic lymphocytic	-0.9*	-1.1*	-0.8	-1.6*	-1.6*	-1.9*
Other lymphocytic	-1.2	-1.1	-2.1	-1.0	-1.5*	-1.1
Myeloid & Monocytic:	0.9*	0.8*	0.9*	-0.2	-0.3	-0.4
Acute myeloid	1.3*	0.9	1.7*	0.3*	0.3	0.1
Chronic myeloid	1.0*	1.1*	0.5	-4.4*	-4.2*	-4.7*
Acute monocytic	-3.1*	-2.1	-4.1*	-3.8*	-4.9*	-3.2
Other myeloid & monocytic	-1.2	0.8	-3.8*	0.3	-0.2	0.2
Other leukemia:	-1.2	-4.2*	-4.1*	-1.6*	-0.2 -1.6*	-1.8*
	-4.1* -4.6*	-4.2*	-4.1* -5.1*	-1.6*	-1.6*	-1.8^ -4.1*
Other acute leukemia						
Aleukemic, subleukemic & NOS	-3.7*	-4.1*	-3.3	0.1	0.3	-0.1
Kaposi Sarcoma ^e	-3.1*	-3.3*	-1.7	-	-	-
Mesothelioma ^e	-1.0	-1.3	-0.4	-	-	-
Ill-defined & unspecified	-2.8*	-2.5*	-3.1*	-2.2*	-2.1*	-2.4*

Trends are based on rates age-adjusted to the 2000 US Std Population

(19 age groups - Census P25-1130). а

SEER 18 areas (San Francisco, Connecticut, Detroit, Hawaii, Iowa, New Mexico, Seattle, Utah, Atlanta, San Jose-Monterey, Los Angeles, Alaska Native Registry, Rural Georgia, California excluding SF/SJM/LA, Kentucky, Louisiana, New Jersey and Georgia excluding ATL/RG).

b US Mortality Files, National Center for Health Statistics, Centers for Disease Control and Prevention. с

Ovary excludes borderline cases or histologies 8442, 8451, 8462, 8472, and 8473. d

e

Due to coding changes, Brain & Nervous System mortality are no longer shown separately. Trend not shown for mortality. Category did not exist in mortality coding until 1999. The APC is significantly different from zero (p<.05). Statistic could not be calculated. Trend based on less than 10 cases for at least one

_ year within the time interval.

				Tal	ole 1.9						
SEER	Incidence	and	U.S.	Mortality	Trends	by	Primary	Cancer	Site	and	Sex
				Whites	, 2002-2	2011	L				

		Incidence	a	Ŭ	JS Mortalit	y ^b
	Total	Males	Females	Total	Males	Females
Site	APC	APC	APC	APC	APC	APC
All Sites	-0.6*	-1.2*	-0.2	-1.4*	-1.7*	-1.4*
Oral Cavity & Pharynx:	0.8*	0.8*	0.4	-0.6*	-0.4	-1.3*
Lip	-3.7*	-4.3*	-2.1*	-1.5	-1.6	-1.9
Tongue	2.1*	2.4*	1.3*	-0.4	-0.1	-1.1*
Salivary gland	0.9	0.7	0.7	-0.1	-0.1	-0.4
Floor of mouth	-2.5*	-3.0*	-1.3	-7.6*	-8.3*	-5.8
Gum & other oral cavity	-0.4	-0.7	0.0	-1.2*	-1.1	-1.2
Nasopharynx	-0.5	-0.8	-0.2	-1.1	-0.6	-2.6*
Tonsil	3.8*	4.0*	2.5*	2.0*	2.3*	0.1
Oropharynx	3.1*	3.4*	1.8	2.0*	1.7*	2.0*
Hypopharynx	-2.1*	-2.4*	-2.0*	-2.9*	-2.6	-4.1*
Other oral cavity & pharynx	-0.5	0.3	-3.3	-1.8*	-1.5*	-3.2*
Digestive System:	-1.2*	-1.2*	-1.3*	-1.0*	-1.0*	-1.2*
Esophagus	-0.3	-0.1	-1.9*	-0.1	-0.1	-0.9*
Stomach	-1.0*	-1.2*	-1.0*	-2.9*	-3.3*	-2.7*
Small intestine	1.9*	1.6*	2.3*	0.0	0.1	-0.3
Colon & Rectum:	-2.9*	-3.2*	-2.7*	-2.9*	-3.1*	-2.9*
Colon	-3.2*	-3.5*	-3.0*	-	-	-
Rectum	-2.3*	-2.6*	-2.1*	-	-	-
Anus, anal canal & anorectum	2.0*	1.0	2.7*	3.6*	3.7*	3.6*
Liver & intrahepatic bile duct	4.1*	4.3*	3.1*	2.6*	2.8*	1.9*
Gallbladder	-0.1	-0.3	0.0	-1.3*	-1.0*	-1.4*
Other biliary	0.9*	1.4*	0.2	-2.2*	-2.4*	-2.2*
Pancreas	0.9*	1.0*	0.7*	0.5*	0.5*	0.5*
Retroperitoneum	-1.4	-1.6	-1.4	-2.0	-2.1	-2.0
Peritoneum, omentum &	-1.1	0.7	-1.1	0.8	2.2	0.6
mesentery		0.7		0.0	2.2	0.0
Other digestive system	1.4*	1.7*	1.1	0.1	0.4	-0.1
Respiratory System:	-1.6*	-2.2*	-0.9*	-1.8*	-2.5*	-1.1*
Nose, nasal cavity &	-0.1	0.0	-0.6	-1.4	-1.6	-1.3
middle ear						
Larynx	-1.6*	-1.8*	-1.5*	-2.0*	-2.4*	-1.8*
Lung & bronchus	-1.6*	-2.3*	-0.9*	-1.8*	-2.5*	-1.1*
Pleura	-3.4*	-1.8	-	-5.6*	-5.7*	-6.1*
Trachea & other	-1.2	-1.3	-1.2	-3.9*	-4.2*	-3.6
respiratory organs						
Bones & joints	0.3	0.4	0.1	0.1	0.2	-0.1
Soft tissue (including heart)	0.9*	1.0*	0.7	0.7*	1.2*	0.2
Skin (excl. basal & squamous):	1.3*	1.6*	0.9	0.7*	1.0*	-0.1
Melanoma of the skin	1.3*	1.6*	0.9	0.6*	0.9*	-0.1
Other non-epithelial skin	1.3*	1.9*	0.2	1.3*	1.5*	0.1
Breast	-0.5	1.8*	-0.3	-2.2*	-0.4	-2.0*
Breast (<i>in situ</i>)	0.6	2.3	0.8	-	-	-

- Trends are based on rates age-adjusted to the 2000 US Std Population
- (19 age groups Census P25-1130).

SEER 18 areas (San Francisco, Connecticut, Detroit, Hawaii, Iowa, New Mexico, Seattle, Utah, Atlanta, San Jose-Monterey, Los Angeles, Alaska Native Registry, Rural Georgia, California excluding SF/SJM/LA, Kentucky, Louisiana, New Jersey and a Georgia excluding ATL/RG).

b

US Mortality Files, National Center for Health Statistics, Centers for Disease Control and Prevention. -

The APC is significantly different from zero (p<.05). Statistic could not be calculated. Trend based on less than 10 cases for at least one year within the time interval.

Table 1.9 - continued											
SEER	Incidence	and	U.S.	Mortality	Trends	by	Primary	Cancer	Site	and	Sex
				Whites	, 2002-2	2011	L				

		Incidence	a	US Mortality ^b				
	Total	Males	Females	Total	Males	Females		
Site	APC	APC	APC	APC	APC	APC		
Female Genital System:	-0.1	_	0.1	-1.2*	_	-1.0*		
Cervix uteri	-1.3*	_	-1.2*	-1.0*	-	-0.8*		
Corpus uteri	0.8*	_	1.0*	-0.7*	-	-0.3		
Uterus, NOS	1.9	_	2.5	1.2*	-	1.7*		
Ovary ^c	-1.8*	_	-1.6*	-2.2*	_	-2.0*		
Vagina	0.7	_	1.1*	-0.7	-	-0.3		
Vulva	1.2*	_	1.4*	1.0*	_	1.5*		
	6.3*	_	6.4*		-	2.4		
Other female genital system	0.3*	-	0.4~	2.1	-	2.4		
Male Genital System:	-2.0*	-2.5*	-	-2.4*	-3.2*	-		
Prostate	-2.2*	-2.6*	-	-2.5*	-3.3*	-		
Testis	0.6*	0.6*	-	0.3	0.2	-		
Penis	0.9	0.4	-	-1.0	-1.2	-		
Other male genital system	0.6	0.3	-	3.6	3.1	-		
Urinary System:	0.3	0.0	0.4	-0.2*	-0.2*	-0.8*		
Urinary bladder	-0.7*	-0.8*	-1.1*	0.3*	0.2	-0.4		
Kidney & renal pelvis	1.9*	1.8*	1.7*	-0.8*	-0.8*	-1.2*		
Ureter	0.0	-0.1	-0.2	-0.5	-0.5	-0.8		
Other urinary system	2.4*	1.9	3.2	2.0	2.7	0.2		
Other armary system	2.1	1.9	5.2	2.0	2.1	0.2		
Eye & Orbit	-0.9*	-1.4*	-0.4	0.9	0.2	1.4		
Brain & Nervous System: ^d	-0.5	-0.5	-0.7*	-0.2	-0.2	-0.4		
Brain	-0.5	-0.5	-0.6*	-	-	-		
Cranial nerves & other nervous system	-1.3	-1.0	-1.6	-	-	-		
Endocrine System:	5.5*	4.7*	5.8*	0.5*	0.8*	0.2		
Thyroid	5.8*	5.4*	6.0*	1.0*	1.7*	0.5		
Other endocrine & thymus	0.3	-0.3	0.8	-0.4	-0.5	-0.5		
Lymphoma:	-0.2	-0.1	-0.4	-2.6*	-2.3*	-3.0*		
Hodgkin lymphoma	-0.5	-0.4	-0.7	-2.7*	-2.2*	-3.4*		
Non-Hodgkin lymphoma	-0.2	-0.1	-0.3	-2.6*	-2.4*	-3.0*		
Myeloma	0.6*	0.7*	0.4	-1.5*	-1.3*	-1.9*		
Leukemia:	0.0	-0.3	0.2	-0.8*	-0.9*	-0.9*		
Lymphocytic:	-0.5	-0.9*	0.0	-1.3*	-1.5*	-1.3*		
Acute lymphocytic	1.5*	0.9	2.4*	-1.0*	-1.6*	-0.2		
Chronic lymphocytic	-1.1*	-1.4*	-1.0	-1.5*	-1.5*	-1.7*		
	-1.7	-1.5	-2.4	-0.9	-1.2	-1.3		
Other lymphocytic	1.0*	0.9*	0.9*	-0.2	-0.2	-0.3		
Myeloid & Monocytic:	1.4*		1.7*			0.2		
Acute myeloid		0.9		0.3*	0.3			
Chronic myeloid	0.9*	1.1*	0.4	-4.2*	-3.9*	-4.7*		
Acute monocytic	-2.6	-1.2	-4.3*	-3.8*	-5.1*	-3.0		
Other myeloid & monocytic	-1.0	1.3	-3.9*	0.3	-0.3	0.4		
Other leukemia:	-3.9*	-4.4*	-3.4*	-1.5*	-1.6*	-1.7*		
Other acute leukemia	-4.3*	-4.6*	-4.3*	-4.1*	-4.3*	-4.0*		
Aleukemic, subleukemic & NOS	-3.5*	-4.3*	-2.9	0.2	0.3	-0.1		
Kaposi Sarcoma ^e	-3.7*	-3.9*	-2.3	-	-	-		
Mesothelioma ^e	-0.9	-1.2	-0.3	-	-	-		
Ill-defined & unspecified	-2.7*	-2.4*	-3.1*	-2.0*	-1.9*	-2.3*		

Trends are based on rates age-adjusted to the 2000 US Std Population

(19 age groups - Census P25-1130).

а SEER 18 areas (San Francisco, Connecticut, Detroit, Hawaii, Iowa, New Mexico, Seattle, Utah, Atlanta, San Jose-Monterey, Los Angeles, Alaska Native Registry, Rural Georgia, California excluding SF/SJM/LA, Kentucky, Louisiana, New Jersey and Georgia excluding ATL/RG).

b US Mortality Files, National Center for Health Statistics, Centers for Disease Control and Prevention. с

Ovary excludes borderline cases or histologies 8442, 8451, 8462, 8472, and 8473. d

e

Due to coding changes, Brain & Nervous System mortality are no longer shown separately. Trend not shown for mortality. Category did not exist in mortality coding until 1999. The APC is significantly different from zero (p<.05). Statistic could not be calculated. Trend based on less than 10 cases for at least one

_ year within the time interval.

				Tab	le 1.10						
SEER	Incidence	and	U.S.	Mortality	Trends	by	Primary	Cancer	Site	and	Sex
				Blacks	, 2002-2	2011	1				

		Incidence	a	Ŭ	US Mortality ^b				
	Total	Males	Females	Total	Males	Females			
Site	APC	APC	APC	APC	APC	APC			
All Sites	-1.0*	-1.9*	-0.2	-2.1*	-2.6*	-1.7*			
Oral Cavity & Pharynx:	-2.5*	-2.9*	-1.7	-3.4*	-3.7*	-2.7*			
Lip	-	-	-	-	-	-			
Tongue	-1.2	-1.2	-1.2	-3.7*	-4.4*	-2.1			
Salivary gland	-0.9	-0.8	-1.1	1.2	2.3	-1.1			
Floor of mouth	-6.5*	-6.7*	-5.4*	-11.6*	-	-			
Gum & other oral cavity	-3.7*	-5.5*	-1.5	-5.4*	-5.8*	-4.3*			
Nasopharynx	-3.1*	-2.9	-3.3	-3.4*	-2.8	-4.8			
Tonsil	-1.1	-1.3	-0.3	-1.5	-1.7*	-1.3			
Oropharynx	-0.2	-0.1	-	-0.5	-1.2	1.0			
Hypopharynx	-5.0*	-5.2*	-4.4	-2.5	-4.6	-			
Other oral cavity & pharynx	-3.9	-6.8*	-	-4.8*	-4.5*	-5.7*			
Digestive System:	-1.5*	-1.5*	-1.6*	-1.6*	-1.5*	-1.9*			
Esophagus	-4.5*	-4.8*	-3.7*	-4.6*	-4.3*	-5.3*			
Stomach	-2.1*	-3.1*	-1.1	-3.3*	-3.3*	-3.5*			
Small intestine	2.1*	1.9	2.4	1.0	1.3	0.5			
Colon & Rectum:	-2.7*	-2.8*	-2.7*	-2.8*	-2.6*	-3.2*			
Colon	-3.1*	-3.3*	-3.0*	-	-	-			
Rectum	-1.3*	-1.0	-1.8*	-	-	-			
Anus, anal canal & anorectum	2.3*	3.3*	1.2	4.4*	7.0*	1.8			
Liver & intrahepatic bile duct	3.4*	3.6*	2.5*	2.5*	2.7*	1.7*			
Gallbladder	1.1*	3.7	0.9	0.9	4.7*	-0.3			
Other biliary	2.4*	2.0	2.8	-1.0	-2.0	-0.2			
Pancreas	-0.2	0.3	-0.5	-0.3	-0.3	-0.2			
Retroperitoneum	0.5	_	-0.4	-6.1	_	_			
Peritoneum, omentum &	-0.4	-	0.2	-1.2	-	-1.2			
mesentery									
Other digestive system	2.6	6.0	-0.2	0.8	-0.6	1.7			
Respiratory System:	-2.0*	-2.8*	-0.8	-2.6*	-3.4*	-1.6*			
Nose, nasal cavity &	-1.7	-2.1	-0.6	1.6	-1.5	6.5*			
middle ear									
Larynx	-3.5*	-3.5*	-3.5	-3.8*	-3.9*	-3.1*			
Lung & bronchus	-1.9*	-2.8*	-0.7	-2.6*	-3.4*	-1.6*			
Pleura	-	-	-	-	-	-			
Trachea & other	-	-	-	-3.7	-	-			
respiratory organs									
Bones & joints	-1.5	-1.4	-1.0	-0.2	0.1	-0.6			
Soft tissue (including heart)	-0.4	-1.5	0.6	0.4	0.2	0.6			
Skin (excl. basal & squamous):	-1.1	-1.2	-1.0	-1.5*	-2.2	-0.4			
Melanoma of the skin	-0.9	-2.4	-0.1	-0.3	-0.9	0.0			
Other non-epithelial skin	-1.1	-0.2	-1.8	-2.6*	-3.0*	-1.3			
other non-epitherrar skill		-0.2	-1.0	-2.0	-3.0	-1.5			
Breast	0.3	1.5	0.5	-1.6*	-0.4	-1.5*			
Breast (<i>in situ</i>)	2.2*	-	2.3*	-	-	-			

- Trends are based on rates age-adjusted to the 2000 US Std Population
- (19 age groups Census P25-1130).

SEER 18 areas (San Francisco, Connecticut, Detroit, Hawaii, Iowa, New Mexico, Seattle, Utah, Atlanta, San Jose-Monterey, Los Angeles, Alaska Native Registry, Rural Georgia, California excluding SF/SJM/LA, Kentucky, Louisiana, New Jersey and a Georgia excluding ATL/RG).

b US Mortality Files, National Center for Health Statistics, Centers for Disease Control and

Prevention. -

The APC is significantly different from zero (p<.05). Statistic could not be calculated. Trend based on less than 10 cases for at least one year within the time interval.

Table 1.10 - continued											
SEER	Incidence	and	U.S.	Mortality	Trends	by	Primary	Cancer	Site	and	Sex
				Blacks	, 2002-2	2011	L				

		Incidence	a	US Mortality ^b			
	Total	Males	Females	Total	Males	Females	
Site	APC	APC	APC	APC	APC	APC	
Female Genital System:	-0.1	-	0.1	-1.0*	_	-0.7*	
Cervix uteri	-3.4*	-	-3.3*	-2.6*	-	-2.3*	
Corpus uteri	2.2*	_	2.4*	-1.2	-	-0.9	
Uterus, NOS	3.8	_	4.1	2.2*		2.4*	
	-1.6*	-	-1.4*	-1.9*	_	-1.7*	
Ovary ^c		-			-		
Vagina	-2.0	-	-1.9	-2.8		-2.5	
Vulva	-0.8	-	-0.7	2.6	-	2.9	
Other female genital system	1.5	-	1.2	-0.3	-	-0.3	
Male Genital System:	-2.0*	-2.5*	_	-3.3*	-3.8*	_	
Prostate	-2.0*	-2.5*	-	-3.3*	-3.8*	_	
Testis	0.7	0.6	-	-5.5*	-5.7*	_	
Penis	-1.1	-1.2	_	0.2	0.1	_	
Other male genital system	-1.1	-1.2	_	-	-	_	
Urinary System:	1.3*	1.2	1.2*	-1.0*	-0.9*	-1.3*	
Urinary bladder	-0.5	0.1	-1.8*	-0.9	-0.7	-1.6*	
Kidney & renal pelvis	2.7*	2.1*	3.2*	-1.0*	-1.2*	-0.7	
Ureter	-2.3	_	-	-	_	_	
Other urinary system	2.1	-	-	-2.8	-	-4.6*	
Eye & Orbit	-1.2	_	_	_	_	_	
	0.0	o =	0 1	0 5			
Brain & Nervous System: ^d	-0.3	-0.5	0.1	-0.7	-1.1	-0.2	
Brain	0.2	-0.3	0.8	-	-	-	
Cranial nerves & other nervous system	-4.6	-3.8	-4.8	-	-	-	
Endocrine System:	5.5*	3.8*	6.0*	1.2	2.5	0.6	
Thyroid	5.9*	4.2*	6.3*	1.1	1.5	1.0	
Other endocrine & thymus	2.5	2.4	2.2	1.4	3.4	-0.2	
Lymphoma:	-0.2	-0.3	-0.1	-2.0*	-1.4	-2.8*	
	0.9	1.1	0.4	-2.1	-2.3	-1.7	
Hodgkin lymphoma							
Non-Hodgkin lymphoma	-0.3	-0.5	-0.2	-2.0*	-1.3	-2.9*	
Myeloma	0.4	-0.3	0.8	-1.6*	-1.3*	-2.0*	
Leukemia:	-1.0	-1.3	-0.8	-1.5*	-1.5*	-1.5*	
Lymphocytic:	-1.8*	-2.2	-1.6	-2.0*	-2.4*	-1.8*	
Acute lymphocytic	1.2	1.0	1.1	-2.0*	-2.9	-0.9	
Chronic lymphocytic	-2.7*	-3.2*	-2.4*	-2.0*	-2.0	-2.4*	
Other lymphocytic	-0.9	-	-	-2.0	-6.6	_	
Myeloid & Monocytic:	0.4	0.1	0.7	-1.2*	-1.0	-1.5*	
Acute myeloid	1.0	0.8	1.2	-0.3	0.1	-0.7	
Chronic myeloid	0.3	-0.7	1.5	-6.1*	-6.8*	-5.0*	
Acute monocytic	-5.0*	-	-	-	-	-	
Other myeloid & monocytic		-	-	-0.4	0.6	-3.7	
Other leukemia:	-5.2*	-4.1	-6.2	-1.3*	-1.3	-1.2	
Other acute leukemia	-7.3*	-	-	-3.8*	-3.4*	-4.2*	
Aleukemic, subleukemic & NOS	-4.0	-3.3	-3.6	-0.1	-0.2	0.2	
Kaposi Sarcoma ^e	-1.3	-1.4	_	_	_	_	
Mesothelioma ^e	0.5	-0.2	-	-	-	-	
Ill-defined & unanegified	-3.3*	-3.7*	-2.9*	-3.3*	-3.3*	-3.4*	
Ill-defined & unspecified	-3.3^	-3./^	-2.9^	-3.3^	-3.3^	-3.4^	

Trends are based on rates age-adjusted to the 2000 US Std Population

(19 age groups - Census P25-1130).

а SEER 18 areas (San Francisco, Connecticut, Detroit, Hawaii, Iowa, New Mexico, Seattle, Utah, Atlanta, San Jose-Monterey, Los Angeles, Alaska Native Registry, Rural Georgia, California excluding SF/SJM/LA, Kentucky, Louisiana, New Jersey and Georgia excluding ATL/RG).

b US Mortality Files, National Center for Health Statistics, Centers for Disease Control and Prevention.

с Ovary excludes borderline cases or histologies 8442, 8451, 8462, 8472, and 8473. d

e

Due to coding changes, Brain & Nervous System mortality are no longer shown separately. Trend not shown for mortality. Category did not exist in mortality coding until 1999. The APC is significantly different from zero (p<.05). Statistic could not be calculated. Trend based on less than 10 cases for at least one

_ year within the time interval.

Age Distribution (%) of Incidence Cases by Site, 2007-2011

All Races, Both Sexes

Age at Diagnosis

	Age at Diagnosis								All			
Site	<20	20-34	35-44	45-54	55-64	65-74	75-84	85+	All Ages	Cases		
All Sites	1.0	2.7	5.2	14.1	24.1	25.4	19.6	7.8	100.0%	2,001,481		
Oral Cavity & Pharynx:	0.6	2.1	5.6	19.8	29.5	21.9	14.4	6.1	100.0%	48,868		
Lip	0.2	1.1	5.4	15.0	18.9	23.9	22.6	12.8	100.0%	2,908		
Tonque	0.1	1.9	5.1	19.7	32.7	23.0	12.9	4.7	100.0%	14,368		
Salivary gland	2.1	б.4	7.7	14.2	19.8	19.7	19.7	10.5	100.0%	5,494		
Floor of mouth	0.0	0.3	3.1	20.0	32.7	25.2	14.1	4.6	100.0%	2,569		
Gum & other oral cavity	0.7	2.1	4.6	12.8	22.9	23.7	21.6	11.6	100.0%	6,582		
Nasopharynx	3.5	5.9	13.3	24.5	25.5	15.6	9.4	2.3	100.0%	2,866		
Tonsil	0.0	0.5	6.1	30.6	38.0	17.0	6.3	1.5	100.0%	8,475		
Oropharynx	0.1	0.2	3.9	20.1	34.0	24.7	12.7	4.2	100.0%	1,797		
Hypopharynx	0.0	0.2	1.4	16.7	31.8	28.1	17.2	4.5	100.0%	2,835		
Other oral cavity & pharynx	0.2	0.8	3.1	16.8	27.8	27.7	15.4	8.1	100.0%	974		
Digestive System:	0.2	1.0	3.6	13.5	22.9	24.4	23.1	11.3	100.0%	365,536		
Esophagus	0.0	0.3	2.0	11.7	27.1	27.7	22.6	8.5	100.0%	19,290		
Stomach	0.1	1.7	4.5	12.2	20.4	24.7	24.3	12.2	100.0%	32,063		
Small intestine	0.1	1.4	5.1	15.7	25.4	25.1	19.5	7.6	100.0%	9,263		
Colon & Rectum:	0.1	1.2	4.1	14.2	21.2	23.9	23.2	12.1	100.0%	188,874		
Colon	0.1	1.0	3.5	11.9	19.7	24.4	25.5	13.9	100.0%	133,670		
Rectum	0.1	1.6	5.6	19.9	24.8	22.5	17.7	7.7	100.0%	55,204		
Colon & Rectum (Male)	0.1	1.2	4.1	15.0	23.6	25.7	21.5	8.8	100.0%	97,463		
Colon & Rectum (Female)	0.1	1.2	4.1	13.4	18.6	22.0	25.1	15.6	100.0%	91,411		
Anus, anal canal & anorectum	0.0	1.1	7.3	25.2	27.6	18.7	13.9	6.2	100.0%	7,834		
Liver & intrahepatic bile duct	0.9	0.8	2.3	16.8	33.5	22.5	17.1	6.0	100.0%	35,695		
Gallbladder	0.0	0.3	2.6	8.6	19.4	25.7	28.5	14.8	100.0%	4,919		
Other biliary	0.1	0.5	2.0	8.4	19.4	25.4	28.5	14.8	100.0%	7,960		
Pancreas	0.1	0.4	2.1	9.4	21.5	26.3	26.8	13.4	100.0%	52,863		
Retroperitoneum	9.0	4.8	7.3	15.3	21.5	20.3	15.2	4.5	100.0%	1,677		
Peritoneum, omentum &	0.4	1.0	2.7	9.7	22.5	33.1	22.9	5.9	100.0%	2,698		
mesentery	0.4	1.0	2.1	5.1	21.1	JJ.T	22.9	5.9		2,000		
Other digestive system	0.3	0.8	2.8	10.8	20.5	23.0	28.1	13.7	100.0%	2,400		
Respiratory System:	0.1	0.4	1.4	9.0	21.9	31.3	27.1	8.7	100.0%	274,682		
Nose, nasal cavity &	1.8	4.0	7.3	14.9	23.7	21.4	17.9	8.9	100.0%	2,995		
middle ear												
Larynx	0.0	0.4	2.6	15.4	30.8	28.7	17.2	4.9	100.0%	14,687		
Lung & bronchus	0.0	0.3	1.3	8.6	21.4	31.7	27.9	8.9	100.0%	256,086		
Lung & bronchus (Male)	0.0	0.2	1.2	8.4	22.5	32.2	27.5	8.1	100.0%	135,278		
Lung & bronchus (Female)	0.0	0.3	1.4	8.9	20.2	31.0	28.3	9.9	100.0%	120,808		
Pleura	5.6	1.6	3.2	10.5	15.3	21.8	27.4	14.5	100.0%	124		
Trachea & other	17.7	20.0	9.1	11.3	13.0	12.8	10.9	5.2	100.0%	790		
respiratory organs												
Bones & joints	26.9	15.7	9.3	12.9	12.4	10.1	9.0	3.7	100.0%	3,972		
Soft tissue (including heart)	9.0	9.2	9.4	14.7	17.9	16.5	16.0	7.3	100.0%	14,395		
Skin (excl. basal & squamous):	0.6	6.2	9.2	16.7	21.4	20.1	17.6	8.1	100.0%	100,797		
Melanoma of the skin	0.5	6.2	9.4	17.3	22.0	20.2	16.8	7.4	100.0%	92,417		
Other non-epithelial skin	1.3	5.4	6.7	9.7	14.8	19.4	26.2	16.5	100.0%	8,380		
_												
Breast (Female)	0.0	1.8	9.3	22.0	25.5	21.3	14.4	5.7	100.0%	292,397		
Breast (Female -in situ)	0.0	0.7	10.4	28.5	27.1	20.3	10.8	2.2	100.0%	74,063		

Source: SEER 18 areas (San Francisco, Connecticut, Detroit, Hawaii, Iowa, New Mexico, Seattle, Utah, Atlanta, San Jose-Monterey, Los Angeles, Alaska Native Registry, Rural Georgia, California excluding SF/SJM/LA, Kentucky, Louisiana, New Jersey and Georgia excluding ATL/RG). Percents may not sum to 100 due to rounding.

Table 1.11 - continued

Age Distribution (%) of Incidence Cases by Site, 2007-2011

All Races, Both Sexes

Age at Diagnosis

	Age at Diagnosis									
Site	<20	20-34	35-44	45-54	55-64	65-74	75-84	85+	All Ages	Cases
Female Genital System:	0.4	4.1	8.9	19.0	27.9	20.6	13.3	5.8	100.0%	115,291
Cervix uteri	0.1	13.8	24.9	24.2	17.6	10.7	5.9	2.7	100.0%	17,223
Corpus uteri	0.0	1.6	5.6	18.4	34.1	23.5	12.5	4.3	100.0%	57,667
Uterus, NOS	0.1	1.7	5.4	15.8	25.4	20.4	16.7	14.6	100.0%	1,644
Ovary ^a	1.2	3.7	7.2	18.6	23.9	20.7	16.6	8.1	100.0%	29,010
Vagina	0.9	1.8	5.1	13.5	23.9	22.4	20.8	11.6	100.0%	1,759
Vulva	0.2	2.1	6.4	15.1	19.7	19.3	22.5	14.6	100.0%	5,699
Other female genital system	0.2	6.3	6.2	14.6	24.9	23.5	16.8	6.9	100.0%	2,289
Male Genital System:	0.3	1.9	1.5	9.9	31.6	34.9	16.3	3.7	100.0%	311,294
Prostate	0.0	0.0	0.6	9.7	32.7	36.3	16.8	3.8	100.0%	297,322
Testis	6.6	48.9	23.8	13.5	5.0	1.3	0.6	0.2	100.0%	11,801
Penis	0.1	1.7	5.8	11.7	22.9	25.0	22.3	10.5	100.0%	1,663
Other male genital system	3.1	2.8	6.9	13.2	21.5	18.7	21.5	12.4	100.0%	508
Urinary System:	0.6	1.0	3.4	11.0	21.7	26.7	25.0	10.5	100.0%	158,403
Urinary bladder	0.1	0.4	1.5	7.1	18.5	27.9	30.5	14.0	100.0%	86,940
Kidney & renal pelvis	1.2	1.8	6.0	16.4	26.2	25.2	17.4	5.7	100.0%	67,743
Ureter	0.0	0.1	0.6	4.1	14.3	28.7	36.1	16.0	100.0%	2,399
Other urinary system	0.0	0.7	1.7	7.1	17.4	24.6	31.5	17.0	100.0%	1,321
Eye & Orbit	12.9	3.2	6.4	14.9	19.9	20.4	15.7	6.5	100.0%	3,454
Brain & Nervous System:	13.1	8.9	8.4	14.8	19.7	17.0	13.3	4.8	100.0%	27,816
Brain	12.4	8.8	8.2	14.7	20.0	17.4	13.6	5.0	100.0%	26,100
Cranial nerves & other	23.7	10.4	11.8	15.9	16.0	10.7	8.6	2.9	100.0%	1,716
nervous system										
Endocrine System:	2.9	14.7	19.0	23.7	19.8	12.5	5.9	1.5	100.0%	59,131
Thyroid	1.8	15.1	19.6	24.2	19.9	12.3	5.7	1.4	100.0%	55,834
Other endocrine & thymus	21.7	7.6	8.2	14.5	19.0	16.6	9.8	2.7	100.0%	3,297
Lymphoma:	3.0	7.1	7.2	13.1	19.5	21.4	20.2	8.5	100.0%	96,350
Hodgkin lymphoma	12.8	31.0	14.5	12.7	10.9	9.1	6.6	2.3	100.0%	11,757
Non-Hodgkin lymphoma	1.6	3.8	6.2	13.2	20.7	23.1	22.1	9.3	100.0%	84,593
Myeloma	0.0	0.6	3.1	11.4	23.0	27.7	24.7	9.5	100.0%	26,346
Leukemia:	10.1	4.6	5.0	10.3	17.1	20.6	21.3	11.0	100.0%	55,604
Lymphocytic:	15.2	2.9	3.1	9.2	17.9	21.1	20.3	10.2	100.0%	27,868
Acute lymphocytic	58.8	10.2	5.5	7.0	7.3	5.5	4.2	1.5	100.0%	7,190
Chronic lymphocytic	0.0	0.2	1.6	9.0	21.5	27.2	26.7	13.7	100.0%	18,934
Other lymphocytic	0.2	2.0	10.4	19.3	23.6	18.3	17.8	8.3	100.0%	1,744
Myeloid & Monocytic:	4.9	6.7	7.2	12.0	16.6	20.5	22.0	10.2	100.0%	25,048
Acute myeloid	5.6	6.4	6.3	11.0	16.2	21.3	23.1	10.1	100.0%	16,173
Chronic myeloid	2.7	7.6	9.2	14.2	17.9	18.9	19.5	10.0	100.0%	7,113
Acute monocytic	9.6	5.7	8.4	12.6	17.9	17.0	19.6	9.1	100.0%	1,082
Other myeloid & monocytic	4.0	5.7	6.8	10.9	11.8	22.6	22.6	15.6	100.0%	680
Other leukemia:	4.5	3.5	4.0	7.6	12.5	16.1	26.1	25.7	100.0%	2,688
Other acute leukemia	8.0	4.3	3.5	7.8	10.7	14.8	24.5	26.4	100.0%	1,036
Aleukemic, subleukemic & NOS	2.3	3.0	4.4	7.4	13.6	16.9	27.1	25.3	100.0%	1,652
Kaposi Sarcoma	0.4	20.3	26.5	21.1	9.7	7.7	8.1	6.3	100.0%	2,258
Mesothelioma	0.1	0.7	1.9	6.0	16.1	27.2	33.8	14.3	100.0%	4,267
Ill-defined & unspecified	0.4	0.9	2.4	9.4	18.4	22.2	27.0	19.3	100.0%	38,242

Source: SEER 18 areas (San Francisco, Connecticut, Detroit, Hawaii, Iowa, New Mexico, Seattle, Utah, Atlanta, San Jose-Monterey, Los Angeles, Alaska Native Registry, Rural Georgia, California excluding SF/SJM/LA, Kentucky, Louisiana, New Jersey and Georgia excluding ATL/RG). Percents may not sum to 100 due to rounding.
^a Ovary excludes borderline cases or histologies 8442, 8451, 8462, 8472, and 8473.

	Table	e 1.12	
Median Age	of Cancer Paties	nts at Diagnosis	^a , 2007-2011
By	Primary Cancer	Site, Race and	Sex

	2	All Race	S		Whites		Blacks			
Site	Total		Females	Total		Females	Total		Females	
All Sites	66.0	66.0	65.0	66.0	67.0	65.0	62.0	63.0	62.0	
Oral Cavity & Pharynx:	62.0	61.0	65.0	62.0	61.0	66.0	58.0	59.0	58.0	
Lip	68.0	67.0	72.0	69.0	68.0	72.0	55.0	56.0	51.5	
Tongue	62.0	61.0	64.0	62.0	61.0	64.0	59.0	59.0	57.0	
Salivary gland	64.0	67.0	61.0	66.0	68.0	63.0	55.0	58.0	53.0	
Floor of mouth	63.0	62.0	66.0	63.0	62.0	67.0	60.0	60.5	60.0	
Gum & other oral cavity	67.0	65.0	71.0	69.0	65.0	73.0	60.0	60.0	60.0	
Nasopharynx	55.0	55.0	55.0	58.0	57.0	60.0	54.0	54.0	51.5	
Tonsil	58.0	57.0	60.0	58.0	57.0	60.0	57.0	58.0	57.0	
Oropharynx	62.0	61.0	64.0	63.0	62.0	66.0	59.5	60.0	58.5	
Hypopharynx	64.0	64.0	67.0	65.0	65.0	67.0	60.0	60.0	63.0	
Other oral cavity & pharynx	65.0	63.0	70.0	65.0	64.0	70.0	63.0	61.0	67.5	
Other ofat Cavity & pharynx	05.0	03.0	70.0	05.0	04.0	70.0	03.0	01.0	07.5	
Digestive System:	68.0	66.0	71.0	69.0	67.0	72.0	64.0	63.0	66.0	
Esophagus	67.0	66.0	72.0	68.0	67.0	73.0	63.0	63.0	65.0	
Stomach	69.0	68.0	71.0	69.0	68.0	72.0	67.0	66.0	69.0	
Small intestine	65.0	65.0	66.0	66.0	65.0	67.0	63.0	63.0	63.0	
Colon & Rectum:	68.0	67.0	70.0	70.0	68.0	72.0	64.0	63.0	65.0	
Colon	70.0	68.0	72.0	71.0	69.0	74.0	65.0	64.0	66.0	
Rectum	64.0	63.0	65.0	64.0	64.0	65.0	60.0	60.0	61.0	
Anus, anal canal & anorectum	60.0	58.0	61.0	61.0	60.0	61.0	54.0	52.0	58.0	
Liver & intrahepatic	63.0	61.0	69.0	63.0	62.0	69.0	60.0	59.0	62.0	
bile duct										
Gallbladder	72.0	72.0	72.0	73.0	73.0	73.0	67.0	69.0	66.0	
Other biliary	72.0	71.0	74.0	73.0	72.0	74.0	68.0	67.0	70.0	
Pancreas	71.0	69.0	73.0	71.0	69.0	74.0	67.0	64.0	69.0	
Retroperitoneum	61.0	61.0	60.5	61.0	61.0	62.0	58.0	58.0	59.0	
Peritoneum, omentum &	68.0	65.0	68.0	68.0	66.0	68.0	66.0	55.0	66.0	
mesentery	00.0	05.0	00.0	00.0	00.0	00.0	00.0	55.0	00.0	
Other digestive system	71.0	70.0	72.0	72.0	71.0	73.0	66.0	66.0	66.0	
other digestive system	/1.0	70.0	72.0	72.0	/1.0	/3.0	00.0	00.0	00.0	
Respiratory System:	70.0	70.0	71.0	71.0	70.0	71.0	66.0	65.0	67.0	
Nose, nasal cavity &	64.0	63.0	66.0	65.0	64.0	67.0	57.0	57.0	57.0	
middle ear										
Larynx	65.0	65.0	64.0	65.0	65.0	64.0	62.0	62.0	60.0	
Lung & bronchus	70.0	70.0	71.0	71.0	71.0	71.0	66.0	66.0	67.0	
Pleura	69.0	69.0	68.0	72.0	69.5	77.5	-	-	-	
Trachea & other	47.0	42.0	57.0	49.0	42.0	59.0	47.0	44.0	53.0	
respiratory organs										
Bones & joints	43.0	40.0	45.0	44.0	41.0	47.0	37.0	35.0	41.0	
5										
Soft tissue (including heart)	59.0	59.0	58.0	60.0	61.0	59.0	51.0	49.0	53.0	
Skin (excl. basal & squamous):	63.0	65.0	58.0	63.0	65.0	59.0	56.0	56.0	57.0	
Melanoma of the skin	62.0	65.0	58.0	62.0	65.0	58.0	62.0	62.0	62.5	
Other non-epithelial skin	71.0	72.0	69.0	73.0	73.0	71.0	50.0	50.5	49.0	
Breast	61.0	68.0	61.0	62.0	69.0	62.0	58.0	64.0	58.0	
Breast (<i>in situ</i>)	58.0	63.0	58.0	59.0	62.0	59.0	58.0	66.0	58.0	
DIEASL (III SILU)	50.0	03.0	0.00	59.0	02.0	0.66	50.0	00.0	50.0	

а SEER 18 areas (San Francisco, Connecticut, Detroit, Hawaii, Iowa, New Mexico, Seattle, Utah, Atlanta, San Jose-Monterey, Los Angeles, Alaska Native Registry, Rural Georgia, California excluding SF/SJM/LA, Kentucky, Louisiana, New Jersey and Georgia excluding ATL/RG). Statistic could not be calculated. Less than 16 cases were diagnosed during the time

-

interval.

Table 1.12 - continued Median Age of Cancer Patients at Diagnosis^a, 2007-2011 By Primary Cancer Site, Race and Sex

	All Races			Whites		Blacks			
Site	Total		Females	Total		Females	Total		Females
Female Genital System:	61.0	-	61.0	61.0	-	61.0	61.0	-	61.0
Cervix uteri	49.0	-	49.0	48.0	-	48.0	51.0	-	51.0
Corpus uteri	62.0	-	62.0	62.0	-	62.0	63.0	-	63.0
Uterus, NOS	65.0	-	65.0	66.0	-	66.0	63.0	-	63.0
Ovary ^b	63.0	-	63.0	63.0	-	63.0	62.0	-	62.0
Vagina	67.0	-	67.0	68.0	-	68.0	61.0	-	61.0
Vulva	68.0	_	68.0	69.0	_	69.0	58.0	-	58.0
Other female genital system	63.0	-	63.0	64.0	-	64.0	60.0	-	60.0
Male Genital System:	66.0	66.0	-	66.0	66.0	_	63.0	63.0	-
Prostate	66.0	66.0	-	66.0	66.0	-	63.0	63.0	-
Testis	33.0	33.0	_	33.0	33.0	_	35.0	35.0	_
Penis	68.0	68.0	_	68.0	68.0	_	62.5	62.5	_
Other male genital system	66.0	66.0	-	67.0	67.0	-	55.0	55.0	-
Urinary System:	69.0	69.0	69.0	70.0	70.0	70.0	65.0	64.0	66.0
Urinary bladder	73.0	72.0	74.0	73.0	73.0	74.0	70.0	68.5	73.0
Kidney & renal pelvis	64.0	63.0	65.0	64.0	64.0	66.0	61.0	60.0	63.0
Ureter	75.0	74.0	76.0	75.0	74.0	77.0	73.0	74.0	69.0
Other urinary system	74.0	75.0	72.0	75.0	75.0	74.0	67.0	67.0	63.0
Eye & Orbit	61.0	61.0	60.0	62.0	62.0	61.0	3.5	3.5	4.5
Brain & Nervous System:	57.0	57.0	58.0	58.0	57.0	59.0	50.0	50.0	49.0
Brain	58.0	57.0	59.0	59.0	58.0	60.0	50.0	50.0	50.0
Cranial nerves & other	47.0	45.0	49.0	48.0	46.0	49.0	45.0	41.0	46.5
nervous system	17.0	45.0	19.0	40.0	10.0	49.0	10.0	11.0	10.5
Endocrine System:	50.0	54.0	49.0	51.0	54.0	49.0	51.0	53.0	50.0
Thyroid	50.0	54.0	49.0	50.0	55.0	49.0	51.0	55.0	50.0
Other endocrine & thymus	53.0	51.0	56.0	54.0	52.0	57.0	51.0	48.0	53.0
Lymphoma:	65.0	63.0	66.0	66.0	64.0	67.0	55.0	53.0	57.0
Hodgkin lymphoma	39.0	40.0	37.0	40.0	41.0	37.0	37.0	38.0	35.0
Non-Hodgkin lymphoma	66.0	65.0	68.0	67.0	66.0	69.0	58.0	56.0	61.0
Myeloma	69.0	68.0	69.0	70.0	69.0	70.0	65.0	65.0	66.0
Leukemia:	66.0	65.0	67.0	67.0	66.0	68.0	60.0	59.0	62.0
Lymphocytic:	65.0	65.0	67.0	66.0	65.0	67.0	62.0	60.0	65.0
Acute lymphocytic	14.0	14.0	13.0	14.0	15.0	13.0	13.0	13.0	13.0
Chronic lymphocytic	71.0	70.0	73.0	72.0	70.0	74.0	69.0	67.0	71.0
Other lymphocytic	62.0	61.0	67.0	62.0	61.0	67.0	65.0	63.0	73.0
Myeloid & Monocytic:	66.0	66.0	66.0	67.0	67.0	67.0	58.0	58.0	59.0
Acute myeloid	67.0	67.0	66.0	68.0	68.0	68.0	60.0	59.0	61.0
Chronic myeloid	67.0	63.0	65.0	68.0 65.0	68.0 64.0	68.0 67.0	56.0	59.0 56.0	57.0
-									
Acute monocytic	62.0	64.0	60.0	63.0	65.0	61.0	52.0	56.0	49.5
Other myeloid & monocytic	70.0	70.0	69.0	71.0	72.0	69.5	62.5	59.5	63.5
Other leukemia:	76.0	73.0	78.0	77.0	75.0	79.0	67.0	63.0	71.5
Other acute leukemia	75.0	72.0	78.0	77.0	74.0	80.0	63.5	58.0	76.0
Aleukemic, subleukemic & NOS	76.0	73.0	78.0	77.0	75.0	79.0	68.0	66.0	70.0
Kaposi Sarcoma	46.0	44.0	75.0	48.0	46.5	80.0	39.0	39.0	46.0
Mesothelioma	74.0	75.0	71.0	74.0	75.0	71.0	70.0	71.5	63.0
Ill-defined & unspecified	73.0	70.0	76.0	74.0	71.0	77.0	67.0	64.0	69.0

a SEER 18 areas (San Francisco, Connecticut, Detroit, Hawaii, Iowa, New Mexico, Seattle, Utah, Atlanta, San Jose-Monterey, Los Angeles, Alaska Native Registry, Rural Georgia, California excluding SF/SJM/LA, Kentucky, Louisiana, New Jersey and Georgia excluding ATL/RG). Ovary excludes borderline cases or histologies 8442, 8451, 8462, 8472, and 8473. Statistic could not be calculated. Less than 16 cases were diagnosed during the time b

interval.

Age Distribution (%) of Deaths by Site, 2007-2011

All Races, Both Sexes

Age at Death

			Age a	t Death						
Site	<20	20-34	35-44	45-54	55-64	65-74	75-84	85+	All Ages	Deaths
All Sites	0.3	0.8	2.2	8.8	18.8	25.0	27.9	16.1	100.0%	2,847,364
Oral Cavity & Pharynx:	0.1	0.8	2.7	13.7	25.7	24.0	20.8	12.3	100.0%	41,139
Lip	0.3	0.9	2.0	8.2	12.2	17.5	28.0	30.9	100.0%	343
Tongue	0.1	1.0	3.2	14.4	26.8	23.7	19.8	10.9	100.0%	10,202
Salivary gland	0.1	0.9	2.6	9.1	16.8	21.2	27.9	21.4	100.0%	3,902
Floor of mouth	0.0	0.0	2.7	11.3	32.2	28.1	17.9	7.8	100.0%	487
Gum & other oral cavity	0.1	0.5	1.9	8.9	18.8	22.0	25.5	22.4	100.0%	5,817
Nasopharynx	0.8	3.3	6.0	19.2	26.5	20.3	16.5	7.5	100.0%	3,315
Tonsil	0.0	0.2	2.8	19.9	33.9	23.9	14.2	5.1	100.0%	3,785
Oropharynx	0.0	0.3	2.3	15.2	29.3	24.8	18.7	9.3	100.0%	3,842
Hypopharynx	0.0	0.3	1.2	12.9	28.8	29.4	21.2	6.4	100.0%	1,543
Other oral cavity & pharynx	0.0	0.2	1.7	13.0	27.4	27.2	20.9	9.6	100.0%	7,903
Digestive System:	0.1	0.5	2.1	9.6	20.3	23.9	26.9	16.7	100.0%	701,293
Esophagus	0.0	0.2	1.7	10.5	25.0	27.4	24.4	10.7	100.0%	70,150
Stomach	0.0	1.3	3.8	10.3	17.5	22.6	26.8	17.6	100.0%	56,349
Small intestine	0.0	0.8	2.6	9.5	20.7	23.3	27.0	16.1	100.0%	5,944
Colon & Rectum:	0.0	0.6	2.5	9.1	17.6	21.9	27.3	20.9	100.0%	261,752
Colon & Rectum (Male)	0.0	0.7	2.7	10.0	20.2	24.6	26.6	15.1	100.0%	134,620
Colon & Rectum (Female)	0.0	0.6	2.3	8.2	14.7	19.1	28.0	27.1	100.0%	127,132
Anus, anal canal & anorectum	0.0	0.5	5.2	19.9	25.1	20.0	17.8	11.4	100.0%	3,856
Liver & intrahepatic bile duct	0.3	0.6	1.7	13.1	27.6	23.0	23.3	10.4	100.0%	96,623
Gallbladder	0.0	0.2	1.4	6.9	17.8	26.4	29.4	17.8	100.0%	10,139
Other biliary	0.0	0.3	1.5	6.5	15.0	22.2	31.9	22.6	100.0%	7,226
Pancreas	0.0	0.2	1.4	7.9	19.7	26.1	28.9	15.9	100.0%	179,211
Retroperitoneum	0.4	1.4	2.4	10.2	20.1	24.4	27.0	14.1	100.0%	1,037
Peritoneum, omentum & mesentery	0.1	0.6	1.8	7.3	19.3	29.0	29.6	12.4	100.0%	4,146
Other digestive system	0.0	0.4	2.1	7.4	16.5	21.7	30.2	21.8	100.0%	4,860
Respiratory System:	0.0	0.1	1.0	7.8	19.8	30.5	29.6	11.1	100.0%	813,550
Nose, nasal cavity & middle ear	0.2	1.8	5.7	12.9	19.4	21.1	23.4	15.4	100.0%	2,432
Larynx	0.0	0.1	1.3	11.4	26.2	28.6	23.2	9.3	100.0%	18,447
Lung & bronchus	0.0	0.1	1.0	7.7	19.7	30.6	29.8	11.2	100.0%	790,557
Lung & bronchus (Male)	0.0	0.1	0.9	7.7	21.0	31.4	29.2	9.7	100.0%	438,998
Lung & bronchus (Female)	0.0	0.1	1.1	7.8	18.1	29.5	30.5	13.0	100.0%	351,559
Pleura	0.3	0.5	1.2	2.9	14.4	26.4	38.5	15.7	100.0%	1,018
Trachea & other	0.9	4.9	4.0	12.7	17.4	21.2	24.3	14.6	100.0%	1,096
respiratory organs										
Bones & joints	12.6	15.1	6.1	10.6	12.5	14.2	16.7	12.2	100.0%	6,904
Soft tissue (including heart)	3.6	6.2	6.5	13.2	19.4	19.3	20.3	11.5	100.0%	21,139
Skin (excl. basal & squamous):	0.1	1.8	4.2	11.4	19.4	21.2	24.9	17.2	100.0%	59,224
Melanoma of the skin	0.1	2.3	5.1	12.7	20.5	21.7	24.0	13.6	100.0%	44,565
Other non-epithelial skin	0.0	0.3	1.4	7.4	16.0	19.5	27.4	28.0	100.0%	14,659
Breast (Female)	0.0	0.9	5.2	14.5	21.7	20.6	21.0	16.2	100.0%	203,790

Source: US Mortality Files, National Center for Health Statistics, Centers for Disease Control and Prevention. Percents may not sum to 100 due to rounding.

Table 1.13 - continued

Age Distribution (%) of Deaths by Site, 2007-2011

All Races, Both Sexes

Age at Death

			Age a	t Death						
Site	<20	20-34	35-44	45-54	55-64	65-74	75-84	85+	All Ages	Deaths
Female Genital System:	0.0	1.2	4.0	11.6	21.3	24.1	23.6	14.1	100.0%	141,166
Cervix uteri	0.0	4.8	14.5	24.0	22.6	15.8	11.8	6.4	100.0%	19,969
Corpus uteri	0.0	0.3	1.8	7.5	23.2	28.4	24.6	14.1	100.0%	17,504
Uterus, NOS	0.0	0.4	2.2	8.7	21.8	26.4	24.4	16.2	100.0%	22,383
Ovary	0.1	0.7	2.5	10.5	21.0	25.2	25.8	14.2	100.0%	72,337
Vagina	0.0	0.8	2.7	6.6	16.0	19.7	28.2	26.0	100.0%	2,042
Vulva	0.0	0.5	2.3	7.5	13.1	16.7	29.7	30.1	100.0%	4,696
Other female genital system	0.1	1.3	2.9	9.9	21.5	25.9	24.2	14.3	100.0%	2,235
Male Genital System:	0.0	0.5	0.4	1.9	8.7	19.9	36.2	32.4	100.0%	145,473
Prostate	0.0	0.0	0.1	1.6	8.5	20.1	36.8	33.0	100.0%	142,182
Testis	1.8	34.8	20.4	18.4	11.2	5.0	5.1	3.3	100.0%	1,839
Penis	0.1	0.8	4.7	10.6	20.6	23.7	22.6	16.8	100.0%	1,223
Other male genital system	0.4	0.9	3.5	9.6	14.0	22.3	26.6	22.7	100.0%	229
Urinary System:	0.2	0.3	1.2	6.6	15.9	23.0	30.7	22.1	100.0%	140,985
Urinary bladder	0.0	0.1	0.6	4.0	11.6	20.8	34.5	28.4	100.0%	71,824
Kidney & renal pelvis	0.4	0.6	1.8	9.6	21.1	25.3	26.2	15.0	100.0%	65,371
Ureter	0.1	0.2	0.4	3.2	9.2	24.1	37.5	25.3	100.0%	1,753
Other urinary system	0.0	0.2	1.3	5.9	12.0	21.6	35.5	23.4	100.0%	2,037
Eye & Orbit	2.0	1.8	4.7	11.2	19.8	21.4	23.6	15.5	100.0%	1,352
Brain & Nervous System:	3.8	3.5	5.9	14.2	23.4	23.1	18.9	7.2	100.0%	69,789
Endocrine System:	6.3	2.5	3.9	9.6	18.0	22.1	23.8	13.8	100.0%	13,007
Thyroid	0.1	0.8	2.1	7.8	17.7	24.4	28.9	18.1	100.0%	8,351
Other endocrine & thymus	17.4	5.6	7.1	12.8	18.7	17.8	14.5	6.0	100.0%	4,656
Lymphoma:	0.4	2.0	2.7	6.7	14.4	21.9	31.7	20.2	100.0%	107,987
Hodgkin lymphoma	1.4	12.3	10.2	11.1	14.9	17.8	21.5	10.8	100.0%	6,091
Non-Hodgkin lymphoma	0.4	1.4	2.3	6.4	14.4	22.1	32.3	20.7	100.0%	101,896
Myeloma	0.0	0.1	1.0	5.8	16.4	26.3	33.1	17.3	100.0%	54,601
Leukemia:	2.5	3.0	2.8	6.3	13.1	21.7	30.5	20.1	100.0%	112,914
Lymphocytic:	4.0	3.7	2.2	4.8	11.4	19.2	29.2	25.4	100.0%	31,589
Acute lymphocytic	17.4	15.4	8.3	11.1	14.0	14.2	13.0	6.6	100.0%	7,133
Chronic lymphocytic	0.0	0.1	0.4	2.8	10.7	20.7	34.2	31.2	100.0%	22,517
Other lymphocytic	1.6	1.6	1.4	5.5	11.1	20.0	31.1	27.7	100.0%	1,939
Myeloid & Monocytic:	1.9	2.9	3.4	7.8	15.2	24.1	30.0	14.7	100.0%	54,525
Acute myeloid	2.1	3.0	3.4	7.9	15.8	24.9	29.8	13.2	100.0%	45,394
Chronic myeloid	0.5	3.5	4.8	8.7	12.2	17.6	28.7	24.1	100.0%	5,097
Acute monocytic	1.2	1.6	1.8	5.7	10.9	24.8	32.4	21.4	100.0%	487
Other myeloid & monocytic	1.5	1.2	1.8	4.9	12.3	23.1	34.7	20.4	100.0%	3,547
Other leukemia:	2.2	2.2	2.3	5.0	10.8	20.0	33.0	24.6	100.0%	26,800
Other acute leukemia	1.2	2.4	2.4	5.0	11.1	20.9	33.5	23.5	100.0%	10,055
Aleukemic, subleukemic & NOS	2.7	2.1	2.2	5.0	10.7	19.4	32.6	25.3	100.0%	16,745
Ill-defined & unspecified	0.2	0.7	1.9	8.0	17.7	23.3	28.8	19.4	100.0%	210,961

Source: US Mortality Files, National Center for Health Statistics, Centers for Disease Control and Prevention. Percents may not sum to 100 due to rounding.

Table 1.14											
Median	Age	of	Cano	cer	Pat	ients	at	Dea	ath ^a ,	2007-	-2011
	By I	Prin	nary	Car	ncer	Site,	Ra	lce	and	Sex	

	All Race	s		Whites		Blacks			
Site	Total		Females	Total		Females	Total		Females
All Sites	72.0	72.0	73.0	73.0	72.0	74.0	67.0	67.0	68.0
Oral Cavity & Pharynx:	67.0	65.0	73.0	68.0	66.0	74.0	62.0	61.0	63.0
Lip	78.0	76.0	85.0	78.0	76.0	85.0	_	-	-
Tongue	66.0	64.0	72.0	67.0	65.0	72.0	61.0	61.0	60.5
Salivary gland	74.0	73.0	76.0	75.0	74.0	78.0	62.0	61.5	65.5
Floor of mouth	66.0	64.0	71.0	67.0	64.0	72.0	61.0	61.0	-
Gum & other oral cavity	74.0	68.0	80.0	75.0	69.0	81.0	64.0	62.0	68.0
Nasopharynx	62.0	61.0	66.0	65.0	63.0	71.0	59.0	58.0	59.0
Tonsil	62.0	61.0	67.0	63.0	61.0	68.0	61.0	61.0	61.0
Oropharynx	65.0	64.0	72.0	67.0	65.0	73.0	61.0	61.0	61.0
Hypopharynx	67.0	66.0	69.5	67.0	67.0	70.0	62.0	61.0	63.5
Other oral cavity & pharynx	67.0	66.0	71.0	68.0	67.0	72.0	63.0	63.0	66.0
Digestive System:	72.0	69.0	75.0	73.0	70.0	76.0	67.0	65.0	70.0
Esophagus	69.0	68.0	74.0	70.0	68.0	75.0	64.0	64.0	66.0
Stomach	72.0	71.0	75.0	73.0	71.0	76.0	70.0	68.0	73.0
Small intestine	72.0	70.0	74.0	73.0	71.0	75.0	65.0	63.0	67.0
Colon & Rectum	74.0	71.0	77.0	75.0	72.0	78.0	68.0	66.0	71.0
Anus, anal canal & anorectum	64.0	62.0	66.0	65.0	63.0	66.0	57.0	54.0	61.0
Liver & intrahepatic	67.0	64.0	74.0	69.0	66.0	74.0	61.0	60.0	67.0
bile duct									
Gallbladder	74.0	73.0	74.0	74.0	73.0	75.0	69.0	70.0	68.0
Other biliary	76.0	74.0	78.0	77.0	75.0	78.0	71.0	70.0	73.0
Pancreas	73.0	70.0	75.0	73.0	71.0	76.0	69.0	66.0	72.0
Retroperitoneum	71.0	70.0	71.0	72.0	71.0	73.0	64.0	64.0	65.5
Peritoneum, omentum &	72.0	68.0	72.0	72.0	68.5	73.0	68.0	64.0	69.0
mesentery									
Other digestive system	75.0	73.0	78.0	76.0	74.0	79.0	69.0	65.0	74.0
Respiratory System:	72.0	71.0	72.0	72.0	71.0	73.0	67.0	67.0	69.0
Nose, nasal cavity &	69.0	66.0	74.0	71.0	67.0	76.0	63.0	62.0	66.0
middle ear									
Larynx	68.0	68.0	70.0	69.0	68.0	71.0	65.0	65.0	65.0
Lung & bronchus	72.0	71.0	72.0	72.0	72.0	73.0	68.0	67.0	69.0
Pleura	75.5	76.0	75.0	76.0	76.0	75.0	70.0	68.0	-
Trachea & other	69.0	66.0	74.0	70.0	66.0	74.0	60.0	61.0	57.5
respiratory organs									
Bones & joints	59.0	57.0	63.0	61.0	58.0	65.0	52.0	50.0	56.0
-									
Soft tissue (including heart)	65.0	65.0	65.0	66.0	66.0	67.0	57.0	55.0	59.0
Skin (excl. basal & squamous):	71.0	70.0	72.0	71.0	71.0	72.0	63.0	61.0	69.0
Melanoma of the skin	69.0	69.0	69.0	69.0	69.0	69.0	68.0	65.0	70.0
Other non-epithelial skin	77.0	75.0	81.0	78.0	76.0	82.0	60.0	59.0	65.0
Breast	68.0	71.0	68.0	69.0	72.0	69.0	62.0	65.0	61.0

US Mortality Files, National Center for Health Statistics, Centers for Disease Control and а

_

Prevention. Statistic could not be calculated. Less than 16 deaths occurred during the time interval.

			Tabl	le 1	1.14	- con	itin	ued	1		
Median	Age	e of	Cano	cer	Pat	ients	at	Dea	ath ^a ,	2007	-2011
	Ву	Prin	nary	Cai	ncer	Site,	Ra	ce	and	Sex	

	1	All Race	S		Whites			Blacks		
Site	Total		Females	Total		Females	Total	Males		
Female Genital System:	69.0	-	69.0	70.0	-	70.0	66.0	-	66.0	
Cervix uteri	57.0	-	57.0	58.0	-	58.0	57.0	-	57.0	
Corpus uteri	70.0	-	70.0	71.0	-	71.0	68.0	-	68.0	
Uterus, NOS	71.0	-	71.0	72.0	-	72.0	68.0	-	68.0	
Ovary	71.0	-	71.0	71.0	-	71.0	67.0	-	67.0	
Vagina	76.0	-	76.0	78.0	-	78.0	70.0	-	70.0	
Vulva	78.0	-	78.0	79.0	-	79.0	69.0	-	69.0	
Other female genital system	70.0	-	70.0	71.0	-	71.0	66.0	-	66.0	
Male Genital System:	80.0	80.0	_	81.0	81.0	-	77.0	77.0	-	
Prostate	80.0	80.0	-	81.0	81.0	-	77.0	77.0	-	
Testis	41.0	41.0	-	41.0	41.0	-	42.0	42.0	-	
Penis	70.0	70.0	-	70.0	70.0	-	69.0	69.0	-	
Other male genital system	74.0	74.0	-	75.0	75.0	-	69.0	69.0	-	
Urinary System:	76.0	74.0	78.0	76.0	75.0	78.0	70.0	68.0	74.0	
Urinary bladder	79.0	78.0	80.0	79.0	78.0	81.0	75.0	73.0	77.0	
Kidney & renal pelvis	71.0	69.0	74.0	72.0	70.0	75.0	67.0	64.0	70.0	
Ureter	78.0	77.0	79.0	78.0	77.5	80.0	75.0	74.0	76.0	
Other urinary system	77.0	77.0	78.0	78.0	77.0	79.0	69.0	69.5	68.0	
Eye & Orbit	69.0	67.0	72.0	70.0	68.0	72.0	55.0	54.5	62.0	
Brain & Nervous System	64.0	63.0	66.0	65.0	63.0	66.0	59.0	58.0	61.0	
Endocrine System:	69.0	66.0	72.0	70.0	67.0	73.0	63.0	60.0	65.0	
Thyroid	73.0	71.0	76.0	70.0	71.0	76.0	69.0	67.0	71.0	
Other endocrine & thymus	58.0	57.0	59.0	59.0	58.0	60.0	53.0	52.0	53.0	
other endocrine & thymus	50.0	57.0	59.0	55.0	50.0	00.0	55.0	52.0	55.0	
Lymphoma:	75.0	73.0	78.0	76.0	74.0	78.0	64.0	62.0	68.0	
Hodgkin lymphoma	65.0	62.0	68.0	66.0	64.0	70.0	51.0	49.0	53.0	
Non-Hodgkin lymphoma	76.0	73.0	78.0	76.0	74.0	78.0	65.0	62.0	69.0	
Myeloma	75.0	74.0	76.0	75.0	74.0	77.0	71.0	69.0	72.0	
Leukemia:	75.0	74.0	76.0	75.0	74.0	77.0	68.0	66.0	70.0	
Lymphocytic:	76.0	74.0	79.0	77.0	75.0	80.0	69.5	67.0	74.0	
Acute lymphocytic	53.0	50.0	56.0	54.0	52.0	57.0	43.0	37.5	49.0	
Chronic lymphocytic	80.0	78.0	83.0	80.0	78.0	83.0	74.0	71.0	77.0	
Other lymphocytic	78.0	76.0	81.0	79.0	76.0	82.0	71.0	67.0	78.0	
Myeloid & Monocytic:	73.0	72.0	73.0	73.0	73.0	74.0	65.0	64.0	66.0	
Acute myeloid	72.0	72.0	72.0	73.0	72.0	73.0	66.0	65.0	67.0	
Chronic myeloid	76.0	73.0	79.0	77.0	75.0	80.0	61.0	58.0	62.0	
Acute monocytic	76.0	76.0	76.0	77.0	77.0	77.0	70.0	_	-	
Other myeloid & monocytic	76.0	75.0	78.0	77.0	76.0	78.0	70.0	67.0	76.0	
Other leukemia:	77.0	76.0	79.0	78.0	77.0	80.0	70.0	69.0	73.0	
Other acute leukemia	77.0	76.0	78.0	77.0	76.0	79.0	71.0	69.0	73.0	
Aleukemic, subleukemic & NOS	77.0	76.0	79.0	78.0	77.0	80.0	70.0	68.0	73.0	
Ill-defined & unspecified	74.0	72.0	76.0	75.0	73.0	77.0	68.0	66.0	70.0	

US Mortality Files, National Center for Health Statistics, Centers for Disease Control and а

Prevention. Statistic could not be calculated. Less than 16 deaths occurred during the time interval. _

Lifetime Risk (Percent) of Being Diagnosed with Cancer by Site and Race/Ethnicity

Both Sexes, 18 SEER Areas, 2009-2011

	All Races	Whites	Blacks
Site	Percent (95% C.I.)	Percent (95% C.I.)	Percent (95% C.I.)
All Sites	40.37 (40.28, 40.46)	40.55 (40.46, 40.65)	37.63 (37.37, 37.89)
Invasive and In Situ	42.73 (42.64, 42.82)	42.91 (42.81, 43.01)	38.95 (38.69, 39.22)
Oral Cavity and Pharynx	1.10 (1.09, 1.11)	1.15 (1.13, 1.16)	0.80 (0.76, 0.83)
Esophagus	0.50 (0.50, 0.51)	0.53 (0.52, 0.54)	0.45 (0.42, 0.48)
Stomach	0.87 (0.85, 0.88)	0.76 (0.75, 0.78)	1.10 (1.06, 1.15)
Colon and Rectum	4.66 (4.63, 4.69)	4.56 (4.53, 4.59)	4.85 (4.75, 4.94)
Invasive and In Situ	4.83 (4.80, 4.86)	4.72 (4.68, 4.75)	5.07 (4.97, 5.17)
Liver and Intrahepatic Bile Duct	0.89 (0.88, 0.90)	0.79 (0.77, 0.80)	0.91 (0.88, 0.95)
Pancreas	1.50 (1.49, 1.52)	1.49 (1.47, 1.51)	1.59 (1.54, 1.65)
Larynx	0.35 (0.35, 0.36)	0.36 (0.35, 0.37)	0.44 (0.41, 0.47)
Invasive and In Situ	0.38 (0.37, 0.39)	0.38 (0.38, 0.39)	0.47 (0.44, 0.50)
Lung and Bronchus	6.75 (6.71, 6.79)	6.92 (6.88, 6.96)	6.47 (6.36, 6.58)
Melanoma of the Skin	2.05 (2.03, 2.07)	2.39 (2.36, 2.41)	0.09 (0.08, 0.11)
Invasive and In Situ	3.44 (3.42, 3.47)	3.90 (3.87, 3.93)	0.12 (0.11, 0.14)
Breast	6.38 (6.35, 6.42)	6.50 (6.47, 6.54)	5.88 (5.79, 5.98)
Invasive and In Situ	7.62 (7.58, 7.65)	7.73 (7.69, 7.77)	7.05 (6.94, 7.15)
Urinary Bladder (Invasive and In Situ)	2.40 (2.38, 2.43)	2.60 (2.58, 2.63)	1.25 (1.20, 1.30)
Kidney and Renal Pelvis	1.60 (1.59, 1.62)	1.65 (1.63, 1.67)	1.60 (1.55, 1.65)
Brain and Other Nervous System	0.62 (0.61, 0.63)	0.68 (0.67, 0.69)	0.35 (0.32, 0.37)
Thyroid	1.13 (1.11, 1.14)	1.18 (1.17, 1.20)	0.64 (0.62, 0.67)
Hodgkin Lymphoma	0.22 (0.22, 0.23)	0.24 (0.23, 0.24)	0.19 (0.18, 0.21)
Non-Hodgkin Lymphoma	2.12 (2.11, 2.14)	2.22 (2.20, 2.25)	1.28 (1.24, 1.33)
Myeloma	0.72 (0.71, 0.73)	0.66 (0.65, 0.67)	1.21 (1.17, 1.26)
Leukemia	1.43 (1.42, 1.45)	1.50 (1.48, 1.51)	0.96 (0.92, 1.01)
Acute Lymphocytic Leukemia	0.13 (0.13, 0.14)	0.14 (0.14, 0.15)	0.07 (0.07, 0.08)
Chronic Lymphocytic Leukemia	0.54 (0.53, 0.55)	0.57 (0.56, 0.58)	0.32 (0.29, 0.35)
Acute Myeloid Leukemia	0.43 (0.42, 0.44)	0.45 (0.44, 0.46)	0.31 (0.29, 0.34)
Chronic Myeloid Leukemia	0.18 (0.18, 0.19)	0.18 (0.18, 0.19)	0.14 (0.13, 0.16)
Kaposi Sarcoma	0.05 (0.04, 0.05)	0.04 (0.04, 0.04)	0.08 (0.07, 0.09)
Mesothelioma	0.12 (0.12, 0.13)	0.14 (0.13, 0.15)	0.06 (0.05, 0.07)
			,,

Devcan Version 6.8.0, August 2014, National Cancer Institute (<u>http://surveillance.cancer.gov/devcan</u>/). Source: SEER 18 areas (San Francisco, Connecticut, Detroit, Hawaii, Iowa, New Mexico, Seattle, Utah, Atlanta, San Jose-Monterey, Los Angeles, Alaska Native Registry, Rural Georgia, California excluding SF/SJM/LA, Kentucky, Louisiana, New Jersey, and Georgia excluding ATL/RG).

Note: Invasive cancer only unless specified otherwise.

A percent or confidence interval value of 0.00 represents a value that is below 0.005.

Table 1.15 - continued

Lifetime Risk (Percent) of Being Diagnosed with Cancer by Site and Race/Ethnicity

Both Sexes, 18 SEER Areas, 2009-2011

	Asian/Pacific Islanders	American Indian/ Alaska Natives ^a	Hispanics ^b		
Site	Percent (95% C.I.)	Percent (95% C.I.)	Percent (95% C.I.)		
All Sites	34.75 (34.40, 35.11)	29.15 (27.99, 30.41)	37.15 (36.83, 37.47)		
Invasive and In Situ	36.13 (35.77, 36.48)	29.95 (28.77, 31.21)	38.36 (38.04, 38.69)		
Oral Cavity and Pharynx	0.88 (0.83, 0.94)	0.80 (0.65, 1.08)	0.81 (0.76, 0.86)		
Esophagus	0.34 (0.30, 0.39)	0.32 (0.21, 0.56)	0.38 (0.34, 0.41)		
Stomach	1.78 (1.69, 1.87)	1.07 (0.86, 1.40)	1.49 (1.43, 1.57)		
Colon and Rectum	4.99 (4.85, 5.14)	4.23 (3.79, 4.77)	4.56 (4.44, 4.68)		
Invasive and In Situ	5.17 (5.03, 5.31)	4.33 (3.89, 4.88)	4.70 (4.58, 4.82)		
Liver and Intrahepatic Bile Duct	1.92 (1.84, 2.01)	1.47 (1.24, 1.81)	1.59 (1.53, 1.66)		
Pancreas	1.58 (1.50, 1.67)	1.23 (0.99, 1.58)	1.64 (1.57, 1.72)		
Larynx	0.18 (0.16, 0.21)	0.25 (0.16, 0.48)	0.30 (0.27, 0.33)		
Invasive and In Situ	0.19 (0.17, 0.22)	0.26 (0.17, 0.49)	0.31 (0.29, 0.35)		
Lung and Bronchus	5.55 (5.41, 5.70)	4.46 (4.00, 5.01)	4.28 (4.17, 4.40)		
Melanoma of the Skin	0.18 (0.15, 0.22)	0.38 (0.28, 0.61)	0.49 (0.46, 0.54)		
Invasive and In Situ	0.24 (0.21, 0.28)	0.59 (0.45, 0.85)	0.78 (0.73, 0.83)		
Breast	5.38 (5.27, 5.50)	4.06 (3.68, 4.54)	5.12 (5.01, 5.23)		
Invasive and In Situ	6.67 (6.55, 6.80)	4.60 (4.20, 5.09)	5.99 (5.88, 6.10)		
Urinary Bladder (Invasive and In Situ)	1.40 (1.33, 1.49)	0.93 (0.71, 1.28)	1.64 (1.56, 1.72)		
Kidney and Renal Pelvis	1.03 (0.97, 1.09)	1.96 (1.71, 2.31)	1.75 (1.69, 1.82)		
Brain and Other Nervous System	0.39 (0.36, 0.43)	0.30 (0.18, 0.57)	0.54 (0.51, 0.58)		
Thyroid	1.21 (1.16, 1.27)	0.75 (0.59, 1.04)	1.04 (1.01, 1.09)		
Hodgkin Lymphoma	0.12 (0.10, 0.14)	0.09 (0.05, 0.30)	0.21 (0.20, 0.24)		
Non-Hodgkin Lymphoma	1.83 (1.75, 1.92)	1.33 (1.10, 1.68)	2.23 (2.16, 2.31)		
Myeloma	0.54 (0.50, 0.59)	0.44 (0.31, 0.71)	0.76 (0.72, 0.81)		
Leukemia	0.96 (0.90, 1.02)	0.78 (0.60, 1.07)	1.21 (1.15, 1.27)		
Acute Lymphocytic Leukemia	0.10 (0.09, 0.12)	0.09 (0.06, 0.30)	0.19 (0.18, 0.21)		
Chronic Lymphocytic Leukemia	0.17 (0.14, 0.20)	0.18 (0.10, 0.43)	0.30 (0.27, 0.33)		
Acute Myeloid Leukemia	0.43 (0.39, 0.47)	0.26 (0.16, 0.51)	0.41 (0.38, 0.45)		
Chronic Myeloid Leukemia	0.15 (0.13, 0.18)	0.10 (0.06, 0.31)	0.18 (0.15, 0.21)		
Kaposi Sarcoma	0.02 (0.01, 0.03)	0.02 (0.00, 0.24)	0.09 (0.07, 0.11)		
Mesothelioma	0.06 (0.04, 0.08)	0.06 (0.01, 0.29)	0.11 (0.10, 0.14)		

Devcan Version 6.8.0, August 2014, National Cancer Institute (<u>http://surveillance.cancer.gov/devcan</u>/).

Source: SEER 18 areas (San Francisco, Connecticut, Detroit, Hawaii, Iowa, New Mexico, Seattle, Utah, Atlanta, San Jose-Monterey, Los Angeles, Alaska Native Registry, Rural Georgia, California excluding SF/SJM/LA, Kentucky, Louisiana, New Jersey, and Georgia excluding ATL/RG).

Note: Invasive cancer only unless specified otherwise.

Underlying incidence data for American Indian/Alaska Native are based on the CHSDA(Contract Health Service Delivery Area) counties.
 Hisparic is not mutually evaluative from whites blacks Asian Bacific Islanders and American Indians/Alaska Natives

Hispanic is not mutually exclusive from whites, blacks, Asian Pacific Islanders, and American Indians/Alaska Natives. Underlying incidence data for Hispanics are based on NHIA and exclude cases from the Alaska Native Registry. A percent or confidence interval value of 0.00 represents a value that is below 0.005.

Lifetime Risk (Percent) of Being Diagnosed with Cancer by Site and Race/Ethnicity

Males, 18 SEER Areas, 2009-2011

	All Races	Whites	Blacks
Site	Percent (95% C.I.)	Percent (95% C.I.)	Percent (95% C.I.)
All Sites	43.31 (43.17, 43.44)	42.90 (42.75, 43.04)	41.63 (41.22, 42.04)
Invasive and In Situ	44.76 (44.63, 44.90)	44.34 (44.19, 44.49)	41.96 (41.55, 42.37)
Oral Cavity and Pharynx	1.55 (1.52, 1.57)	1.61 (1.58, 1.63)	1.15 (1.09, 1.22)
Esophagus	0.80 (0.78, 0.82)	0.85 (0.83, 0.87)	0.63 (0.59, 0.69)
Stomach	1.08 (1.06, 1.11)	0.98 (0.96, 1.01)	1.23 (1.16, 1.31)
Colon and Rectum	4.84 (4.80, 4.89)	4.73 (4.69, 4.78)	4.93 (4.79, 5.07)
Invasive and In Situ	5.03 (4.99, 5.08)	4.91 (4.86, 4.96)	5.17 (5.03, 5.32)
Liver and Intrahepatic Bile Duct	1.27 (1.25, 1.29)	1.13 (1.10, 1.15)	1.36 (1.30, 1.43)
Pancreas	1.52 (1.50, 1.55)	1.52 (1.49, 1.55)	1.51 (1.43, 1.59)
Larynx	0.59 (0.58, 0.61)	0.59 (0.58, 0.61)	0.75 (0.70, 0.81)
Invasive and In Situ	0.64 (0.62, 0.65)	0.64 (0.62, 0.65)	0.80 (0.75, 0.86)
Lung and Bronchus	7.43 (7.38, 7.49)	7.46 (7.40, 7.52)	7.78 (7.60, 7.96)
Melanoma of the Skin	2.56 (2.53, 2.59)	2.95 (2.91, 2.99)	0.08 (0.06, 0.11)
Invasive and In Situ	4.26 (4.22, 4.30)	4.77 (4.72, 4.82)	0.11 (0.09, 0.14)
Breast	0.13 (0.12, 0.14)	0.13 (0.12, 0.14)	0.15 (0.12, 0.18)
Invasive and In Situ	0.14 (0.14, 0.15)	0.14 (0.13, 0.15)	0.17 (0.14, 0.20)
Prostate	15.02 (14.95, 15.10)	14.16 (14.08, 14.24)	19.08 (18.82, 19.35)
Testis	0.38 (0.37, 0.39)	0.46 (0.45, 0.47)	0.09 (0.08, 0.11)
Urinary Bladder (Invasive and In Situ)	3.83 (3.79, 3.87)	4.15 (4.10, 4.19)	1.79 (1.70, 1.89)
Kidney and Renal Pelvis	2.04 (2.02, 2.07)	2.10 (2.07, 2.13)	1.94 (1.86, 2.03)
Brain and Other Nervous System	0.69 (0.67, 0.71)	0.76 (0.74, 0.78)	0.36 (0.33, 0.40)
Thyroid	0.57 (0.56, 0.59)	0.61 (0.60, 0.63)	0.26 (0.23, 0.29)
Hodgkin Lymphoma	0.24 (0.23, 0.25)	0.26 (0.25, 0.27)	0.21 (0.19, 0.24)
Non-Hodgkin Lymphoma	2.36 (2.33, 2.39)	2.46 (2.43, 2.50)	1.41 (1.34, 1.49)
Myeloma	0.83 (0.81, 0.85)	0.78 (0.76, 0.80)	1.25 (1.18, 1.33)
Leukemia	1.70 (1.67, 1.72)	1.77 (1.74, 1.80)	1.07 (1.00, 1.14)
Acute Lymphocytic Leukemia	0.14 (0.14, 0.15)	0.16 (0.15, 0.17)	0.08 (0.07, 0.10)
Chronic Lymphocytic Leukemia	0.66 (0.65, 0.68)	0.70 (0.68, 0.71)	0.37 (0.33, 0.42)
Acute Myeloid Leukemia	0.49 (0.48, 0.51)	0.51 (0.50, 0.53)	0.34 (0.30, 0.38)
Chronic Myeloid Leukemia	0.22 (0.21, 0.23)	0.22 (0.21, 0.23)	0.15 (0.13, 0.18)
Kaposi Sarcoma	0.08 (0.07, 0.08)	0.07 (0.06, 0.07)	0.14 (0.12, 0.16)
Mesothelioma	0.21 (0.20, 0.22)	0.23 (0.22, 0.24)	0.10 (0.08, 0.13)

Devcan Version 6.8.0, August 2014, National Cancer Institute (<u>http://surveillance.cancer.gov/devcan</u>/). Source: SEER 18 areas (San Francisco, Connecticut, Detroit, Hawaii, Iowa, New Mexico, Seattle, Utah, Atlanta, San Jose-Monterey, Los Angeles, Alaska Native Registry, Rural Georgia, California excluding SF/SJM/LA, Kentucky, Louisiana, New Jersey, and Georgia excluding ATL/RG).

Note: Invasive cancer only unless specified otherwise.

A percent or confidence interval value of 0.00 represents a value that is below 0.005.

Table 1.16 - continued

Lifetime Risk (Percent) of Being Diagnosed with Cancer by Site and Race/Ethnicity

Males, 18 SEER Areas, 2009-2011

	Asian/Pacific Islanders	American Indian/ Alaska Natives ^a	Hispanics ^b		
Site	Percent (95% C.I.)	Percent (95% C.I.)	Percent (95% C.I.)		
All Sites	36.92 (36.39, 37.47)	29.98 (28.29, 31.89)	40.34 (39.85, 40.85)		
Invasive and In Situ	37.26 (36.72, 37.81)	30.25 (28.55, 32.16)	40.78 (40.28, 41.29)		
Oral Cavity and Pharynx	1.25 (1.16, 1.36)	1.24 (0.96, 1.81)	1.12 (1.04, 1.21)		
Esophagus	0.53 (0.47, 0.62)	0.49 (0.30, 1.03)	0.61 (0.56, 0.68)		
Stomach	2.11 (1.97, 2.26)	1.36 (0.99, 2.01)	1.80 (1.70, 1.93)		
Colon and Rectum	5.33 (5.13, 5.54)	4.62 (3.96, 5.52)	4.92 (4.75, 5.11)		
Invasive and In Situ	5.53 (5.32, 5.74)	4.73 (4.07, 5.64)	5.08 (4.90, 5.26)		
Liver and Intrahepatic Bile Duct	2.66 (2.53, 2.80)	2.09 (1.69, 2.76)	2.21 (2.11, 2.33)		
Pancreas	1.56 (1.45, 1.69)	1.28 (0.98, 1.87)	1.55 (1.45, 1.66)		
Larynx	0.35 (0.30, 0.41)	0.40 (0.25, 0.90)	0.56 (0.51, 0.64)		
Invasive and In Situ	0.37 (0.32, 0.44)	0.41 (0.25, 0.91)	0.59 (0.54, 0.66)		
Lung and Bronchus	6.83 (6.59, 7.09)	4.89 (4.21, 5.81)	4.96 (4.77, 5.17)		
Melanoma of the Skin	0.21 (0.17, 0.28)	0.39 (0.24, 0.89)	0.49 (0.44, 0.55)		
Invasive and In Situ	0.29 (0.24, 0.36)	0.57 (0.38, 1.08)	0.78 (0.71, 0.87)		
Breast	0.09 (0.07, 0.14)	0.02 (0.00, 0.52)	0.07 (0.06, 0.11)		
Invasive and In Situ	0.10 (0.07, 0.15)	0.03 (0.01, 0.53)	0.08 (0.06, 0.12)		
Prostate	10.10 (9.84, 10.37)	7.01 (6.25, 8.01)	14.03 (13.76, 14.31)		
Testis	0.14 (0.12, 0.18)	0.34 (0.25, 0.79)	0.35 (0.33, 0.39)		
Urinary Bladder (Invasive and In Situ)	2.23 (2.09, 2.39)	1.64 (1.18, 2.38)	2.65 (2.50, 2.81)		
Kidney and Renal Pelvis	1.37 (1.27, 1.47)	2.43 (2.03, 3.08)	2.19 (2.08, 2.30)		
Brain and Other Nervous System	0.45 (0.40, 0.52)	0.28 (0.19, 0.75)	0.55 (0.51, 0.61)		
Thyroid	0.58 (0.53, 0.65)	0.35 (0.23, 0.82)	0.46 (0.42, 0.51)		
Hodgkin Lymphoma	0.13 (0.11, 0.17)	0.09 (0.04, 0.57)	0.24 (0.21, 0.29)		
Non-Hodgkin Lymphoma	2.06 (1.93, 2.19)	1.36 (1.06, 1.94)	2.39 (2.28, 2.52)		
Myeloma	0.63 (0.56, 0.71)	0.48 (0.28, 1.02)	0.88 (0.80, 0.97)		
Leukemia	1.17 (1.08, 1.27)	0.84 (0.58, 1.40)	1.37 (1.28, 1.47)		
Acute Lymphocytic Leukemia	0.12 (0.10, 0.16)	0.09 (0.04, 0.57)	0.19 (0.17, 0.23)		
Chronic Lymphocytic Leukemia	0.23 (0.19, 0.29)	0.21 (0.08, 0.73)	0.35 (0.30, 0.42)		
Acute Myeloid Leukemia	0.49 (0.44, 0.57)	0.22 (0.11, 0.71)	0.45 (0.41, 0.52)		
Chronic Myeloid Leukemia	0.21 (0.17, 0.26)	0.08 (0.03, 0.57)	0.20 (0.17, 0.26)		
Kaposi Sarcoma	0.04 (0.02, 0.07)	0.04 (0.00, 0.54)	0.12 (0.10, 0.16)		
Mesothelioma	0.10 (0.07, 0.15)	0.08 (0.01, 0.59)	0.19 (0.16, 0.24)		

Devcan Version 6.8.0, August 2014, National Cancer Institute (<u>http://surveillance.cancer.gov/devcan</u>/).

Source: SEER 18 areas (San Francisco, Connecticut, Detroit, Hawaii, Iowa, New Mexico, Seattle, Utah, Atlanta, San Jose-Monterey, Los Angeles, Alaska Native Registry, Rural Georgia, California excluding SF/SJM/LA, Kentucky, Louisiana, New Jersey, and Georgia excluding ATL/RG).

Note: Invasive cancer only unless specified otherwise.

Underlying incidence data for American Indian/Alaska Native are based on the CHSDA(Contract Health Service Delivery Area) counties.
 Hisparic is not mutually evaluative from whites blacks Asian Bacific Islanders and American Indians/Alaska Natives

Hispanic is not mutually exclusive from whites, blacks, Asian Pacific Islanders, and American Indians/Alaska Natives. Underlying incidence data for Hispanics are based on NHIA and exclude cases from the Alaska Native Registry. A percent or confidence interval value of 0.00 represents a value that is below 0.005.

Lifetime Risk (Percent) of Being Diagnosed with Cancer by Site and Race/Ethnicity

Females, 18 SEER Areas, 2009-2011

	All Races	Whites	Blacks		
Site	Percent (95% C.I.)	Percent (95% C.I.)	Percent (95% C.I.)		
All Sites	37.81 (37.70, 37.93)	38.53 (38.40, 38.67)	34.18 (33.84, 34.53)		
Invasive and In Situ	41.08 (40.96, 41.20)	41.83 (41.69, 41.97)	36.42 (36.08, 36.77)		
Oral Cavity and Pharynx	0.67 (0.66, 0.69)	0.70 (0.68, 0.72)	0.48 (0.45, 0.52)		
Esophagus	0.23 (0.22, 0.24)	0.23 (0.22, 0.24)	0.28 (0.25, 0.32)		
Stomach	0.67 (0.65, 0.69)	0.56 (0.54, 0.57)	1.00 (0.94, 1.07)		
Colon and Rectum	4.49 (4.45, 4.53)	4.40 (4.35, 4.44)	4.79 (4.66, 4.92)		
Invasive and In Situ	4.64 (4.60, 4.69)	4.54 (4.49, 4.58)	4.99 (4.86, 5.12)		
Liver and Intrahepatic Bile Duct	0.53 (0.51, 0.54)	0.46 (0.44, 0.47)	0.51 (0.47, 0.55)		
Pancreas	1.48 (1.46, 1.51)	1.45 (1.43, 1.48)	1.66 (1.58, 1.74)		
Larynx	0.13 (0.13, 0.14)	0.14 (0.13, 0.15)	0.17 (0.14, 0.19)		
Invasive and In Situ	0.14 (0.13, 0.15)	0.15 (0.14, 0.16)	0.17 (0.15, 0.20)		
Lung and Bronchus	6.17 (6.12, 6.22)	6.46 (6.41, 6.52)	5.39 (5.26, 5.53)		
Melanoma of the Skin	1.61 (1.58, 1.63)	1.89 (1.86, 1.91)	0.10 (0.09, 0.13)		
Invasive and In Situ	2.74 (2.71, 2.77)	3.13 (3.09, 3.17)	0.13 (0.11, 0.15)		
Breast	12.33 (12.27, 12.40)	12.70 (12.63, 12.77)	10.99 (10.82, 11.18)		
Invasive and In Situ	14.75 (14.68, 14.82)	15.13 (15.05, 15.20)	13.19 (13.00, 13.39)		
Cervix Uteri	0.65 (0.63, 0.66)	0.64 (0.62, 0.65)	0.78 (0.73, 0.82)		
Corpus and Uterus, NOS	2.73 (2.71, 2.76)	2.81 (2.78, 2.85)	2.48 (2.39, 2.57)		
Invasive and In Situ	2.76 (2.73, 2.79)	2.84 (2.80, 2.87)	2.51 (2.42, 2.60)		
Ovary ^a	1.33 (1.31, 1.35)	1.41 (1.38, 1.43)	0.99 (0.93, 1.05)		
Urinary Bladder (Invasive and In Situ)	1.14 (1.12, 1.16)	1.21 (1.19, 1.24)	0.81 (0.76, 0.87)		
Kidney and Renal Pelvis	1.19 (1.17, 1.21)	1.23 (1.20, 1.25)	1.29 (1.23, 1.36)		
Brain and Other Nervous System	0.55 (0.54, 0.56)	0.60 (0.59, 0.62)	0.34 (0.31, 0.37)		
Thyroid	1.68 (1.66, 1.70)	1.77 (1.75, 1.80)	1.00 (0.95, 1.05)		
Hodgkin Lymphoma	0.20 (0.19, 0.21)	0.22 (0.21, 0.22)	0.18 (0.16, 0.20)		
Non-Hodgkin Lymphoma	1.91 (1.89, 1.94)	2.01 (1.98, 2.04)	1.17 (1.11, 1.23)		
Myeloma	0.62 (0.61, 0.64)	0.55 (0.53, 0.57)	1.19 (1.12, 1.25)		
Leukemia	1.19 (1.17, 1.22)	1.24 (1.22, 1.27)	0.87 (0.82, 0.93)		
Acute Lymphocytic Leukemia	0.12 (0.11, 0.12)	0.13 (0.12, 0.14)	0.07 (0.06, 0.09)		
Chronic Lymphocytic Leukemia	0.43 (0.42, 0.44)	0.45 (0.44, 0.47)	0.27 (0.24, 0.31)		
Acute Myeloid Leukemia	0.38 (0.37, 0.39)	0.39 (0.38, 0.40)	0.29 (0.26, 0.33)		
Chronic Myeloid Leukemia	0.15 (0.14, 0.15)	0.15 (0.14, 0.16)	0.13 (0.11, 0.16)		
Kaposi Sarcoma	0.01 (0.01, 0.02)	0.01 (0.01, 0.02)	0.01 (0.01, 0.02)		
Mesothelioma	0.05 (0.05, 0.06)	0.06 (0.05, 0.06)	0.03 (0.02, 0.04)		

Devcan Version 6.8.0, August 2014, National Cancer Institute (<u>http://surveillance.cancer.gov/devcan</u>/). Source: SEER 18 areas (San Francisco, Connecticut, Detroit, Hawaii, Iowa, New Mexico, Seattle, Utah, Atlanta, San Jose-Monterey, Los Angeles, Alaska Native Registry, Rural Georgia, California excluding SF/SJM/LA, Kentucky, Louisiana, New Jersey, and Georgia excluding ATL/RG).

Note:

Invasive cancer only unless specified otherwise. Ovary excludes borderline cases or histologies 8442, 8451, 8462, 8472, and 8473.

A percent or confidence interval value of 0.00 represents a value that is below 0.005.

Table 1.17 - continued

Lifetime Risk (Percent) of Being Diagnosed with Cancer by Site and Race/Ethnicity

Females, 18 SEER Areas, 2009-2011

	Asian/Pacific Islanders	American Indian/ Alaska Natives ^a	Hispanics ^b		
Site	Percent (95% C.I.)	Percent (95% C.I.)	Percent (95% C.I.)		
All Sites	33.08 (32.62, 33.55)	28.63 (27.02, 30.41)	34.72 (34.30, 35.14)		
Invasive and In Situ	35.36 (34.89, 35.84)	29.94 (28.32, 31.74)	36.68 (36.26, 37.12)		
Oral Cavity and Pharynx	0.56 (0.50, 0.63)	0.40 (0.27, 0.81)	0.54 (0.48, 0.60)		
Esophagus	0.18 (0.14, 0.24)	0.16 (0.08, 0.56)	0.17 (0.14, 0.21)		
Stomach	1.50 (1.39, 1.63)	0.84 (0.60, 1.31)	1.23 (1.15, 1.33)		
Colon and Rectum	4.71 (4.51, 4.91)	3.89 (3.29, 4.68)	4.24 (4.08, 4.41)		
Invasive and In Situ	4.86 (4.67, 5.07)	3.98 (3.38, 4.78)	4.38 (4.22, 4.55)		
Liver and Intrahepatic Bile Duct	1.30 (1.20, 1.41)	0.90 (0.66, 1.39)	1.03 (0.96, 1.11)		
Pancreas	1.60 (1.49, 1.72)	1.17 (0.83, 1.74)	1.71 (1.60, 1.83)		
Larynx	0.04 (0.03, 0.07)	0.11 (0.04, 0.50)	0.07 (0.05, 0.10)		
Invasive and In Situ	0.04 (0.03, 0.07)	0.12 (0.05, 0.52)	0.07 (0.06, 0.10)		
Lung and Bronchus	4.50 (4.33, 4.69)	4.11 (3.50, 4.91)	3.75 (3.60, 3.90)		
Melanoma of the Skin	0.15 (0.12, 0.20)	0.37 (0.24, 0.77)	0.50 (0.45, 0.57)		
Invasive and In Situ	0.20 (0.17, 0.26)	0.61 (0.42, 1.05)	0.79 (0.72, 0.86)		
Breast	9.93 (9.72, 10.15)	7.92 (7.18, 8.83)	9.76 (9.56, 9.96)		
Invasive and In Situ	12.33 (12.11, 12.58)	8.97 (8.21, 9.91)	11.45 (11.24, 11.66)		
Cervix Uteri	0.65 (0.59, 0.72)	0.60 (0.45, 1.00)	0.91 (0.86, 0.98)		
Corpus and Uterus, NOS	2.16 (2.07, 2.27)	2.05 (1.75, 2.57)	2.34 (2.26, 2.44)		
Invasive and In Situ	2.18 (2.08, 2.28)	2.07 (1.76, 2.59)	2.36 (2.27, 2.46)		
Ovary ^c	1.03 (0.96, 1.12)	1.13 (0.82, 1.68)	1.26 (1.18, 1.34)		
Urinary Bladder (Invasive and In Situ)	0.72 (0.64, 0.82)	0.32 (0.17, 0.75)	0.80 (0.73, 0.89)		
Kidney and Renal Pelvis	0.74 (0.67, 0.82)	1.51 (1.22, 2.03)	1.38 (1.30, 1.46)		
Brain and Other Nervous System	0.34 (0.30, 0.40)	0.31 (0.12, 0.81)	0.52 (0.47, 0.58)		
Thyroid	1.77 (1.69, 1.86)	1.15 (0.85, 1.67)	1.62 (1.56, 1.70)		
Hodgkin Lymphoma	0.11 (0.09, 0.14)	0.10 (0.04, 0.49)	0.19 (0.17, 0.23)		
Non-Hodgkin Lymphoma	1.64 (1.54, 1.75)	1.31 (0.97, 1.87)	2.10 (1.99, 2.21)		
Myeloma	0.46 (0.41, 0.53)	0.41 (0.24, 0.85)	0.67 (0.61, 0.74)		
Leukemia	0.78 (0.71, 0.86)	0.73 (0.50, 1.20)	1.07 (1.00, 1.15)		
Acute Lymphocytic Leukemia	0.09 (0.07, 0.12)	0.10 (0.05, 0.49)	0.18 (0.16, 0.21)		
Chronic Lymphocytic Leukemia	0.11 (0.09, 0.15)	0.16 (0.06, 0.58)	0.25 (0.21, 0.30)		
Acute Myeloid Leukemia	0.37 (0.33, 0.43)	0.30 (0.15, 0.73)	0.37 (0.33, 0.43)		
Chronic Myeloid Leukemia	0.10 (0.08, 0.14)	0.11 (0.06, 0.50)	0.15 (0.12, 0.20)		
Kaposi Sarcoma	0.00 (0.00, 0.03)	0.01 (0.00, 0.42)	0.05 (0.03, 0.08)		
Mesothelioma	0.02 (0.01, 0.05)	0.04 (0.00, 0.45)	0.05 (0.04, 0.08)		

Devcan Version 6.8.0, August 2014, National Cancer Institute (<u>http://surveillance.cancer.gov/devcan</u>/).

Source: SEER 18 areas (San Francisco, Connecticut, Detroit, Hawaii, Iowa, New Mexico, Seattle, Utah, Atlanta, San Jose-Monterey, Los Angeles, Alaska Native Registry, Rural Georgia, California excluding SF/SJM/LA, Kentucky, Louisiana, New Jersey, and Georgia excluding ATL/RG).

Note: Invasive cancer only unless specified otherwise.

^a Underlying incidence data for American Indian/Alaska Native are based on the CHSDA(Contract Health Service Delivery Area) counties.
 ^b Hispanic is not mutually evaluative from whites blacks Asian Pacific Islanders and American Indians/Alaska Natives

^b Hispanic is not mutually exclusive from whites, blacks, Asian Pacific Islanders, and American Indians/Alaska Natives.
 Underlying incidence data for Hispanics are based on NHIA and exclude cases from the Alaska Native Registry.
 Overviewed Verderlying appear or bictelogics 2442, 2451, 2462, 2472, and 2472.

Ovary excludes borderline cases or histologies 8442, 8451, 8462, 8472, and 8473. A percent or confidence interval value of 0.00 represents a value that is below 0.005.

Lifetime Risk (Percent) of Dying from Cancer by Site and Race/Ethnicity

Both Sexes, Total U.S., 2009-2011

	All Races	Whites	Blacks
Site	Percent (95% C.I.)	Percent (95% C.I.)	Percent (95% C.I.)
All Sites	20.84 (20.81, 20.87)	20.95 (20.92, 20.98)	21.17 (21.08, 21.27)
Oral Cavity and Pharynx	0.28 (0.28, 0.29)	0.28 (0.28, 0.29)	0.28 (0.27, 0.29)
Esophagus	0.49 (0.48, 0.49)	0.51 (0.50, 0.52)	0.40 (0.39, 0.42)
Stomach	0.41 (0.40, 0.41)	0.36 (0.35, 0.36)	0.67 (0.65, 0.69)
Colon and Rectum	1.94 (1.93, 1.95)	1.90 (1.89, 1.91)	2.29 (2.26, 2.33)
Liver and Intrahepatic Bile Duct	0.68 (0.67, 0.69)	0.64 (0.63, 0.64)	0.78 (0.76, 0.80)
Pancreas	1.34 (1.33, 1.35)	1.33 (1.32, 1.34)	1.43 (1.40, 1.46)
Larynx	0.12 (0.12, 0.13)	0.12 (0.12, 0.12)	0.18 (0.17, 0.19)
Lung and Bronchus	5.66 (5.64, 5.68)	5.79 (5.77, 5.81)	5.21 (5.17, 5.26)
Melanoma of the Skin	0.32 (0.31, 0.32)	0.36 (0.36, 0.37)	0.04 (0.04, 0.05)
Breast	1.43 (1.42, 1.44)	1.41 (1.40, 1.42)	1.78 (1.75, 1.81)
Urinary Bladder	0.60 (0.60, 0.61)	0.63 (0.62, 0.64)	0.42 (0.40, 0.43)
Kidney and Renal Pelvis	0.47 (0.47, 0.48)	0.49 (0.48, 0.49)	0.39 (0.38, 0.41)
Brain and Other Nervous System	0.45 (0.45, 0.46)	0.49 (0.49, 0.50)	0.23 (0.22, 0.24)
Thyroid	0.06 (0.06, 0.07)	0.06 (0.06, 0.07)	0.05 (0.05, 0.06)
Hodgkin Lymphoma	0.04 (0.04, 0.04)	0.04 (0.04, 0.04)	0.03 (0.03, 0.03)
Non-Hodgkin Lymphoma	0.78 (0.77, 0.78)	0.82 (0.81, 0.82)	0.45 (0.43, 0.46)
Myeloma	0.42 (0.41, 0.42)	0.39 (0.39, 0.40)	0.66 (0.64, 0.68)
Leukemia	0.86 (0.86, 0.87)	0.90 (0.89, 0.91)	0.62 (0.60, 0.64)
Acute Lymphocytic Leukemia	0.04 (0.04, 0.04)	0.04 (0.04, 0.05)	0.03 (0.02, 0.03)
Chronic Lymphocytic Leukemia	0.19 (0.19, 0.19)	0.20 (0.20, 0.20)	0.14 (0.13, 0.15)
Acute Myeloid Leukemia	0.33 (0.33, 0.34)	0.35 (0.34, 0.35)	0.21 (0.20, 0.22)
Chronic Myeloid Leukemia	0.04 (0.04, 0.04)	0.04 (0.04, 0.04)	0.03 (0.03, 0.04)

Devcan Version 6.8.0, August 2014, National Cancer Institute (<u>http://surveillance.cancer.gov/devcan</u>/). Source: NCHS public use data file for the total US.

A percent or confidence interval value of 0.00 represents a value that is below 0.005.

Table 1.18 - continued

Lifetime Risk (Percent) of Dying from Cancer by Site and Race/Ethnicity

Both Sexes, Total U.S., 2009-2011

	Asian/Pacific Islanders	American Indian/ Alaska Natives ^a	Hispanics ^b
Site	Percent (95% C.I.)	Percent (95% C.I.)	Percent (95% C.I.)
All Sites	18.69 (18.45, 18.93)	16.99 (16.51, 17.50)	17.71 (17.57, 17.86)
Oral Cavity and Pharynx	0.30 (0.27, 0.34)	0.24 (0.19, 0.33)	0.21 (0.19, 0.23)
Esophagus	0.30 (0.27, 0.34)	0.35 (0.28, 0.45)	0.35 (0.32, 0.37)
Stomach	1.14 (1.07, 1.21)	0.55 (0.46, 0.67)	0.79 (0.76, 0.83)
Colon and Rectum	2.02 (1.94, 2.12)	1.87 (1.70, 2.08)	1.89 (1.84, 1.95)
Liver and Intrahepatic Bile Duct	1.55 (1.48, 1.62)	1.00 (0.89, 1.14)	1.20 (1.17, 1.24)
Pancreas	1.40 (1.33, 1.47)	0.93 (0.81, 1.08)	1.33 (1.29, 1.37)
Larynx	0.07 (0.05, 0.09)	0.10 (0.07, 0.16)	0.12 (0.11, 0.14)
Lung and Bronchus	4.30 (4.19, 4.41)	4.12 (3.88, 4.38)	3.01 (2.95, 3.07)
Melanoma of the Skin	0.06 (0.04, 0.08)	0.12 (0.08, 0.19)	0.12 (0.11, 0.13)
Breast	0.97 (0.91, 1.03)	0.91 (0.79, 1.06)	1.11 (1.07, 1.15)
Urinary Bladder	0.39 (0.35, 0.44)	0.29 (0.21, 0.39)	0.43 (0.40, 0.46)
Kidney and Renal Pelvis	0.35 (0.31, 0.39)	0.74 (0.63, 0.88)	0.50 (0.47, 0.52)
Brain and Other Nervous System	0.27 (0.24, 0.30)	0.19 (0.16, 0.26)	0.33 (0.32, 0.35)
Thyroid	0.12 (0.10, 0.15)	0.05 (0.03, 0.12)	0.08 (0.08, 0.10)
Hodgkin Lymphoma	0.03 (0.02, 0.05)	0.02 (0.01, 0.08)	0.05 (0.05, 0.06)
Non-Hodgkin Lymphoma	0.74 (0.69, 0.80)	0.50 (0.41, 0.61)	0.77 (0.74, 0.80)
Myeloma	0.29 (0.27, 0.33)	0.31 (0.25, 0.41)	0.42 (0.40, 0.45)
Leukemia	0.66 (0.62, 0.71)	0.50 (0.41, 0.62)	0.68 (0.65, 0.71)
Acute Lymphocytic Leukemia	0.04 (0.03, 0.06)	0.02 (0.01, 0.08)	0.07 (0.06, 0.07)
Chronic Lymphocytic Leukemia	0.06 (0.04, 0.08)	0.10 (0.05, 0.19)	0.09 (0.08, 0.10)
Acute Myeloid Leukemia	0.32 (0.30, 0.36)	0.19 (0.15, 0.27)	0.26 (0.24, 0.28)
Chronic Myeloid Leukemia	0.04 (0.03, 0.06)	0.03 (0.01, 0.09)	0.04 (0.03, 0.04)

Devcan Version 6.8.0, August 2014, National Cancer Institute (<u>http://surveillance.cancer.gov/devcan</u>/).

- Source: NCHS public use data file for the total US.
- Underlying mortality data for American Indian/Alaska Native are based on the CHSDA (Contract Health Service Delivery Area) counties.
- Hispanic is not mutually exclusive from whites, blacks, Asian Pacific Islanders, and American Indians/Alaska Natives. A percent or confidence interval value of 0.00 represents a value that is below 0.005.

b

Lifetime Risk (Percent) of Dying from Cancer by Site and Race/Ethnicity

Males, Total U.S., 2009-2011

	All Races	Whites	Blacks
Site	Percent (95% C.I.)	Percent (95% C.I.)	Percent (95% C.I.)
All Sites	22.83 (22.78, 22.87)	22.87 (22.82, 22.92)	23.66 (23.51, 23.81)
Oral Cavity and Pharynx	0.39 (0.38, 0.39)	0.38 (0.38, 0.39)	0.44 (0.42, 0.46)
Esophagus	0.79 (0.78, 0.80)	0.83 (0.82, 0.84)	0.62 (0.59, 0.64)
Stomach	0.49 (0.48, 0.50)	0.43 (0.42, 0.44)	0.82 (0.79, 0.85)
Colon and Rectum	2.04 (2.02, 2.05)	2.00 (1.98, 2.01)	2.41 (2.36, 2.46)
Liver and Intrahepatic Bile Duct	0.90 (0.89, 0.91)	0.84 (0.83, 0.85)	1.08 (1.05, 1.11)
Pancreas	1.35 (1.34, 1.37)	1.36 (1.35, 1.38)	1.35 (1.31, 1.39)
Larynx	0.20 (0.20, 0.21)	0.19 (0.19, 0.20)	0.32 (0.30, 0.34)
Lung and Bronchus	6.47 (6.45, 6.50)	6.53 (6.51, 6.56)	6.49 (6.42, 6.58)
Melanoma of the Skin	0.43 (0.43, 0.44)	0.49 (0.49, 0.50)	0.04 (0.04, 0.05)
Breast	0.03 (0.03, 0.03)	0.03 (0.03, 0.03)	0.05 (0.04, 0.05)
Prostate	2.66 (2.65, 2.68)	2.48 (2.46, 2.50)	4.57 (4.49, 4.66)
Testis	0.02 (0.02, 0.02)	0.02 (0.02, 0.02)	0.01 (0.01, 0.01)
Urinary Bladder	0.91 (0.90, 0.92)	0.97 (0.95, 0.98)	0.51 (0.48, 0.54)
Kidney and Renal Pelvis	0.61 (0.60, 0.62)	0.63 (0.62, 0.64)	0.48 (0.46, 0.51)
Brain and Other Nervous System	0.51 (0.50, 0.51)	0.55 (0.54, 0.56)	0.24 (0.23, 0.26)
Thyroid	0.05 (0.05, 0.06)	0.06 (0.05, 0.06)	0.03 (0.03, 0.04)
Hodgkin Lymphoma	0.05 (0.04, 0.05)	0.05 (0.05, 0.05)	0.04 (0.03, 0.04)
Non-Hodgkin Lymphoma	0.87 (0.86, 0.88)	0.92 (0.90, 0.93)	0.51 (0.48, 0.53)
Myeloma	0.47 (0.46, 0.47)	0.45 (0.44, 0.45)	0.68 (0.66, 0.71)
Leukemia	1.03 (1.01, 1.04)	1.07 (1.06, 1.08)	0.69 (0.66, 0.72)
Acute Lymphocytic Leukemia	0.05 (0.04, 0.05)	0.05 (0.05, 0.05)	0.03 (0.02, 0.03)
Chronic Lymphocytic Leukemia	0.24 (0.23, 0.24)	0.25 (0.24, 0.25)	0.17 (0.15, 0.19)
Acute Myeloid Leukemia	0.39 (0.39, 0.40)	0.41 (0.41, 0.42)	0.23 (0.22, 0.25)
Chronic Myeloid Leukemia	0.04 (0.04, 0.05)	0.05 (0.04, 0.05)	0.04 (0.03, 0.05)

Devcan Version 6.8.0, August 2014, National Cancer Institute (<u>http://surveillance.cancer.gov/devcan</u>/). Source: NCHS public use data file for the total US.

A percent or confidence interval value of 0.00 represents a value that is below 0.005.

Table 1.19 - continued

Lifetime Risk (Percent) of Dying from Cancer by Site and Race/Ethnicity

Males, Total U.S., 2009-2011

	Asian/Pacific Islanders	American Indian/ Alaska Natives ^a	Hispanics ^b
Site	Percent (95% C.I.)	Percent (95% C.I.)	Percent (95% C.I.)
All Sites	21.04 (20.68, 21.41)	18.46 (17.69, 19.29)	20.28 (20.05, 20.52)
Oral Cavity and Pharynx	0.41 (0.36, 0.48)	0.33 (0.24, 0.54)	0.29 (0.27, 0.32)
Esophagus	0.47 (0.41, 0.54)	0.48 (0.37, 0.70)	0.58 (0.54, 0.62)
Stomach	1.31 (1.22, 1.42)	0.69 (0.55, 0.92)	0.94 (0.89, 1.00)
Colon and Rectum	2.12 (1.99, 2.25)	1.83 (1.57, 2.16)	2.13 (2.05, 2.21)
Liver and Intrahepatic Bile Duct	2.02 (1.92, 2.14)	1.30 (1.12, 1.57)	1.55 (1.49, 1.61)
Pancreas	1.33 (1.24, 1.43)	0.86 (0.72, 1.10)	1.30 (1.24, 1.36)
Larynx	0.13 (0.10, 0.17)	0.16 (0.10, 0.34)	0.23 (0.21, 0.26)
Lung and Bronchus	5.43 (5.25, 5.63)	4.63 (4.25, 5.09)	3.97 (3.87, 4.08)
Melanoma of the Skin	0.06 (0.04, 0.09)	0.19 (0.12, 0.38)	0.15 (0.12, 0.18)
Breast	0.03 (0.01, 0.07)	0.02 (0.00, 0.20)	0.03 (0.02, 0.05)
Prostate	2.20 (2.04, 2.37)	2.38 (2.02, 2.84)	3.08 (2.96, 3.21)
Testis	0.00 (0.00, 0.03)	0.02 (0.01, 0.20)	0.03 (0.02, 0.04)
Urinary Bladder	0.58 (0.50, 0.67)	0.50 (0.33, 0.78)	0.65 (0.59, 0.71)
Kidney and Renal Pelvis	0.45 (0.40, 0.52)	0.93 (0.76, 1.20)	0.64 (0.60, 0.69)
Brain and Other Nervous System	0.31 (0.27, 0.36)	0.23 (0.17, 0.41)	0.36 (0.34, 0.39)
Thyroid	0.07 (0.06, 0.11)	0.05 (0.02, 0.23)	0.06 (0.05, 0.08)
Hodgkin Lymphoma	0.04 (0.02, 0.07)	0.04 (0.01, 0.22)	0.06 (0.05, 0.08)
Non-Hodgkin Lymphoma	0.83 (0.76, 0.93)	0.52 (0.41, 0.73)	0.82 (0.77, 0.87)
Myeloma	0.36 (0.31, 0.42)	0.31 (0.21, 0.51)	0.48 (0.45, 0.53)
Leukemia	0.78 (0.71, 0.85)	0.68 (0.52, 0.95)	0.77 (0.73, 0.83)
Acute Lymphocytic Leukemia	0.05 (0.03, 0.08)	0.02 (0.01, 0.20)	0.07 (0.06, 0.09)
Chronic Lymphocytic Leukemia	0.07 (0.05, 0.11)	0.15 (0.07, 0.35)	0.10 (0.08, 0.12)
Acute Myeloid Leukemia	0.37 (0.33, 0.43)	0.23 (0.15, 0.43)	0.29 (0.27, 0.32)
Chronic Myeloid Leukemia	0.04 (0.03, 0.08)	0.06 (0.03, 0.24)	0.05 (0.04, 0.07)

Devcan Version 6.8.0, August 2014, National Cancer Institute (http://surveillance.cancer.gov/devcan/).

- Source: NCHS public use data file for the total US. ^a Underlying mortality data for American Indian/Alaska Native are based on the CHSDA (Contract Health Service Delivery Area) counties. b
 - Hispanic is not mutually exclusive from whites, blacks, Asian Pacific Islanders, and American Indians/Alaska Natives. A percent or confidence interval value of 0.00 represents a value that is below 0.005.

Lifetime Risk (Percent) of Dying from Cancer by Site and Race/Ethnicity

Females, Total U.S., 2009-2011

	All Races	Whites	Blacks
Site	Percent (95% C.I.)	Percent (95% C.I.)	Percent (95% C.I.)
All Sites	19.26 (19.22, 19.30)	19.41 (19.37, 19.45)	19.37 (19.25, 19.50)
Oral Cavity and Pharynx	0.18 (0.18, 0.19)	0.18 (0.18, 0.19)	0.15 (0.13, 0.16)
Esophagus	0.21 (0.21, 0.22)	0.21 (0.21, 0.22)	0.22 (0.20, 0.23)
Stomach	0.33 (0.32, 0.33)	0.29 (0.28, 0.29)	0.55 (0.52, 0.57)
Colon and Rectum	1.85 (1.84, 1.87)	1.81 (1.79, 1.82)	2.21 (2.16, 2.26)
Liver and Intrahepatic Bile Duct	0.47 (0.46, 0.48)	0.44 (0.44, 0.45)	0.51 (0.49, 0.53)
Pancreas	1.32 (1.31, 1.33)	1.31 (1.29, 1.32)	1.50 (1.46, 1.54)
Larynx	0.05 (0.05, 0.05)	0.05 (0.05, 0.05)	0.07 (0.06, 0.07)
Lung and Bronchus	4.95 (4.93, 4.97)	5.14 (5.11, 5.16)	4.15 (4.09, 4.21)
Melanoma of the Skin	0.21 (0.21, 0.22)	0.24 (0.24, 0.25)	0.05 (0.04, 0.05)
Breast	2.72 (2.70, 2.74)	2.69 (2.67, 2.71)	3.26 (3.21, 3.32)
Cervix Uteri	0.23 (0.22, 0.23)	0.21 (0.20, 0.21)	0.38 (0.37, 0.40)
Corpus and Uterus, NOS	0.57 (0.56, 0.58)	0.53 (0.53, 0.54)	0.89 (0.86, 0.92)
Ovary	0.98 (0.97, 0.99)	1.02 (1.01, 1.03)	0.76 (0.73, 0.78)
Urinary Bladder	0.34 (0.34, 0.35)	0.35 (0.34, 0.35)	0.35 (0.33, 0.37)
Kidney and Renal Pelvis	0.35 (0.34, 0.35)	0.35 (0.35, 0.36)	0.32 (0.30, 0.34)
Brain and Other Nervous System	0.40 (0.39, 0.41)	0.43 (0.43, 0.44)	0.22 (0.21, 0.24)
Thyroid	0.07 (0.07, 0.07)	0.07 (0.07, 0.07)	0.07 (0.06, 0.08)
Hodgkin Lymphoma	0.03 (0.03, 0.04)	0.04 (0.03, 0.04)	0.03 (0.02, 0.03)
Non-Hodgkin Lymphoma	0.69 (0.69, 0.70)	0.73 (0.72, 0.74)	0.40 (0.38, 0.42)
Myeloma	0.38 (0.37, 0.38)	0.35 (0.34, 0.35)	0.64 (0.62, 0.67)
Leukemia	0.72 (0.72, 0.73)	0.75 (0.74, 0.76)	0.57 (0.55, 0.59)
Acute Lymphocytic Leukemia	0.04 (0.04, 0.04)	0.04 (0.04, 0.04)	0.02 (0.02, 0.03)
Chronic Lymphocytic Leukemia	0.15 (0.15, 0.16)	0.16 (0.15, 0.16)	0.12 (0.11, 0.13)
Acute Myeloid Leukemia	0.28 (0.27, 0.28)	0.29 (0.29, 0.30)	0.19 (0.18, 0.21)
Chronic Myeloid Leukemia	0.04 (0.03, 0.04)	0.04 (0.03, 0.04)	0.03 (0.03, 0.04)

Devcan Version 6.8.0, August 2014, National Cancer Institute (<u>http://surveillance.cancer.gov/devcan</u>/). Source: NCHS public use data file for the total US.

A percent or confidence interval value of 0.00 represents a value that is below 0.005.

Table 1.20 - continued

Lifetime Risk (Percent) of Dying from Cancer by Site and Race/Ethnicity

Females, Total U.S., 2009-2011

	Asian/Pacific Islanders	American Indian/ Alaska Natives ^a	Hispanics ^b
Site	Percent (95% C.I.)	Percent (95% C.I.)	Percent (95% C.I.)
All Sites	16.84 (16.52, 17.16)	15.93 (15.28, 16.62)	15.81 (15.63, 15.99)
Oral Cavity and Pharynx	0.20 (0.17, 0.25)	0.16 (0.10, 0.28)	0.14 (0.12, 0.16)
Esophagus	0.16 (0.13, 0.20)	0.23 (0.15, 0.37)	0.15 (0.13, 0.18)
Stomach	0.99 (0.90, 1.09)	0.42 (0.31, 0.58)	0.67 (0.63, 0.71)
Colon and Rectum	1.95 (1.83, 2.09)	1.92 (1.67, 2.21)	1.71 (1.64, 1.78)
Liver and Intrahepatic Bile Duct	1.15 (1.07, 1.24)	0.71 (0.58, 0.90)	0.89 (0.85, 0.94)
Pancreas	1.45 (1.36, 1.56)	0.98 (0.80, 1.20)	1.35 (1.30, 1.42)
Larynx	0.02 (0.01, 0.04)	0.03 (0.01, 0.13)	0.03 (0.02, 0.05)
Lung and Bronchus	3.39 (3.25, 3.53)	3.70 (3.39, 4.05)	2.23 (2.16, 2.31)
Melanoma of the Skin	0.05 (0.04, 0.09)	0.06 (0.03, 0.16)	0.10 (0.08, 0.11)
Breast	1.75 (1.65, 1.87)	1.73 (1.51, 2.00)	2.05 (1.98, 2.12)
Cervix Uteri	0.24 (0.21, 0.29)	0.31 (0.23, 0.45)	0.31 (0.29, 0.34)
Corpus and Uterus, NOS	0.46 (0.41, 0.52)	0.39 (0.30, 0.54)	0.51 (0.48, 0.54)
Ovary	0.69 (0.63, 0.75)	0.83 (0.68, 1.03)	0.82 (0.78, 0.86)
Urinary Bladder	0.25 (0.20, 0.31)	0.14 (0.09, 0.25)	0.26 (0.24, 0.30)
Kidney and Renal Pelvis	0.27 (0.22, 0.33)	0.56 (0.43, 0.75)	0.37 (0.35, 0.41)
Brain and Other Nervous System	0.24 (0.21, 0.28)	0.16 (0.11, 0.26)	0.31 (0.28, 0.33)
Thyroid	0.16 (0.13, 0.21)	0.06 (0.02, 0.16)	0.11 (0.09, 0.13)
Hodgkin Lymphoma	0.02 (0.01, 0.05)	0.01 (0.00, 0.10)	0.04 (0.03, 0.05)
Non-Hodgkin Lymphoma	0.67 (0.61, 0.75)	0.47 (0.36, 0.64)	0.73 (0.69, 0.77)
Myeloma	0.24 (0.21, 0.28)	0.32 (0.23, 0.47)	0.38 (0.35, 0.41)
Leukemia	0.57 (0.51, 0.64)	0.36 (0.27, 0.51)	0.60 (0.57, 0.65)
Acute Lymphocytic Leukemia	0.04 (0.03, 0.06)	0.03 (0.01, 0.12)	0.06 (0.05, 0.07)
Chronic Lymphocytic Leukemia	0.05 (0.03, 0.08)	0.06 (0.02, 0.18)	0.08 (0.07, 0.11)
Acute Myeloid Leukemia	0.29 (0.25, 0.34)	0.17 (0.11, 0.28)	0.23 (0.21, 0.25)
Chronic Myeloid Leukemia	0.04 (0.02, 0.07)	0.01 (0.00, 0.10)	0.03 (0.02, 0.04)

Devcan Version 6.8.0, August 2014, National Cancer Institute (http://surveillance.cancer.gov/devcan/).

- Source: NCHS public use data file for the total US. ^a Underlying mortality data for American Indian/Alaska Native are based on the CHSDA (Contract Health Service Delivery Area) counties. b
 - Hispanic is not mutually exclusive from whites, blacks, Asian Pacific Islanders, and American Indians/Alaska Natives. A percent or confidence interval value of 0.00 represents a value that is below 0.005.

Table 1.21 U.S. and SEER Death Rates by Primary Cancer Site and Race/Ethnicity, 2007-2011

	Total United States ^a								SEER 18 Areas ^{ab}						
Site		Total	White	Black	AI/AN ^c	API ^d	Hisp ^e	W-NHisp ^e	Total	White	Black	AI/AN ^c	API ^d	Hisp ^e	W-NHisp ^e
All Sites	Both Sexes	173.8	173.3	206.4	158.0	107.8	120.3	177.3	168.2	170.0	206.5	136.3	114.1	121.5	175.8
	Male	211.6	209.8	269.3	190.0	131.0	150.1	214.0	202.6	203.4	266.6	161.3	139.2	148.8	209.6
	Female	147.4	147.5	169.0	135.2	91.5	99.9	151.2	144.4	146.8	170.5	118.3	96.5	102.7	152.2
Oral Cavity	Both Sexes	2.5	2.4	3.0	2.3	1.9	1.5	2.5	2.5	2.4	3.0	2.0	2.1	1.5	2.6
& Pharynx	Male	3.8	3.7	5.1	3.4	2.9	2.4	3.8	3.7	3.6	5.2	2.9	3.1	2.3	3.8
	Female	1.4	1.4	1.4	1.4	1.2	0.8	1.4	1.4	1.4	1.4	-	1.3	0.8	1.5
Esophagus	Both Sexes	4.2	4.3	4.2	3.3	1.7	2.3	4.5	3.8	4.1	4.1	2.3	1.8	2.2	4.3
	Male	7.5	7.8	7.4	5.7	3.0	4.3	8.1	6.8	7.2	6.9	3.8	3.1	4.2	7.6
	Female	1.6	1.5	2.0	1.5	0.8	0.8	1.6	1.5	1.5	2.0	1.3	0.8	0.8	1.6
Stomach	Both Sexes	3.5	3.0	6.5	5.2	6.3	5.6	2.7	4.0	3.4	6.8	6.1	6.5	6.2	2.9
	Male	4.7	4.1	9.6	7.0	8.3	7.5	3.8	5.4	4.6	9.5	8.0	8.5	8.1	4.1
	Female	2.5	2.1	4.5	3.8	4.8	4.2	1.9	2.9	2.4	5.0	4.4	5.0	4.8	2.0
Colon &	Both Sexes	15.9	15.5	22.1	17.2	11.0	12.4	15.7	15.5	15.2	22.4	16.5	11.6	12.0	15.5
Rectum	Male	19.1	18.5	27.7	19.2	13.1	15.8	18.7	18.5	18.1	27.7	18.9	13.9	15.4	18.4
	Female	13.5	13.0	18.5	15.6	9.5	9.9	13.2	13.2	12.9	18.9	14.6	9.9	9.4	13.3
Liver &	Both Sexes	5.8	5.3	7.6	9.5	9.8	8.7	5.0	6.2	5.6	7.6	9.0	10.0	8.8	5.1
Intrahepatic	Male	8.5	7.8	12.1	13.8	14.5	12.6	7.3	9.1	8.2	12.1	13.2	14.9	12.6	7.4
Bile Duct	Female	3.4	3.2	4.2	6.0	6.0	5.5	3.0	3.8	3.4	4.2	5.7	6.1	5.6	3.1
Pancreas	Both Sexes	10.9	10.8	13.6	8.8	7.7	8.6	10.9	10.9	10.9	13.6	9.9	8.4	9.3	11.1
	Male	12.5	12.5	15.3	9.9	8.5	9.7	12.7	12.5	12.5	15.3	11.3	9.3	10.2	12.8
	Female	9.6	9.4	12.4	8.0	7.2	7.7	9.5	9.6	9.5	12.3	8.8	7.6	8.5	9.7
Larynx	Both Sexes	1.1	1.0	1.9	1.0	0.4	0.8	1.1	1.0	1.0	1.7	0.7	0.4	0.7	1.0
_	Male	2.0	1.8	3.8	1.8	0.8	1.7	1.9	1.8	1.7	3.3	-	0.9	1.5	1.7
	Female	0.4	0.4	0.6	-	0.1	0.2	0.4	0.4	0.4	0.7	-	0.1	0.2	0.4
Lung &	Both Sexes	48.4	49.1	52.0	39.9	25.2	20.9	51.5	44.2	45.4	51.6	29.1	26.6	19.8	48.9
Bronchus	Male	61.6	61.4	75.7	50.0	34.7	30.5	63.9	55.4	55.6	73.7	36.0	37.2	27.8	59.3
	Female	38.5	39.8	36.5	32.4	18.4	14.0	42.1	35.9	37.7	37.0	24.0	19.0	14.0	41.1
Melanoma	Both Sexes	2.7	3.1	0.4	1.1	0.4	0.8	3.4	2.6	3.1	0.4	-	0.4	0.8	3.5
of the Skin	Male	4.1	4.6	0.5	1.6	0.4	1.1	5.0	3.9	4.6	0.6	-	0.5	1.0	5.1
	Female	1.7	2.0	0.4	0.7	0.3	0.6	2.1	1.6	1.9	0.3	-	0.3	0.6	2.2
Breast	Female	22.2	21.7	30.6	15.2	11.3	14.5	22.2	22.3	22.2	31.1	13.1	12.5	14.7	23.2

а US Mortality Files, National Center for Health Statistics, Centers for Disease Control and Prevention. Rates are per 100,000 b

and are age-adjusted to the 2000 US Std Population (19 age groups - Census P25-1130). The SEER 18 areas are San Francisco, Connecticut, Detroit, Hawaii, Iowa, New Mexico, Seattle, Utah, Atlanta, San Jose-Monterey, Los Angeles, Alaska Native Registry, Rural Georgia, California excluding SF/SJM/LA, Kentucky, Louisiana, New Jersey and Georgia excluding ATL/RG. С

Rates for American Indian/Alaska Native (AI/AN) are based on the CHSDA(Contract Health Service Delivery Area) counties.

d Asian/Pacific Islander.

e Hispanic (Hisp) and White Non-Hispanic (W-NHisp) are not mutually exclusive from whites, blacks, Asian/Pacific Islanders, and American Indians/Alaska Natives.

Statistic could not be calculated due to less than 16 cases in the time interval. _

Table 1.21 - continued U.S. and SEER Death Rates by Primary Cancer Site and Race/Ethnicity, 2007-2011

	Total United States ^a								SEER 18 Areas ^{ab}						
Site		Total	White	Black	AI/AN ^c	API ^d	Hisp ^e	W-NHisp ^e	Total	White	Black	AI/AN ^c	API ^d	Hisp ^e	W-NHisp ^e
Cervix	Female	2.3	2.1	4.1	3.4	1.8	2.8	2.0	2.3	2.2	3.9	3.0	1.9	2.9	2.0
Corpus & Uterus, NOS	Female	4.3	4.0	7.5	3.4	2.7	3.4	4.0	4.3	4.1	7.3	2.8	3.1	3.4	4.1
Ovary	Female	7.9	8.2	6.6	6.9	4.7	5.6	8.4	8.0	8.5	6.6	6.2	4.9	6.1	8.8
Prostate	Male	22.3	20.6	48.9	21.2	10.0	18.5	20.7	22.4	21.3	48.2	18.6	10.7	19.0	21.5
Testis	Male	0.2	0.3	0.1	-	0.1	0.3	0.3	0.2	0.3	0.1	-	0.1	0.3	0.3
Urinary Bladder	Both Sexes Male Female	4.4 7.7 2.2	4.6 8.1 2.2	3.6 5.4 2.6	2.5 4.4 1.3	1.7 2.9 0.9	2.4 4.0 1.3	4.8 8.4 2.3	4.3 7.4 2.2	4.6 8.1 2.3	3.7 5.5 2.7	2.1 3.6 -	1.7 3.0 0.9	2.3 3.8 1.3	4.9 8.6 2.4
Kidney & Renal Pelvis	Both Sexes Male Female	4.0 5.8 2.6	4.0 5.9 2.6	3.9 5.6 2.6	6.7 9.5 4.4	2.0 3.0 1.3	3.6 5.1 2.3	4.1 5.9 2.6	3.8 5.5 2.4	3.9 5.7 2.5	4.0 5.9 2.6	6.4 9.0 4.4	2.2 3.2 1.4	3.7 5.3 2.5	3.9 5.7 2.5
Brain & Nervous System	Both Sexes Male Female	4.3 5.2 3.5	4.6 5.6 3.8	2.5 3.0 2.1	2.6 2.9 2.3	1.9 2.3 1.5	2.8 3.3 2.4	4.8 5.9 3.9	4.2 5.2 3.4	4.7 5.8 3.8	2.6 3.1 2.2	1.9 2.3 1.5	2.0 2.4 1.6	3.0 3.4 2.6	5.0 6.2 4.1
Thyroid	Both Sexes Male Female	0.5 0.5 0.5	0.5 0.5 0.5	0.5 0.4 0.6	0.6 - 0.5	0.7 0.6 0.9	0.6 0.5 0.7	0.5 0.5 0.5	0.5 0.5 0.5	0.5 0.5 0.5	0.4 0.4 0.5	- - -	0.8 0.6 1.0	0.6 0.4 0.7	0.5 0.5 0.5
Hodgkin Lymphoma	Both Sexes Male Female	0.4 0.5 0.3	0.4 0.5 0.3	0.3 0.4 0.3	- - -	0.1 0.2 0.1	0.4 0.5 0.3	0.4 0.5 0.3	0.4 0.5 0.3	0.4 0.5 0.3	0.4 0.5 0.3	- - -	0.1 0.2 0.1	0.5 0.6 0.3	0.4 0.5 0.3
Non-Hodgkin Lymphoma	Both Sexes Male Female	6.3 8.1 5.0	6.6 8.4 5.2	4.5 5.8 3.5	4.6 5.3 3.9	4.1 5.2 3.4	5.3 6.4 4.4	6.6 8.5 5.2	6.1 7.9 4.9	6.5 8.3 5.1	4.4 6.0 3.3	3.8 4.2 3.5	4.5 5.6 3.7	5.5 6.8 4.5	6.6 8.4 5.2
Myeloma	Both Sexes Male Female	3.4 4.3 2.7	3.1 4.0 2.5	6.3 7.7 5.3	2.7 3.4 2.2	1.8 2.3 1.4	2.8 3.5 2.3	3.1 4.0 2.4	3.3 4.2 2.6	3.1 4.0 2.4	6.4 7.9 5.4	1.8 2.6 -	1.9 2.5 1.4	2.9 3.6 2.4	3.1 4.1 2.3
Leukemia	Both Sexes Male Female	7.0 9.4 5.3	7.3 9.7 5.4	6.0 8.0 4.8	4.7 6.6 3.5	4.0 5.0 3.2	4.8 6.0 3.9	7.4 9.9 5.5	6.8 9.1 5.1	7.2 9.6 5.4	6.1 7.9 4.9	3.7 4.4 3.3	4.2 5.3 3.4	4.8 6.0 3.9	7.3 9.9 5.4

а US Mortality Files, National Center for Health Statistics, Centers for Disease Control and Prevention. Rates are per 100,000 b

and are age-adjusted to the 2000 US Std Population (19 age groups - Census P25-1130). The SEER 18 areas are San Francisco, Connecticut, Detroit, Hawaii, Iowa, New Mexico, Seattle, Utah, Atlanta, San Jose-Monterey, Los Angeles, Alaska Native Registry, Rural Georgia, California excluding SF/SJM/LA, Kentucky, Louisiana, New Jersey and Georgia excluding ATL/RG. С

Rates for American Indian/Alaska Native (AI/AN) are based on the CHSDA(Contract Health Service Delivery Area) counties.

d Asian/Pacific Islander.

e Hispanic (Hisp) and White Non-Hispanic (W-NHisp) are not mutually exclusive from whites, blacks, Asian/Pacific Islanders, and American Indians/Alaska Natives.

Statistic could not be calculated due to less than 16 cases in the time interval. _

Table 1.22 U.S. Prevalence Counts, Invasive Cancers Only, January 1, 2011ª Using Different Tumor Inclusion Criteriab

		5-Y	ear Limited Dura	ation	36-year Lim	ited Duration
Site	Sex	lst Invasive Tumor Ever ^c	lst Per Site in Previous 36 Years ^d	lst Per Site in Previous 5 Years ^e	lst Invasive Tumor Ever ^c	lst Per Site in Previous 36 Years ^d
All Sites	Both Sexes	4,594,732	4,684,591	5,152,908	12,998,655	13,213,910
	Male	2,373,342	2,409,199	2,638,979	6,188,275	6,261,129
	Female	2,221,390	2,275,392	2,513,929	6,810,380	6,952,781
Oral Cavity & Pharynx	Both Sexes Male Female	104,979 73,503 31,476	121,403 84,311 37,092	125,889 87,116 38,773	272,072 181,087 90,985	300,609 198,337 102,272
Esophagus	Both Sexes	21,165	26,055	26,172	34,428	40,593
	Male	16,662	20,345	20,416	26,846	31,461
	Female	4,503	5,710	5,756	7,582	9,132
Stomach	Both Sexes	36,380	43,398	43,779	72,734	83,348
	Male	21,307	26,023	26,197	41,394	47,992
	Female	15,073	17,375	17,582	31,340	35,356
Colon & Rectum	Both Sexes	403,434	465,899	475,547	1,140,625	1,270,460
	Male	206,233	237,736	242,463	567,872	629,107
	Female	197,201	228,163	233,084	572,753	641,353
Liver &	Both Sexes	33,682	38,427	38,461	45,468	50,961
Intrahepatic	Male	24,703	27,797	27,819	32,312	35,854
Bile Duct	Female	8,979	10,630	10,642	13,156	15,107
Pancreas	Both Sexes	32,756	39,642	39,654	43,238	51,264
	Male	16,110	19,950	19,962	20,679	25,032
	Female	16,646	19,692	19,692	22,559	26,232
Larynx	Both Sexes	31,318	37,292	37,636	87,652	98,482
	Male	25,568	30,391	30,700	70,687	79,142
	Female	5,750	6,901	6,936	16,965	19,340
Lung & Bronchus	Both Sexes	231,126	297,286	306,343	395,186	487,438
	Male	107,276	140,106	143,631	179,129	222,121
	Female	123,850	157,180	162,712	216,057	265,317
Melanoma of the Skin	Both Sexes Male Female	282,028 151,313 130,715	320,618 176,523 144,095	335,149 185,633 149,516	924,397 459,686 464,711	998,829 503,903 494,926
Breast	Female	855,411	924,344	982,776	2,847,146	3,016,451
Cervix	Female	39,578	41,703	41,825	211,309	217,336
Corpus & Uterus, NOS	Female	177,109	198,883	199,045	584,180	637,931
Ovary ^f	Female	58,762	67,726	67,789	173,192	194,691

U.S. 2011 cancer prevalence counts are based on 2011 cancer prevalence proportions from the SEER 9 registries and 1/1/2011 U.S. population estimates based on the average of 2010 and 2011 population estimates from the U.S. Bureau of the Census.

b Prevalence estimates are ambiguous for those with multiple cancers, unless the tumor inclusion criteria are understood. Depending on the application, different inclusion criteria may be appropriate. This table provides three different methods of tumor inclusion:

С d

(c) First invasive tumor ever
 (d) First invasive tumor for each cancer site diagnosed during the previous 36 years (1975-2010)
 (e) First invasive tumor for each cancer site diagnosed during the previous 5 years (2006-2010)
 For definitions (d) and (e) all sites is treated as a separate cancer "site".

Consider a woman who had three invasive cancers: Melanoma in 1981; Breast cancer in 2006; Melanoma in 2007. In method (c) the melanoma is the woman's first cancer, and is counted for the melanoma and all sites 36-year limited duration prevalence. For 5-year limited duration prevalence, the woman is not counted at all since her first cancer occurred more than 5 years prior to 1/1/2011. In method (d) the 1981 melanoma is counted for the melanoma and all sites 36-year limited duration prevalence. The 2006 breast cancer is counted for the breast 5-year and 36-year limited duration prevalence. In method (e) the 2006 breast cancer is counted for the breast cancer and all sites 5-year limited duration prevalence. The 2007 melanoma is counted for 5-year limited duration prevalence for melanoma.

f

а

e

Ovary excludes borderline cases or histologies 8442, 8451, 8462, 8472, and 8473.

Table 1.22 - continued U.S. Prevalence Counts, Invasive Cancers Only, January 1, 2011ª Using Different Tumor Inclusion Criteriab

5-Year Limited Duration

		5-Y	ear Limited Dura	ation	36-year Limited Duration			
Site	Sex	lst Invasive Tumor Ever ^c	lst Per Site in Previous 36 Years ^d	lst Per Site in Previous 5 Years ^e	lst Invasive Tumor Ever ^c	lst Per Site in Previous 36 Years ^d		
Prostate	Male	1,047,674	1,131,608	1,131,634	2,706,583	2,883,365		
Testis	Male	42,114	42,942	43,546	212,079	214,995		
Urinary Bladder	Both Sexes	200,849	251,314	256,779	560,445	653,826		
	Male	152,753	192,335	196,676	418,562	489,001		
	Female	48,096	58,979	60,103	141,883	164,825		
Kidney & Renal Pelvis	Both Sexes Male Female	149,969 91,313 58,656	182,913 113,044 69,869	185,010 114,471 70,539	347,916 206,735 141,181	406,298 243,752 162,546		
Brain & Nervous System	Both Sexes Male Female	44,217 24,099 20,118	47,403 25,776 21,627	47,827 26,028 21,799	131,408 69,809 61,599	135,930 72,101 63,829		
Thyroid	Both Sexes	174,786	194,868	195,449	527,919	564,773		
	Male	38,665	45,067	45,197	114,267	125,269		
	Female	136,121	149,801	150,252	413,652	439,504		
Hodgkin Lymphoma	Both Sexes	38,529	41,043	41,065	172,689	177,526		
	Male	20,912	22,396	22,407	89,550	92,081		
	Female	17,617	18,647	18,658	83,139	85,445		
Non-Hodgkin Lymphoma	Both Sexes Male Female	203,992 109,031 94,961	239,804 129,460 110,344	242,418 130,668 111,750	522,113 274,206 247,907	584,133 307,672 276,461		
Myeloma	Both Sexes	52,505	62,629	62,743	83,118	95,874		
	Male	28,573	34,693	34,761	45,299	52,835		
	Female	23,932	27,936	27,982	37,819	43,039		
Leukemia	Both Sexes	115,823	134,637	135,040	297,129	327,520		
	Male	66,545	78,337	78,566	168,443	186,907		
	Female	49,278	56,300	56,474	128,686	140,613		
Acute	Both Sexes	16,260	16,720	16,720	68,728	69,433		
Lymphocytic	Male	9,051	9,286	9,286	37,729	38,032		
Leukemia	Female	7,209	7,434	7,434	30,999	31,401		
Childhood (Ages 0-19)	Both Sexes Male Female	63,918 33,711 30,207	63,993 33,746 30,247	64,508 33,991 30,517	314,920 162,108 152,812	315,399 162,335 153,064		
Kaposi Sarcoma	Both Sexes	7,194	7,680	7,680	25,504	26,815		
	Male	6,655	7,078	7,078	24,134	25,255		
	Female	539	602	602	1,370	1,560		
Mesothelioma	Both Sexes	3,004	3,922	3,922	4,663	5,734		
	Male	2,172	2,854	2,854	2,829	3,595		
	Female	832	1,068	1,068	1,834	2,139		

U.S. 2011 cancer prevalence counts are based on 2011 cancer prevalence proportions from the SEER 9 registries and 1/1/2011 U.S. population estimates based on the average of 2010 and 2011 population estimates from the U.S. Bureau of the Census.

b

с d е

а

Prevalence estimates are ambiguous for those with multiple cancers, unless the tumor inclusion criteria are understood. Depending on the application, different inclusion criteria may be appropriate. This table provides three different methods of tumor inclusion: (c) First invasive tumor ever (d) First invasive tumor for each cancer site diagnosed during the previous 36 years (1975-2010) (e) First invasive tumor for each cancer site diagnosed during the previous 5 years (2006-2010) For definitions (d) and (e) all sites is treated as a separate gameer "site"

For definitions (d) and (e) all sites is treated as a separate cancer "site"

Consider a woman who had three invasive cancers: Melanoma in 1981; Breast cancer in 2006;

Melanoma in 2007. In method (c) the melanoma is the woman's first cancer, and is counted for the melanoma and all sites 36-year limited duration prevalence. For 5-year limited duration prevalence, the woman is not counted at all since her first cancer occurred more than 5 years prior to 1/1/2011. In method (d) the 1981 melanoma is counted for the melanoma and all sites 36-year limited duration prevalence. The 2006 breast cancer is counted for the breast 5-year and 36-year limited duration prevalence.

In method (e) the 2006 breast cancer is counted for the breast cancer and all sites 5-year limited duration prevalence. The 2007 melanoma is counted for 5-year limited duration prevalence for melanoma.

36-year Limited Duration

Table 1.23 U.S. Complete Prevalence Counts, Invasive Cancers Only, January 1, 2011^a By Age at Prevalence

	Age at Prevalence										
Site/Sex	All Ages ^{c}	0-9	10-19	20-29	30-39	40-49	50-59	60-69	70+		
All Sites											
Males	6,271,036	19,001	41,846	85,407	155,919	356,211	883,826	1,658,545	3,070,283		
Females	7,126,123	16,447	36,376	88,223	226,162	638,879	1,316,044	1,758,236	3,045,756		
Oral Cavity & Pharynx											
Males	186,491	62	423	1,548	3,544	14,256	45,288	58,097	63,273		
Females	95,100	90	522	1,483	3,689	8,554	18,160	23,692	38,911		
Esophagus											
Males	26,909	0	0	22	144	1,109	4,697	9,747	11,190		
Females	7,642	0	0	11	11	240	1,378	1,832	4,170		
Stomach											
Males	42,127	0	29	62	603	2,502	7,119	11,218	20,593		
Females	31,908	4	23	154	618	2,252	4,650	6,664	17,543		
Colon & Rectum											
Males	575,457	11	69	1,276	5,758	25,233	82,732	139,236	321,141		
Females	586,969	0	67	1,472	5,551	23,983	70,097	112,784	373,015		
Liver & Intrahep											
Males	32,389	556	508	616	547	1,590	10,492	11,364	6,716		
Females	13,553	396	521	359	403	889	3,016	3,463	4,506		
Pancreas											
Males	20,801	23	12	130	276	1,653	4,466	6,626	7,615		
Females	22,737	0	62	212	431	1,446	3,801	6,179	10,604		
Larynx											
Males	71,999	0	0	90	219	1,822	10,381	21,280	38,207		
Females	17,266	0	0	36	135	841	3,336	4,868	8,051		
Lung & Bronchus											
Males	183,215	45	110	405	1,195	5,366	24,255	54,762	97,077		
Females	219,111	23	71	367	1,387	7,548	29,898	59,021	120,797		
Melanoma of the Skin											
Males	471,220	56	594	5,039	17,738	46,603	97,543	130,762	172,885		
Females	489,011	68	878	10,864	34,091	73,459	115,119	113,905	140,628		

а b

С

U.S. 2011 cancer prevalence counts are based on 2011 cancer prevalence proportions from the SEER 9 registries (San Francisco, Connecticut, Detroit, Hawaii, Iowa, New Mexico, Seattle, Utah, and Atlanta) and 1/1/2011 U.S. population estimates based on the average of 2010 and 2011 population estimates from the U.S. Bureau of the Census. Prevalence was calculated using the First Malignant Primary Only for a person. Cases diagnosed more than 36 years ago were estimated using the completeness index method (Capocaccia et. al. 1997, Merrill et. al. 2000).

Due to rounding, the sum of the age specific estimates may not equal the all ages estimate.

Table 1.23 - continued U.S. Complete Prevalence Counts, Invasive Cancers Only, January 1, 2011^a By Age at Prevalence

	Age at Prevalence									
Site/Sex	All Ages $^{\circ}$	0-9	10-19	20-29	30-39	40-49	50-59	60-69	70+	
Breast										
Males	14,871	0	0	22	112	591	1,838	4,277	8,030	
Females	2,899,726	0	47	2,669	35,679	215,380	550,120	795,921	1,299,910	
remares	2,000,120	0	17	2,009	55,075	215,500	550,120	195,921	1,200,010	
Cervix										
Females	249,632	0	57	2,142	15,934	41,969	59,700	57,440	72,38	
Corpus & Uterus, NOS										
Females	610,804	12	12	581	5,712	25,316	89,564	165,807	323,80	
Ovary ^d										
Females	188,867	45	1,063	3,475	6,807	18,458	41,719	49,281	68,021	
Prostate										
Males	2,707,821	34	47	113	332	20,758	235,407	776,168	1,674,962	
								·		
Urinary Bladder										
Males	425,722	57	88	567	2,476	10,749	43,060	102,624	266,10	
Females	145,796	56	46	219	1,015	4,094	13,710	30,955	95,70	
Kidney & Renal Pelvis										
Males	212,703	1,458	2,458	2,630	5,254	18,386	41,683	61,520	79,31	
Females	145,900	1,374	2,649	2,922	4,870	12,287	25,458	35,857	60,48	
Hodgkin Lymphoma										
Males	96,100	186	2,493	9,242	16,547	23,764	22,436	14,185	7,24	
Females	89,693	67	2,083	9,407	16,399	23,306	19,845	11,219	7,36	
Non-Hodgkin Lymphoma										
Males	278,836	842	3,598	7,821	13,627	29,722	54,887	71,106	97,23	
Females	252,083	519	1,758	4,663	8,663	21,422	42,382	60,220	112,45	
Myeloma	45 200	0	<i>c</i>	0.0	400	2 054	0.000	14 015	10.04	
Males	45,388	0	6	99	490	3,274	8,260	14,917	18,34	
Females	37,979	0	0	28	312	2,325	6,791	11,124	17,39	
Leukemia										
Males	170,969	6,686	13,047	12,524	11,179	14,206	23,300	35,001	55,02	
Females	131,831	5,729	10,353	11,342	9,836	10,094	15,793	22,736	45,94	
Acute Lymphocytic Leuk	5									
Males	39,514	5,599	11,159	9,649	6,330	3,974	1,505	836	46	
Females	32,723	4,763	8,785	8,138	5,414	3,188	1,137	816	48	

U.S. 2011 cancer prevalence counts are based on 2011 cancer prevalence proportions from the SEER 9 registries (San Francisco, Connecticut, Detroit, Hawaii, Iowa, New Mexico, Seattle, Utah, and Atlanta) and 1/1/2011 U.S. population estimates based on the average of 2010 and 2011 population estimates from the U.S. Bureau of the Census. Prevalence was calculated using the First Malignant Primary Only for a person. Cases diagnosed more than 36 years ago were estimated using the completeness index method (Capocaccia et. al. 1997, Merrill et. al. 2000).

b С

Due to rounding, the sum of the age specific estimates may not equal the all ages estimate.

а

Age-Adjusted SEER Incidence Rates and Trends for the Top 15 Cancer Sites^a by Race/Ethnicity

Both Sexes

All Rac	ces		Whit	e		Blac	k	
	Rate ^b	APC ^c		Rate ^b	APC ^c		Rate ^b	APC ^c
	2007-2011	2002-2011		2007-2011	2002-2011		2007-2011	2002-2011
All Sites	460.4	-0.6*	All Sites	468.9	-0.6*	All Sites	480.8	-1.0*
Breast	67.1	-0.3	Breast	68.3	-0.5	Prostate ^f	94.8	-2.0*
Prostate ^f	67.0	-1.9*	Prostate ^f	64.2	-2.2*	Breast	70.0	0.3
Lung and Bronchus	60.1	-1.7*	Lung and Bronchus	61.7	-1.6*	Lung and Bronchus	68.0	-1.9*
Colon and Rectum	43.7	-2.8*	Colon and Rectum	42.9	-2.9*	Colon and Rectum	53.6	-2.7*
Melanoma of the Skin	21.3	1.4*	Melanoma of the Skin	25.2	1.3*	Kidney and Renal Pelvis	17.9	2.7*
Urinary Bladder	20.5	-0.8*	Urinary Bladder	22.4	-0.7*	Pancreas	15.6	-0.2
Non-Hodgkin Lymphoma	19.7	-0.1	Non-Hodgkin Lymphoma	20.6	-0.2	Non-Hodgkin Lymphoma	14.3	-0.3
Kidney and Renal Pelvis	15.5	1.9*	Kidney and Renal Pelvis	15.9	1.9*	Corpus and Uterus, NOS ^f	13.3	2.3*
Corpus and Uterus, NOS ^f	13.1	1.0*	Leukemia	13.7	0.0	Urinary Bladder	12.6	-0.5
Leukemia	13.0	0.0	Thyroid	13.7	5.8*	Myeloma	12.2	0.4
Thyroid	12.9	5.8*	Corpus and Uterus, NOS ^f	13.4	0.8*	Stomach	11.2	-2.1*
Pancreas	12.3	0.7*	Pancreas	12.2	0.9*	Leukemia	10.0	-1.0
Oral Cavity and Pharynx	11.0	0.4*	Oral Cavity and Pharynx	11.4	0.8*	Liver & IBD ^g	9.4	3.4*
Liver & IBD ^g	7.9	3.4*	Brain and ONS ^g	7.1	-0.5	Oral Cavity and Pharynx	9.4	-2.5*
Stomach	7.5	-1.3*	Liver & IBD ^g	7.0	4.1*	Thyroid	7.6	5.9*
Asian/Pacific	Islander		American Indian/Alaska Native ^d			Hispanic ^e		
	Rate ^b	APC ^c		Rate ^b	APC ^c		Rate ^b	APC ^c
	2007-2011	2002-2011		2007-2011	2002-2011		2007-2011	2002-2011
All Sites	306.7	-0.9*	All Sites	319.3	-0.8	All Sites	353.2	-0.9*
Breast	51.6	0.6	Breast	42.8	0.1	Prostate ^f	53.6	-3.0*
Lung and Bronchus	37.0	-1.4*	Lung and Bronchus	40.9	-2.7*	Breast	49.1	-0.2
Colon and Rectum	36.9	-2.2*	Colon and Rectum	40.0	-1.7*	Colon and Rectum	36.6	-1.9*
Prostate ^f	34.5	-3.6*	Prostate ^f	31.6	-3.6*	Lung and Bronchus	31.3	-2.1*
Liver & IBD ^g	14.1	-0.9	Kidney and Renal Pelvis	19.4	1.6	Non-Hodgkin Lymphoma	17.7	0.3
Non-Hodgkin Lymphoma	13.4	0.0	Liver & IBD ^g	13.7	3.9	Kidney and Renal Pelvis	15.2	1.7*
Thyroid	12.5	5.5*	Non-Hodgkin Lymphoma	12.2	1.8	Liver & IBD ^g	12.2	2.8*
Stomach	11.5	-3.7*	Pancreas	10.7	0.6	Urinary Bladder	11.4	-1.3*
Corpus and Uterus, NOS ^f	10.5	2.5*	Corpus and Uterus, NOS ^f	10.6	2.2	Pancreas	11.1	-0.1
Pancreas	9.7	0.9	Stomach	9.6	-4.0*	Stomach	11.1	-2.0*
Urinary Bladder	8.8	-1.6*	Urinary Bladder	8.1	0.7	Thyroid	10.9	5.3*
Kidney and Renal Pelvis	8.1	2.3*	Oral Cavity and Pharynx	8.1	3.0	Corpus and Uterus, NOS ^f	10.6	1.7*
Leukemia	7.6	-0.2	Thyroid	7.7	4.2*	Leukemia	10.3	0.1
Oral Cavity and Pharynx	7.5	-1.1	Leukemia	7.3	0.1	Oral Cavity and Pharynx	6.7	1.2*
Ovary ^{fh}	5.1	-1.5*	Ovary ^{fh}	5.4	-2.1	Ovary ^{fh}	5.9	-2.1*

Source: SEER 18 areas (San Francisco, Connecticut, Detroit, Hawaii, Iowa, New Mexico, Seattle, Utah, Atlanta, San Jose-Monterey, Los Angeles, Alaska Native Registry, Rural Georgia, California excluding SF/SJM/LA, Kentucky, Louisiana, New Jersey and Georgia excluding ATL/RG).

- а Top 15 cancer sites selected based on 2007-2011 age-adjusted rates for the race/ethnic group. b
- Rates are per 100,000 and age-adjusted to the 2000 US Std Population (19 age groups Census P25-1130). с
- The APC is the Annual Percent Change over the time interval.
- Trends are based on rates age-adjusted to the 2000 US Std Population (19 age groups Census P25-1130). d
- Rates for American Indian/Alaska Native are based on the CHSDA(Contract Health Service Delivery Area) counties.
- е Hispanic is not mutually exclusive from whites, blacks, Asian/Pacific Islanders, and American Indians/Alaska Natives.
- Incidence data for Hispanics are based on NHIA and exclude cases from the Alaska Native Registry. f
- The rates for sex-specific cancer sites are calculated using the population for both sexes combined. q
- IBD = Intrahepatic Bile Duct. ONS = Other Nervous System. h
- Ovary excludes borderline cases or histologies 8442, 8451, 8462, 8472, and 8473.
- The APC is significantly different from zero (p<.05).
- Statistic not shown. Rate based on less than 16 cases for the time interval. Trend based on less than 10 cases for at least one year within the time interval.

Age-Adjusted SEER Incidence Rates and Trends for the Top 15 Cancer Sites^a by Race/Ethnicity

Males

$ \begin{array}{c c c c c c c c c c c c c c c c c c c $	APC ^c <u>2002-2011</u> -1.9* -2.5* -2.8* 2.1*
All Sites 529.4 -1.2* All Sites 532.1 -1.2* All Sites 600.9 Prostate 147.8 -2.3* Prostate 139.9 -2.6* Prostate 223.9 Lung and Bronchus 72.2 -2.4* Lung and Bronchus 72.4 -2.3* Lung and Bronchus 93.0	-1.9* -2.5* -2.8* -2.8*
Prostate 147.8 -2.3* Prostate 139.9 -2.6* Prostate 223.9 Lung and Bronchus 72.2 -2.4* Lung and Bronchus 72.4 -2.3* Lung and Bronchus 93.0	-2.5* -2.8* -2.8*
Lung and Bronchus 72.2 -2.4* Lung and Bronchus 72.4 -2.3* Lung and Bronchus 93.0	-2.8* -2.8*
	-2.8*
Colon and Rectum 50.6 -3.0* Colon and Rectum 49.6 -3.2* Colon and Rectum 62.3	
	2 1*
Urinary Bladder 36.2 -0.8* Urinary Bladder 39.4 -0.8* Kidney and Renal Pelvis 24.7	2.1"
Melanoma of the Skin 27.7 1.6* Melanoma of the Skin 32.3 1.6* Urinary Bladder 21.3	0.1
Non-Hodgkin Lymphoma 23.9 0.0 Non-Hodgkin Lymphoma 24.9 -0.1 Non-Hodgkin Lymphoma 17.4	-0.5
Kidney and Renal Pelvis 21.2 1.8* Kidney and Renal Pelvis 21.7 1.8* Pancreas 17.2	0.3
Leukemia 16.7 -0.3 Leukemia 17.5 -0.3 Liver & IBD ^f 15.6	3.6*
Oral Cavity and Pharynx 16.5 0.4* Oral Cavity and Pharynx 17.0 0.8* Stomach 15.3	-3.1*
Pancreas 14.0 0.9* Pancreas 14.0 1.0* Myeloma 14.8	-0.3
Liver & IBD ^f 12.4 3.7* Liver & IBD ^f 10.8 4.3* Oral Cavity and Pharynx 14.7	-2.9*
Stomach 10.3 -1.6* Stomach 9.2 -1.2* Leukemia 12.9	-1.3
Myeloma 7.7 0.7* Brain and ONS ^f 8.4 -0.5 Larynx 9.0	-3.5*
Esophagus 7.7 -0.7 Esophagus 8.0 -0.1 Esophagus 7.9	-4.8*
Brain and ONS^{f} 7.6 -0.5* Myeloma 7.2 0.7* Brain and ONS^{f} 4.7	-0.5
Asian/Pacific Islander American Indian/Alaska Native ^d Hispanic ^e	
Rate ^b APC ^c Rate ^b APC ^c Rate ^b	APC ^c
<u>2007-2011</u> <u>2002-2011</u> <u>2007-2011</u> <u>2002-2011</u> <u>2007-201</u>	L 2002-2011
All Sites 331.0 -1.8* All Sites 348.1 -0.7 All Sites 405.0	-1.6*
Prostate 79.3 -3.8* Prostate 71.5 -3.9* Prostate 121.8	-3.4*
Lung and Bronchus 49.4 -2.2* Lung and Bronchus 49.5 -2.4 Colon and Rectum 44.3	-2.0*
Colon and Rectum 43.1 -2.4* Colon and Rectum 45.5 0.2 Lung and Bronchus 39.6	-3.1*
Liver & IBD ^f 21.6 -0.7 Kidney and Renal Pelvis 25.3 2.4 Non-Hodgkin Lymphoma 20.6	0.4
Non-Hodgkin Lymphoma 16.3 0.0 Liver & IBD ^f 20.7 5.1 Kidney and Renal Pelvis 20.2	1.4
Urinary Bladder 15.5 -2.0* Urinary Bladder 15.4 - Urinary Bladder 20.0	-1.1*
Stomach 14.9 -4.1* Non-Hodgkin Lymphoma 14.1 1.4 Liver & IBD ^f 18.7	3.2*
Kidney and Renal Pelvis 11.5 2.6* Stomach 12.9 -4.3* Stomach 14.8	-1.7*
Oral Cavity and Pharynx 10.9 -0.3 Pancreas 12.6 - Leukemia 12.4	-0.3
Pancreas 10.7 0.8 Oral Cavity and Pharynx 12.5 4.1 Pancreas 12.2	1.0
Leukemia 9.4 0.1 Leukemia 8.6 1.7 Oral Cavity and Pharynx 9.8	0.9
Thyroid 5.7 5.7* Esophagus 4.8 - Myeloma 6.9	-0.6
Myeloma 4.5 1.8^{\star} Testis 4.5 - Brain and ONS^{f} 5.8	-1.4*
Brain and ONS ^f 4.2 0.8 Larynx 4.3 - Esophagus 5.2	-1.1
Esophagus 3.7 -1.2 Myeloma 4.3 - Testis 4.8	3.0*

- Source: SEER 18 areas (San Francisco, Connecticut, Detroit, Hawaii, Iowa, New Mexico, Seattle, Utah, Atlanta, San Jose-Monterey, Los Angeles, Alaska Native Registry, Rural Georgia, California excluding SF/SJM/LA, Kentucky, Louisiana, New Jersey and Georgia excluding ATL/RG). а
- Top 15 cancer sites selected based on 2007-2011 age-adjusted rates for the race/ethnic group.
- b Rates are per 100,000 and age-adjusted to the 2000 US Std Population (19 age groups - Census P25-1130).
- С The APC is the Annual Percent Change over the time interval.
- Trends are based on rates age-adjusted to the 2000 US Std Population (19 age groups Census P25-1130). d
 - Rates for American Indian/Alaska Native are based on the CHSDA(Contract Health Service Delivery Area) counties.
- P Hispanic is not mutually exclusive from whites, blacks, Asian/Pacific Islanders, and American Indians/Alaska Natives. Incidence data for Hispanics are based on NHIA and exclude cases from the Alaska Native Registry. f
- IBD = Intrahepatic Bile Duct. ONS = Other Nervous System.
- The APC is significantly different from zero (p<.05).
- Statistic not shown. Rate based on less than 16 cases for the time interval. Trend based on less than 10 cases for at least one year within the time interval.

Age-Adjusted SEER Incidence Rates and Trends for the Top 15 Cancer Sites^a by Race/Ethnicity

Females

All Rad	ces		Whit	e		Blac	k	
	Rate ^b	APC ^c		Rate ^b	APC ^c		Rate ^b	APC ^c
	2007-2011	2002-2011		2007-2011	2002-2011		2007-2011	2002-2011
All Sites	411.3	-0.2	All Sites	424.4	-0.2	All Sites	398.8	-0.2
Breast	124.6	-0.2	Breast	128.0	-0.3	Breast	122.8	0.5
Lung and Bronchus	51.1	-1.0*	Lung and Bronchus	53.8	-0.9*	Lung and Bronchus	51.2	-0.7
Colon and Rectum	38.2	-2.6*	Colon and Rectum	37.3	-2.7*	Colon and Rectum	47.5	-2.7*
Corpus and Uterus, NOS	24.6	1.2*	Corpus and Uterus, NOS	25.4	1.1*	Corpus and Uterus, NOS	23.2	2.5*
Thyroid	19.1	6.0*	Thyroid	20.4	6.0*	Pancreas	14.2	-0.5
Melanoma of the Skin	16.7	0.9	Melanoma of the Skin	20.0	0.9	Kidney and Renal Pelvis	12.7	3.2*
Non-Hodgkin Lymphoma	16.3	-0.3	Non-Hodgkin Lymphoma	17.2	-0.3	Non-Hodgkin Lymphoma	11.9	-0.2
Ovary ^g	12.3	-1.7*	Ovary ^g	13.0	-1.6*	Thyroid	11.3	6.3*
Pancreas	10.9	0.6*	Kidney and Renal Pelvis	11.0	1.7*	Myeloma	10.5	0.8
Kidney and Renal Pelvis	10.7	1.8*	Pancreas	10.7	0.7*	Ovary ^g	9.8	-1.4*
Leukemia	10.2	0.1	Leukemia	10.7	0.2	Cervix Uteri	9.4	-3.3*
Urinary Bladder	8.8	-1.2*	Urinary Bladder	9.5	-1.1*	Stomach	8.5	-1.1
Cervix Uteri	7.8	-1.6*	Cervix Uteri	7.8	-1.2*	Leukemia	8.0	-0.8
Oral Cavity and Pharynx	6.2	0.1	Oral Cavity and Pharynx	6.4	0.4	Urinary Bladder	6.9	-1.8*
Brain and ONS ^f	5.4	-0.7*	Brain and ONS ^f	6.0	-0.7*	Oral Cavity and Pharynx	5.3	-1.7
Asian/Pacific	: Islander		American Indian/Alaska Native ^d			Hispanic ^e		
	Rate ^b	APC ^c		Rate ^b	APC ^c		Rate ^b	APC ^c
	2007-2011	2002-2011		2007-2011	2002-2011		2007-2011	2002-2011
All Sites	293.0	0.0	All Sites	301.9	-0.6	All Sites	320.4	-0.3*
Breast	93.6	0.6	Breast	79.3	0.2	Breast	91.3	0.0
Colon and Rectum	32.0	-2.1*	Colon and Rectum	35.5	-3.5*	Colon and Rectum	30.6	-1.8*
Lung and Bronchus	28.1	-0.2	Lung and Bronchus	34.7	-2.8	Lung and Bronchus	25.5	-0.9
Corpus and Uterus, NOS	19.1	2.5*	Corpus and Uterus, NOS	20.0	2.6	Corpus and Uterus, NOS	19.8	1.9*
Thyroid	18.5	5.3*	Kidney and Renal Pelvis	14.5	0.8	Thyroid	17.1	5.4*
Non-Hodgkin Lymphoma	11.2	-0.2	Thyroid	11.8	4.7*	Non-Hodgkin Lymphoma	15.4	0.2
Ovary ^g	9.3	-1.5*	Non-Hodgkin Lymphoma	10.6	1.3	Kidney and Renal Pelvis	11.2	2.0*
Stomach	9.0	-3.2*	Ovary ^g	9.8	-1.9	Ovary ^g	10.9	-1.8*
Pancreas	8.9	1.0	Pancreas	9.1	-0.4	Cervix Uteri	10.2	-4.5*
Liver & IBD ^f	8.1	-1.3	Liver & IBD ^f	8.0	1.9	Pancreas	10.2	-1.0
Cervix Uteri	6.4	-3.5*	Cervix Uteri	7.6	0.1	Leukemia	8.7	0.4
Leukemia	6.1	-0.6	Stomach	7.3	-3.6	Stomach	8.3	-2.5*
Kidney and Renal Pelvis	5.5	1.8	Leukemia	6.5	-0.5	Liver & IBD ^f	6.7	1.7
Oral Cavity and Pharynx	4.7	-2.4*	Oral Cavity and Pharynx	4.4	-	Urinary Bladder	5.1	-1.9*
Urinary Bladder	3.9	-0.5	Melanoma of the Skin	4.0	-	Myeloma	4.7	-0.6

Source: SEER 18 areas (San Francisco, Connecticut, Detroit, Hawaii, Iowa, New Mexico, Seattle, Utah, Atlanta, San Jose-Monterey, Los Angeles, Alaska Native Registry, Rural Georgia, California excluding SF/SJM/LA, Kentucky, Louisiana, New Jersey and Georgia excluding ATL/RG).

Top 15 cancer sites selected based on 2007-2011 age-adjusted rates for the race/ethnic group.

- ^b Rates are per 100,000 and age-adjusted to the 2000 US Std Population (19 age groups Census P25-1130).
- ^c The APC is the Annual Percent Change over the time interval.
- Trends are based on rates age-adjusted to the 2000 US Std Population (19 age groups Census P25-1130).
- ^d Rates for American Indian/Alaska Native are based on the CHSDA(Contract Health Service Delivery Area) counties.
- ^e Hispanic is not mutually exclusive from whites, blacks, Asian/Pacific Islanders, and American Indians/Alaska Natives.
- Incidence data for Hispanics are based on NHIA and exclude cases from the Alaska Native Registry.
- ^f IBD = Intrahepatic Bile Duct. ONS = Other Nervous System.
- ^g Ovary excludes borderline cases or histologies 8442, 8451, 8462, 8472, and 8473.
- The APC is significantly different from zero (p<.05).
- Statistic not shown. Rate based on less than 16 cases for the time interval. Trend based on less than 10 cases for at least one year within the time interval.

Age-Adjusted U.S. Death Rates and Trends for the Top 15 Cancer Sites^a by Race/Ethnicity

Both Sexes

All Races			Whit	е		Blac	k	
	Rate ^b	APC ^c		Rate ^b	APC ^c		Rate ^b	APC ^c
200	7-2011	2002-2011		2007-2011	2002-2011		2007-2011	2002-2011
All Sites 1	L73.8	-1.5*	All Sites	173.3	-1.4*	All Sites	206.4	-2.1*
Lung and Bronchus	48.4	-1.9*	Lung and Bronchus	49.1	-1.8*	Lung and Bronchus	52.0	-2.6*
Colon and Rectum	15.9	-2.9*	Colon and Rectum	15.5	-2.9*	Colon and Rectum	22.1	-2.8*
Breast	12.4	-2.1*	Breast	12.1	-2.2*	Breast	18.0	-1.6*
Pancreas	10.9	0.4*	Pancreas	10.8	0.5*	Prostate ^f	17.4	-3.3*
Prostate ^f	8.8	-2.6*	Prostate ^f	8.1	-2.5*	Pancreas	13.6	-0.3
Leukemia	7.0	-0.9*	Leukemia	7.3	-0.8*	Liver & IBD ^g	7.6	2.5*
Non-Hodgkin Lymphoma	6.3	-2.6*	Non-Hodgkin Lymphoma	6.6	-2.6*	Stomach	6.5	-3.3*
Liver & IBD ^g	5.8	2.5*	Liver & IBD ^g	5.3	2.6*	Myeloma	6.3	-1.6*
Urinary Bladder	4.4	0.1	Brain and ONS ^g	4.6	-0.2	Leukemia	6.0	-1.5*
Ovary ^f	4.4	-2.2*	Urinary Bladder	4.6	0.3*	Corpus and Uterus, NOS ^f	4.5	0.8
Brain and ONS ^g	4.3	-0.4	Ovary ^f	4.5	-2.2*	Non-Hodgkin Lymphoma	4.5	-2.0*
Esophagus	4.2	-0.7*	Esophagus	4.3	-0.1	Esophagus	4.2	-4.6*
Kidney and Renal Pelvis	4.0	-0.9*	Kidney and Renal Pelvis	4.0	-0.8*	Ovary ^f	3.9	-1.9*
Stomach	3.5	-2.9*	Melanoma of the Skin	3.1	0.6*	Kidney and Renal Pelvis	3.9	-1.0*
Myeloma	3.4	-1.5*	Myeloma	3.1	-1.5*	Urinary Bladder	3.6	-0.9
Asian/Pacific Is	lander		American Indian/	Alaska Nati	ve ^d	Hispan	ic ^e	
	Rate ^b	APC ^c		Rate ^b	APC ^c		Rate ^b	APC ^c
200	7-2011	2002-2011		2007-2011	2002-2011		2007-2011	2002-2011
All Sites 1	L07.8	-1.1*	All Sites	158.0	-1.0*	All Sites	120.3	-1.3*
Lung and Bronchus	25.2	-1.1*	Lung and Bronchus	39.9	-1.0	Lung and Bronchus	20.9	-2.3*
Colon and Rectum	11.0	-1.6*	Colon and Rectum	17.2	-0.6	Colon and Rectum	12.4	-1.7*
Liver & IBD ^g	9.8	-1.2*	Liver & IBD ^g	9.5	2.0	Liver & IBD ^g	8.7	1.5*
Pancreas	7.7	0.5	Pancreas	8.8	1.0	Pancreas	8.6	0.3
Breast	6.4	-1.5*	Breast	8.6	-3.0*	Breast	8.1	-1.6*
Stomach	6.3	-3.7*	Prostate ^f	8.4	-0.9	Prostate ^f	7.3	-2.4*
Non-Hodgkin Lymphoma	4.1	-1.9*	Kidney and Renal Pelvis	6.7	-0.5	Stomach	5.6	-2.9*
Prostate ^f	4.0	-2.6*	Stomach	5.2	-5.5*	Non-Hodgkin Lymphoma	5.3	-1.4*
Leukemia	4.0	0.8	Leukemia	4.7	-1.3	Leukemia	4.8	-0.8*
Ovary ^f	2.6	-1.0	Non-Hodgkin Lymphoma	4.6	-1.4	Kidney and Renal Pelvis	3.6	-1.0
Kidney and Renal Pelvis	2.0	2.2	Ovary ^f	3.9	-0.3	Ovary ^f	3.2	-1.7*
Oral Cavity and Pharynx	1.9	-2.1*	Esophagus	3.3	-3.1*	Myeloma	2.8	-1.4
Brain and ONS ^g	1.9	-0.5	Myeloma	2.7	-5.6*	Brain and ONS ^g	2.8	-0.4
Myeloma	1.8	0.6	Brain and ONS ^g	2.6	2.1	Urinary Bladder	2.4	-1.2*
Esophagus	1.7	-1.3	Urinary Bladder	2.5	3.4	Esophagus	2.3	0.0

- а Top 15 cancer sites selected based on 2007-2011 age-adjusted rates for the race/ethnic group.
- b Rates are per 100,000 and age-adjusted to the 2000 US Std Population (19 age groups - Census P25-1130). С
 - The APC is the Annual Percent Change over the time interval.
- Trends are based on rates age-adjusted to the 2000 US Std Population (19 age groups Census P25-1130).
- d Rates for American Indian/Alaska Native are based on the CHSDA(Contract Health Service Delivery Area) counties. ρ
- Hispanic is not mutually exclusive from whites, blacks, Asian/Pacific Islanders, and American Indians/Alaska Natives. f
- The rates for sex-specific cancer sites are calculated using the population for both sexes combined. g
 - IBD = Intrahepatic Bile Duct. ONS = Other Nervous System.
 - The APC is significantly different from zero (p<.05).
 - Statistic not shown. Rate based on less than 16 cases for the time interval. Trend based on less than 10 cases for at least one year within the time interval.

Age-Adjusted U.S. Death Rates and Trends for the Top 15 Cancer Sites^a by Race/Ethnicity

Males

All sites211.6-1.8*All Sites209.8-1.7*All Sites269.3-2.6*Lung and Bronchus61.6-2.6*Lung and Bronchus61.4-2.5*Lung and Bronchus75.7-3.8*Prostate22.3-3.3*Prostate20.6-3.3*Prostate48.9-3.8*Colon and Rectum19.1-3.0*Colon and Rectum18.5-3.1*Colon and Rectum27.7-2.6*Pancreas12.50.5*Pancreas15.3-0.3-0.3*Pancreas15.3-0.3Leukemia9.4-1.0*Leukemia9.7-0.9*Liver & IBD ⁶ 12.12.7*Non-Hodgkin Lymphoma8.1-2.3*Urinary Bladder8.10.2Leukemia8.0-1.5*Non-Hodgkin Lymphoma8.1-2.4*Stomach8.0-1.5*Esophagus7.4-4.3*Kidney and Renal Pelvis5.8-0.6*Esophagus7.4-4.3*Stomach4.1-3.3*Oral Cavity and Panrynx5.1-0.7Melanoma of the Skin4.10.7*Myeloma4.0-1.3*Cavity and Panrynx5.4-0.7-0.7Melanoma of the Skin4.10.7*Myeloma4.0-1.3*Cavity and Panrynx5.1-3.7*-0.4Brain and ONS ⁴ 5.9-0.8*Non-Hodgkin Lymphoma5.4-0.7-3.3*Cavity and Panrynx5.4-0.7Melanoma of the Skin4.10.3*-0.2Kidney an	All Ra	ces		Whit	e		Blac	k		
$ \begin{array}{c c c c c c c c c c c c c c c c c c c $										
$ \begin{array}{c c c c c c c c c c c c c c c c c c c $									2002-2011	
$ \begin{array}{c ccccccccccccccccccccccccccccccccccc$										
$ \begin{array}{cccccccccccccccccccccccccccccccccccc$							5			
$\begin{array}{c c c c c c c c c c c c c c c c c c c $	Prostate									
$ \begin{array}{c ccccccccccccccccccccccccccccccccccc$	Colon and Rectum			Colon and Rectum			Colon and Rectum		-2.6*	
$ \begin{array}{c ccccccccccccccccccccccccccccccccccc$	Pancreas	12.5		Pancreas			Pancreas	15.3		
$\begin{array}{c ccccccccccccccccccccccccccccccccccc$	Leukemia	9.4	-1.0*	Leukemia	9.7	-0.9*	Liver & IBD ^f			
Urinary Bladder 7.7 0.1 Liver & IBD ^f 7.8 2.8* Myeloma 7.7 -1.3* Esophagus 7.5 -0.6* Esophagus 7.8 -0.1 Esophagus 7.4 -4.3* Kidney and Renal Pelvis 5.8 -0.8* Kidney and Renal Pelvis 5.9 -0.8* Non-Hodgkin Lymphoma 5.8 -1.3 Brain and ONS ^f 5.2 -0.4 Brain and ONS ^f 5.6 -0.2 Kidney and Renal Pelvis 5.6 -1.2* Stomach 4.7 -3.2* Melanoma of the Skin 4.6 0.9* Urinary Bladder 5.4 -0.7 Myeloma 4.3 -1.3* Stomach 4.1 -3.3* Oral Cavity and Pharynx 5.1 -3.7* Melanoma of the Skin 4.1 0.7* Myeloma 4.0 -1.3* Larynx 3.8 -3.9* Oral Cavity and Pharynx 3.8 -0.9* Oral Cavity and Pharynx 3.7 -0.4 Brain and ONS ^f 3.0 -1.1 All Sites 131.0 -1.3* All Sites 100.0 -0.5 All Sites 150.1	Liver & IBD ^f	8.5	2.6*		8.4	-2.4*	Stomach	9.6	-3.3*	
Esophagus 7.5 -0.6^* Esophagus 7.8 -0.1 Esophagus 7.4 -4.3^* Kidney and Renal Pelvis 5.8 -0.8^* Kidney and Renal Pelvis 5.9 -0.8^* Non-Hodgkin Lymphoma 5.8 -1.2^* Brain and ONS ^f 5.2 -0.4 Brain and ONS ^f 5.6 -1.2^* Non-Hodgkin Lymphoma 5.6 -1.2^* Stomach 4.7 -3.2^* Melanoma of the Skin 4.6 0.9^* Urinary Bladder 5.4 -0.7 Myeloma 4.3 -1.3^* Stomach 4.1 -3.3^* Oral Cavity and Pharynx 5.8 -3.9^* Oral Cavity and Pharynx 3.8 -0.9^* Oral Cavity and Pharynx 3.7 -0.4 Brain and ONS ^f 3.0 -1.1 Asian/Pacific Islander American Indian/Alaska Native ^d Hispanic ^e Hispanic ^e $2007-2011$ $2002-2011$ $2002-2011$ $2007-2011$ $2002-201$ All Sites 131.0 -1.3^* All Sites 190.0 -0.5 All Sites 150.1 -1.6^* Liver & IBD ^f 1	Non-Hodgkin Lymphoma	8.1	-2.3*	Urinary Bladder	8.1	0.2	Leukemia	8.0	-1.5*	
Kidney and Renal Pelvis 5.8 -0.8* Kidney and Renal Pelvis 5.9 -0.8* Non-Hodgkin Lymphoma 5.8 -1.3 Brain and ONS ^f 5.6 -0.8* Non-Hodgkin Lymphoma 5.8 -1.3 Brain and ONS ^f 5.6 -0.2 Kidney and Renal Pelvis 5.6 -1.3* Stomach 4.7 -3.2* Melanoma of the Skin 4.6 0.9* Urinary Bladder 5.4 -0.7 Myeloma 4.3 -1.3* Stomach 4.1 -3.3* Oral Cavity and Pharynx 3.7 -0.4 Brain and ONS ^f -1.3 Melanoma of the Skin 4.0 -1.3* Melanoma of the Skin 4.0 -1.3* American Indian/Alaska Native ^d Hispanic ^e American Indian/Alaska Native ^d <th colsp<="" td=""><td>Urinary Bladder</td><td>7.7</td><td>0.1</td><td>Liver & IBD^f</td><td>7.8</td><td>2.8*</td><td>Myeloma</td><td>7.7</td><td>-1.3*</td></th>	<td>Urinary Bladder</td> <td>7.7</td> <td>0.1</td> <td>Liver & IBD^f</td> <td>7.8</td> <td>2.8*</td> <td>Myeloma</td> <td>7.7</td> <td>-1.3*</td>	Urinary Bladder	7.7	0.1	Liver & IBD ^f	7.8	2.8*	Myeloma	7.7	-1.3*
$ \begin{array}{c ccccccccccccccccccccccccccccccccccc$	Esophagus	7.5	-0.6*	Esophagus	7.8	-0.1	Esophagus	7.4	-4.3*	
Stomach 4.7 -3.2^* Melanoma of the Skin 4.6 0.9^* Urinary Bladder 5.4 -0.7 Myeloma 4.3 -1.3^* Stomach 4.1 -3.3^* Oral Cavity and Pharynx 5.1 -3.7^* Melanoma of the Skin 4.1 0.7^* Myeloma 4.0 -1.3^* Larynx 3.8 -3.9^* Oral Cavity and Pharynx 3.8 -0.9^* Oral Cavity and Pharynx 3.7 -0.4 Brain and ONS ^f 3.0 -1.1 Asian/Pacific IslanderMelanoma of the Skin 4.0 -1.3^* Larynx 3.8 -3.9^* Oral Cavity and Pharynx 3.7 -0.4 Brain and ONS ^f 3.0 -1.1 Asian/Pacific IslanderMeerican Indian/Alaska Native ^d Hispanic ^e Meerican Indian/Alaska Native ^d Hispanic ^e AdvectorRate ^b APC ^c $2007-2011$ $2002-2011$ $2002-2011$ $2007-2011$ $2002-201$ All Sites 190.0 -0.5 All Sites 150.1 -1.6^* Lung and Bronchus 34.7 -1.7^* Lung and Bronchus 50.0 -0.7 Lung and Bronchus 30.5 -2.9^* Liver & IBD ^f 14.5 -1.0^* Prostate 21.2 -1.4 Colon and Rectum 15.8 -1.5^* Prostate 10.0 -2.4^* Liver & IBD ^f 13.8 5.2^* Liver & IBD ^f 12.6 1.7^* <td>Kidney and Renal Pelvis</td> <td>5.8</td> <td>-0.8*</td> <td>Kidney and Renal Pelvis</td> <td>5.9</td> <td>-0.8*</td> <td>Non-Hodgkin Lymphoma</td> <td>5.8</td> <td>-1.3</td>	Kidney and Renal Pelvis	5.8	-0.8*	Kidney and Renal Pelvis	5.9	-0.8*	Non-Hodgkin Lymphoma	5.8	-1.3	
Myeloma4.3 -1.3^* Stomach4.1 -3.3^* Oral Cavity and Pharynx 5.1 -3.7^* Melanoma of the Skin4.1 0.7^* Myeloma 4.0 -1.3^* Larynx 3.8 -3.9^* Oral Cavity and Pharynx 3.8 -0.9^* Oral Cavity and Pharynx 3.7 -0.4 Brain and ONS ^f 3.0 -1.1 Asian/Pacific IslanderAmerican Indian/Alaska Native ^d Hispanic ^e Agian/Pacific IslanderAmerican Indian/Alaska Native ^d Hispanic ^e All Sites 131.0 -1.3^* All Sites 190.0 -0.5 All Sites 150.1 -1.6^* Lung and Bronchus 34.7 -1.7^* Lung and Bronchus 50.0 -0.7 Lung and Bronchus 30.5 -2.9^* Liver & IBD ^f 14.5 -1.0^* Prostate 21.2 -1.2 Prostate 13.5^* -1.5^* Prostate 10.0 -2.4^* Liver & IBD ^f 13.8 5.2^* Liver & IBD ^f 12.6 1.7^* Pancreas 8.5 0.5 Pancreas 9.9 0.6 Pancreas 9.7 0.7 Stomach 8.3 -3.6^* Kidney and Renal Pelvis 9.5 -0.5 Stomach 7.5 -3.0^*	Brain and ONS ^f	5.2	-0.4	Brain and ONS ^f	5.6	-0.2	Kidney and Renal Pelvis	5.6	-1.2*	
Melanoma of the Skin4.1 0.7^* Oral Cavity and PharynxMyeloma 4.0 -1.3^* Oral Cavity and PharynxLarynx 3.8 -3.9^* Melanoma of the Skin 4.1 0.7^* Oral Cavity and PharynxMyeloma 4.0 -1.3^* Oral Cavity and PharynxLarynx 3.8 -3.9^* Main and ONS ^f 3.0 -1.1 Asian/Pacific IslanderAmerican Indian/Alaska NativedHispaniceMerican Indian/Alaska NativedHispanice2007-2011 2002-201Anerican Indian/Alaska NativedHispaniceLiver & IBD ^f 131.0-1.7*Lung and Bronchus50.0-0.7<	Stomach	4.7	-3.2*	Melanoma of the Skin	4.6	0.9*	Urinary Bladder	5.4	-0.7	
Melanoma of the Skin4.1 0.7^* Oral Cavity and PharynxMyeloma 4.0 -1.3^* Oral Cavity and PharynxLarynx 3.8 -3.9^* Melanoma of the Skin 4.1 0.7^* Oral Cavity and PharynxMyeloma 4.0 -1.3^* Brain and ONS ^f Larynx 3.8 -3.9^* Asian/Pacific IslanderAmerican Indian/Alaska Native ^d Larynx 3.8 -3.9^* Asian/Pacific IslanderAmerican Indian/Alaska Native ^d Hispanic ^e Rate ^b APC ^c Rate ^b APC ^c 2007-2011 $2002-2011$ $2002-2011$ $2002-2011$ All Sites 131.0 -1.3^* All Sites 190.0 -0.5 All Sites $2107-2011$ Liver & IBD ^f 14.5 -1.0^* Prostate 21.2 -1.2 Prostate 18.5 -3.0^* Colon and Rectum 13.1 -1.9^* Colon and Rectum 19.2 -1.4 Colon and Rectum 15.8 -1.5^* Prostate 10.0 -2.4^* Liver & IBD ^f 13.8 5.2^* Liver & IBD ^f 12.6 1.7^* Pancreas 8.5 0.5 Pancreas 9.9 0.6 Pancreas 9.7 0.7 Non-Hodgkin Lymphoma 5.2 -1.8^* Stomach 7.0 -5.5^* Non-Hodgkin Lymphoma 6.4 -1.1^*	Myeloma	4.3	-1.3*	Stomach	4.1	-3.3*	Oral Cavity and Pharynx	5.1	-3.7*	
Asian/Pacific IslanderAmerican Indian/Alaska NativedHispaniceAsian/Pacific IslanderAmerican Indian/Alaska NativedHispaniceRatebAPC°RatebAPC°RatebAPC°2007-20112002-2011 $2002-2011$ $2007-2011$ $2002-2011$ All Sites131.0-1.3*All Sites190.0-0.5All Sites150.1Lung and Bronchus34.7-1.7*Lung and Bronchus50.0-0.7Lung and Bronchus30.5Liver & IBD ^f 14.5-1.0*Prostate21.2-1.2Prostate18.5-3.0*Colon and Rectum13.1-1.9*Colon and Rectum19.2-1.4Colon and Rectum15.8-1.5*Prostate10.0-2.4*Liver & IBD ^f 13.85.2*Liver & IBD ^f 12.61.7*Pancreas8.50.5Pancreas9.90.6Pancreas9.70.7Stomach8.3-3.6*Kidney and Renal Pelvis9.5-0.5Stomach7.5-3.0*Non-Hodgkin Lymphoma5.2-1.8*Stomach7.0-5.5*Non-Hodgkin Lymphoma6.4-1.1*	Melanoma of the Skin	4.1	0.7*	Myeloma	4.0	-1.3*			-3.9*	
$ \begin{array}{c c c c c c c c c c c c c c c c c c c $	Oral Cavity and Pharynx	3.8	-0.9*	Oral Cavity and Pharynx	3.7	-0.4	Brain and ONS^{f}	3.0	-1.1	
$\begin{array}{c c c c c c c c c c c c c c c c c c c $	Asian/Pacific	c Islander		American Indian/Alaska Native ^d			Hispanic ^e			
All Sites 131.0 -1.3^* All Sites 190.0 -0.5 All Sites 150.1 -1.6^* Lung and Bronchus 34.7 -1.7^* Lung and Bronchus 50.0 -0.7 Lung and Bronchus 30.5 -2.9^* Liver & IBD ^f 14.5 -1.0^* Prostate 21.2 -1.2 Prostate 18.5 -3.0^* Colon and Rectum 13.1 -1.9^* Colon and Rectum 19.2 -1.4 Colon and Rectum 15.8 -1.5^* Prostate 10.0 -2.4^* Liver & IBD ^f 13.8 5.2^* Liver & IBD ^f 12.6 1.7^* Pancreas 8.5 0.5 Pancreas 9.9 0.6 Pancreas 9.7 0.7 Stomach 8.3 -3.6^* Kidney and Renal Pelvis 9.5 -0.5 Stomach 7.5 -3.0^* Non-Hodgkin Lymphoma 5.2 -1.8^* Stomach 7.0 -5.5^* Non-Hodgkin Lymphoma 6.4 -1.1^*		Rate ^b	APC ^c		Rate ^b	APC ^c		Rate ^b	APC ^c	
Lung and Bronchus 34.7 $-1.7*$ Lung and Bronchus 50.0 -0.7 Lung and Bronchus 30.5 $-2.9*$ Liver & IBD ^f 14.5 $-1.0*$ Prostate 21.2 -1.2 Prostate 18.5 $-3.0*$ Colon and Rectum 13.1 $-1.9*$ Colon and Rectum 19.2 -1.4 Colon and Rectum 15.8 $-1.5*$ Prostate 10.0 $-2.4*$ Liver & IBD ^f 13.8 $5.2*$ Liver & IBD ^f 12.6 $1.7*$ Pancreas 8.5 0.5 Pancreas 9.9 0.6 Pancreas 9.7 0.7 Stomach 8.3 $-3.6*$ Kidney and Renal Pelvis 9.5 -0.5 Stomach 7.5 $-3.0*$ Non-Hodgkin Lymphoma 5.2 $-1.8*$ Stomach 7.0 $-5.5*$ Non-Hodgkin Lymphoma 6.4 $-1.1*$		2007-2011			2007-2011	2002-2011		2007-2011	2002-2011	
Liver & IBD^f 14.5-1.0*Prostate21.2-1.2Prostate18.5-3.0*Colon and Rectum13.1-1.9*Colon and Rectum19.2-1.4Colon and Rectum15.8-1.5*Prostate10.0-2.4*Liver & IBD^f13.85.2*Liver & IBD^f12.61.7*Pancreas8.50.5Pancreas9.90.6Pancreas9.70.7Stomach8.3-3.6*Kidney and Renal Pelvis9.5-0.5Stomach7.5-3.0*Non-Hodgkin Lymphoma5.2-1.8*Stomach7.0-5.5*Non-Hodgkin Lymphoma6.4-1.1*	All Sites	131.0	-1.3*	All Sites	190.0	-0.5	All Sites	150.1	-1.6*	
Colon and Rectum 13.1 -1.9* Colon and Rectum 19.2 -1.4 Colon and Rectum 15.8 -1.5* Prostate 10.0 -2.4* Liver & IBD ^f 13.8 5.2* Liver & IBD ^f 12.6 1.7* Pancreas 8.5 0.5 Pancreas 9.9 0.6 Pancreas 9.7 0.7 Stomach 8.3 -3.6* Kidney and Renal Pelvis 9.5 -0.5 Stomach 7.5 -3.0* Non-Hodgkin Lymphoma 5.2 -1.8* Stomach 7.0 -5.5* Non-Hodgkin Lymphoma 6.4 -1.1*	Lung and Bronchus	34.7	-1.7*	Lung and Bronchus	50.0	-0.7	Lung and Bronchus	30.5	-2.9*	
Prostate 10.0 -2.4* Liver & IBD ^f 13.8 5.2* Liver & IBD ^f 12.6 1.7* Pancreas 8.5 0.5 Pancreas 9.9 0.6 Pancreas 9.7 0.7 Stomach 8.3 -3.6* Kidney and Renal Pelvis 9.5 -0.5 Stomach 7.5 -3.0* Non-Hodgkin Lymphoma 5.2 -1.8* Stomach 7.0 -5.5* Non-Hodgkin Lymphoma 6.4 -1.1*	Liver & IBD ^f	14.5	-1.0*	Prostate	21.2	-1.2	Prostate	18.5	-3.0*	
Pancreas 8.5 0.5 Pancreas 9.9 0.6 Pancreas 9.7 0.7 Stomach 8.3 -3.6* Kidney and Renal Pelvis 9.5 -0.5 Stomach 7.5 -3.0* Non-Hodgkin Lymphoma 5.2 -1.8* Stomach 7.0 -5.5* Non-Hodgkin Lymphoma 6.4 -1.1*	Colon and Rectum	13.1	-1.9*	Colon and Rectum	19.2	-1.4	Colon and Rectum	15.8	-1.5*	
Stomach 8.3 -3.6* Kidney and Renal Pelvis 9.5 -0.5 Stomach 7.5 -3.0* Non-Hodgkin Lymphoma 5.2 -1.8* Stomach 7.0 -5.5* Non-Hodgkin Lymphoma 6.4 -1.1*	Prostate	10.0	-2.4*	Liver & IBD ^f	13.8	5.2*	Liver & IBD ^f	12.6	1.7*	
Non-Hodgkin Lymphoma 5.2 -1.8* Stomach 7.0 -5.5* Non-Hodgkin Lymphoma 6.4 -1.1*	Pancreas	8.5	0.5	Pancreas	9.9	0.6	Pancreas	9.7	0.7	
	Stomach	8.3	-3.6*	Kidney and Renal Pelvis	9.5	-0.5	Stomach	7.5	-3.0*	
	Non-Hodgkin Lymphoma	5.2	-1.8*	Stomach	7.0	-5.5*	Non-Hodgkin Lymphoma	6.4	-1.1*	
Leukemia 5.0 0.4 Leukemia 6.6 1.4 Leukemia 6.0 -0.9	Leukemia	5.0	0.4	Leukemia	6.6	1.4	Leukemia	6.0	-0.9	
Esophagus 3.0 -1.0 Esophagus 5.7 -4.4 Kidney and Renal Pelvis 5.1 -1.4	Esophaqus	3.0	-1.0	Esophaqus	5.7	-4.4	Kidney and Renal Pelvis	5.1	-1.4	
Kidney and Renal Pelvis 3.0 2.6 Non-Hodgkin Lymphoma 5.3 0.9 Esophagus 4.3 0.4	Kidney and Renal Pelvis	3.0	2.6	Non-Hodgkin Lymphoma	5.3	0.9	Esophagus	4.3	0.4	
Oral Cavity and Pharynx 2.9 -1.9* Urinary Bladder 4.4 - Urinary Bladder 4.0 -1.4	Oral Cavity and Pharynx	2.9	-1.9*	Urinary Bladder	4.4	-	Urinary Bladder	4.0	-1.4	
Urinary Bladder 2.9 -0.1 Myeloma 3.4 -5.9* Myeloma 3.5 -0.8	Urinary Bladder	2.9	-0.1	Myeloma	3.4	-5.9*	Myeloma	3.5	-0.8	
Brain and ONS ^f 2.3 -1.8 Oral Cavity and Pharynx 3.4 0.3 Brain and ONS ^f 3.3 -0.3		2.3	-1.8		3.4	0.3		3.3	-0.3	
Myeloma 2.3 2.5 Brain and ONS ^f 2.9 0.6 Oral Cavity and Pharynx 2.4 -1.9*										
Soft Tissue including Heart 1.0 3.2 Larynx 1.8 - Larynx 1.7 -2.3*	1									

- ^a Top 15 cancer sites selected based on 2007-2011 age-adjusted rates for the race/ethnic group.
- Rates are per 100,000 and age-adjusted to the 2000 US Std Population (19 age groups Census P25-1130).
- ^c The APC is the Annual Percent Change over the time interval.
- Trends are based on rates age-adjusted to the 2000 US Std Population (19 age groups Census P25-1130).
- d Rates for American Indian/Alaska Native are based on the CHSDA(Contract Health Service Delivery Area) counties.
- Hispanic is not mutually exclusive from whites, blacks, Asian/Pacific Islanders, and American Indians/Alaska Natives.
 - IBD = Intrahepatic Bile Duct. ONS = Other Nervous System.
 - The APC is significantly different from zero (p<.05).
 - Statistic not shown. Rate based on less than 16 cases for the time interval. Trend based on less than 10 cases for at least one year within the time interval.

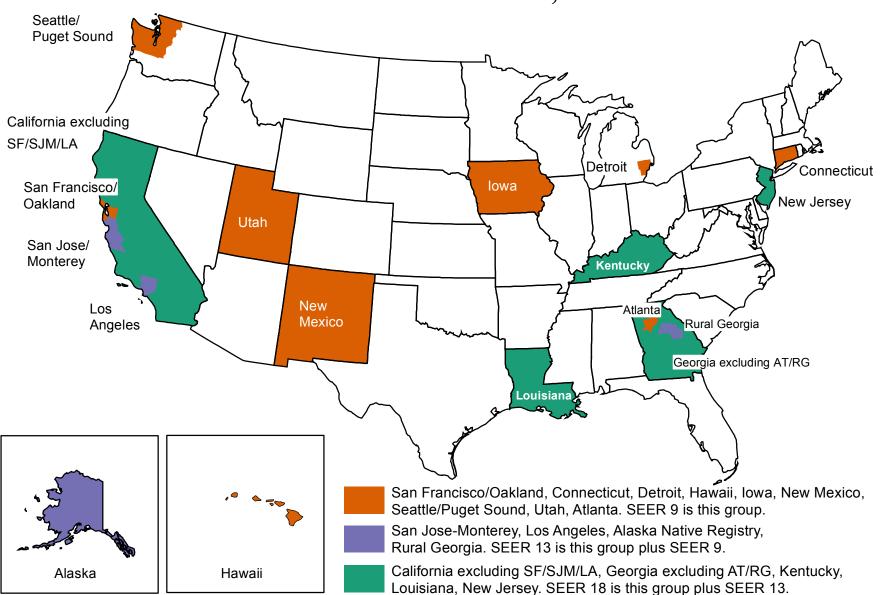
Age-Adjusted U.S. Death Rates and Trends for the Top 15 Cancer Sites^a by Race/Ethnicity

Females

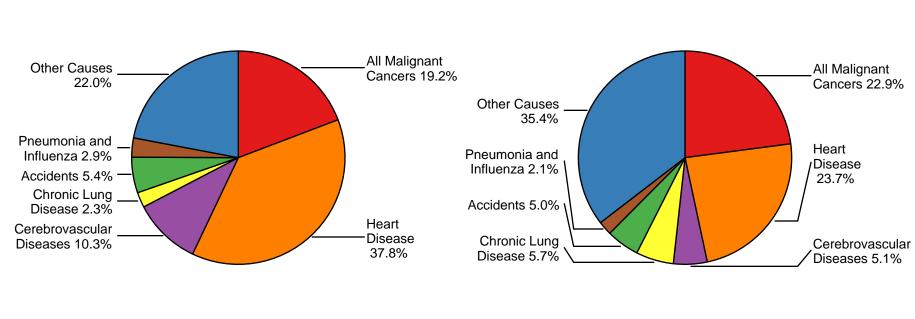
All Rac	ces		Whit	e		Blac	k	
	Rate ^b	APC ^c		Rate ^b	APC ^c		Rate ^b	APC ^c
	2007-2011	2002-2011		2007-2011	2002-2011		2007-2011	2002-2011
All Sites	147.4	-1.4*	All Sites	147.5	-1.4*	All Sites	169.0	-1.7*
Lung and Bronchus	38.5	-1.2*	Lung and Bronchus	39.8	-1.1*	Lung and Bronchus	36.5	-1.6*
Breast	22.2	-2.0*	Breast	21.7	-2.0*	Breast	30.6	-1.5*
Colon and Rectum	13.5	-2.9*	Colon and Rectum	13.0	-2.9*	Colon and Rectum	18.5	-3.2*
Pancreas	9.6	0.4*	Pancreas	9.4	0.5*	Pancreas	12.4	-0.2
Ovary	7.9	-2.0*	Ovary	8.2	-2.0*	Corpus and Uterus, NOS	7.5	1.0*
Leukemia	5.3	-1.0*	Leukemia	5.4	-0.9*	Ovary	6.6	-1.7*
Non-Hodgkin Lymphoma	5.0	-3.0*	Non-Hodgkin Lymphoma	5.2	-3.0*	Myeloma	5.3	-2.0*
Corpus and Uterus, NOS	4.3	0.8*	Corpus and Uterus, NOS	4.0	0.7*	Leukemia	4.8	-1.5*
Brain and ONS ^f	3.5	-0.5*	Brain and ONS^{f}	3.8	-0.4	Stomach	4.5	-3.5*
Liver & IBD ^f	3.4	1.8*	Liver & IBD ^f	3.2	1.9*	Liver & IBD ^f	4.2	1.7*
Myeloma	2.7	-1.9*	Kidney and Renal Pelvis	2.6	-1.2*	Cervix Uteri	4.1	-2.3*
Kidney and Renal Pelvis	2.6	-1.2*	Myeloma	2.5	-1.9*	Non-Hodgkin Lymphoma	3.5	-2.9*
Stomach	2.5	-2.7*	Urinary Bladder	2.2	-0.4	Kidney and Renal Pelvis	2.6	-0.7
Cervix Uteri	2.3	-1.1*	Stomach	2.1	-2.7*	Urinary Bladder	2.6	-1.6*
Urinary Bladder	2.2	-0.6*	Cervix Uteri	2.1	-0.8*	Brain and ONS^{f}	2.1	-0.2
Asian/Pacific	Islander		American Indian/Alaska Native ^d			Hispanic ^e		
	Rate ^b	APC ^c		Rate ^b	APC ^c		Rate ^b	APC ^c
	2007-2011	2002-2011		2007-2011	2002-2011		2007-2011	2002-2011
All Sites	91.5	-0.8*	All Sites	135.2	-1.6*	All Sites	99.9	-1.2*
Lung and Bronchus	18.4	-0.1	Lung and Bronchus	32.4	-1.5*	Breast	14.5	-1.5*
Breast	11.3	-1.6*	Colon and Rectum	15.6	0.0	Lung and Bronchus	14.0	-1.4*
Colon and Rectum	9.5	-1.3*	Breast	15.2	-2.8*	Colon and Rectum	9.9	-2.1*
Pancreas	7.2	0.5	Pancreas	8.0	0.8	Pancreas	7.7	0.1
Liver & IBD ^f	6.0	-1.4	Ovary	6.9	-0.3	Ovary	5.6	-1.6*
Stomach	4.8	-4.0*	Liver & IBD ^f	6.0	-2.4	Liver & IBD ^f	5.5	0.9*
Ovary	4.7	-1.0	Kidney and Renal Pelvis		-0.9	Non-Hodgkin Lymphoma	4.4	-1.8*
Non-Hodgkin Lymphoma	3.4	-2.0*	Non-Hodgkin Lymphoma	3.9	-3.2	Stomach	4.2	-2.8*
Leukemia	3.2	1.2	Stomach	3.8	-6.3*	Leukemia	3.9	-0.5
Corpus and Uterus, NOS	2.7	1.9*	Leukemia	3.5	-3.7	Corpus and Uterus, NOS	3.4	1.5
Cervix Uteri	1.8	-3.0*	Cervix Uteri	3.4	-2.2	Cervix Uteri	2.8	-2.4*
Brain and ONS^{f}	1.5	0.9	Corpus and Uterus, NOS	3.4	-	Brain and ONS^{f}	2.4	-0.6
Myeloma	1.4	-1.4	Brain and ONS ^f	2.3	-	Kidney and Renal Pelvis	2.3	-0.7
wide and paral palate	1.3	1.5	Myeloma	2.2	-5.6	Myeloma	2.3	-2.0*
Kidney and Renal Pelvis	1.3	1.5	Myeroma	2.2	5.0	nyerolla	2.5	2.0

- ^a Top 15 cancer sites selected based on 2007-2011 age-adjusted rates for the race/ethnic group.
- Rates are per 100,000 and age-adjusted to the 2000 US Std Population (19 age groups Census P25-1130).
- ^c The APC is the Annual Percent Change over the time interval.
- Trends are based on rates age-adjusted to the 2000 US Std Population (19 age groups Census P25-1130).
- d Rates for American Indian/Alaska Native are based on the CHSDA(Contract Health Service Delivery Area) counties.
- Hispanic is not mutually exclusive from whites, blacks, Asian/Pacific Islanders, and American Indians/Alaska Natives.
 - IBD = Intrahepatic Bile Duct. ONS = Other Nervous System.
 - The APC is significantly different from zero (p<.05).
 - Statistic not shown. Rate based on less than 16 cases for the time interval. Trend based on less than 10 cases for at least one year within the time interval.

Surveillance, Epidemiology, and End Results (SEER) Program: SEER 9, 13, & 18 Geographic Areas National Cancer Institute, USA



Leading Causes of Death in US, 1975 vs 2011 Percent of All Causes of Death



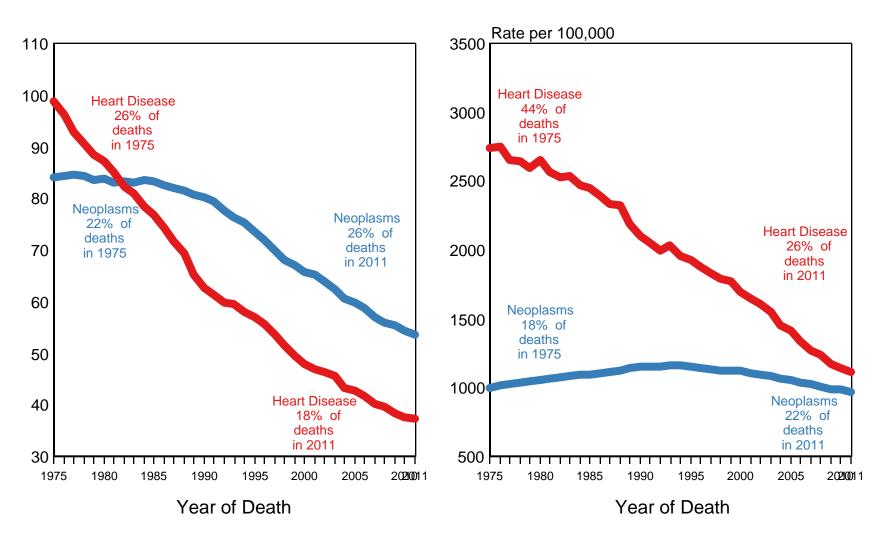
1975

2011

Us Death Rates, 1975-2011 Heart Disease compared to Neoplasms, by age at death

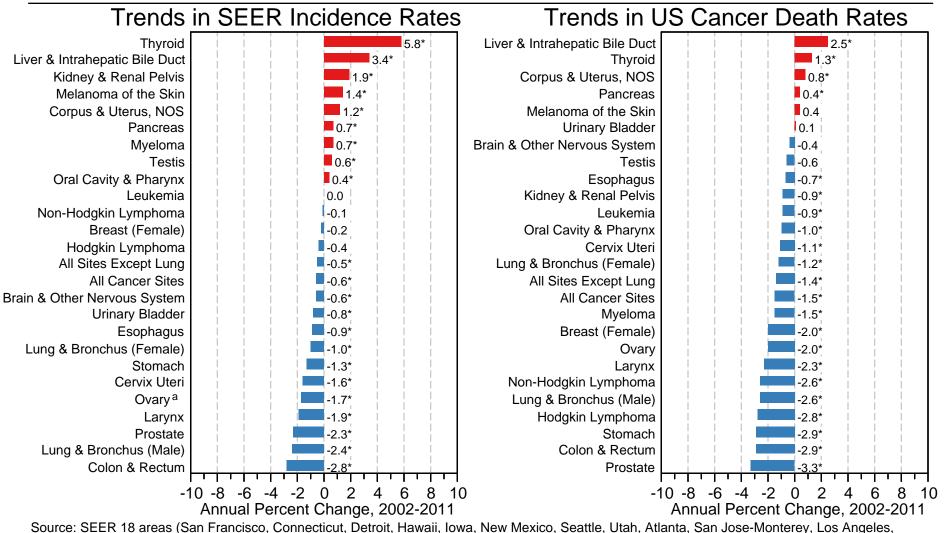
Ages Less Than 65

Ages 65 and Over



Source: US Mortality Files, National Center for Health Statistics, Centers for Disease Control and Prevention. Rates are per 100,000 and age-adjusted to the 2000 US Std Population (19 age groups - Census P25-1103).

Trends in SEER Incidence and US Death Rates by Primary Cancer Site 2002-2011



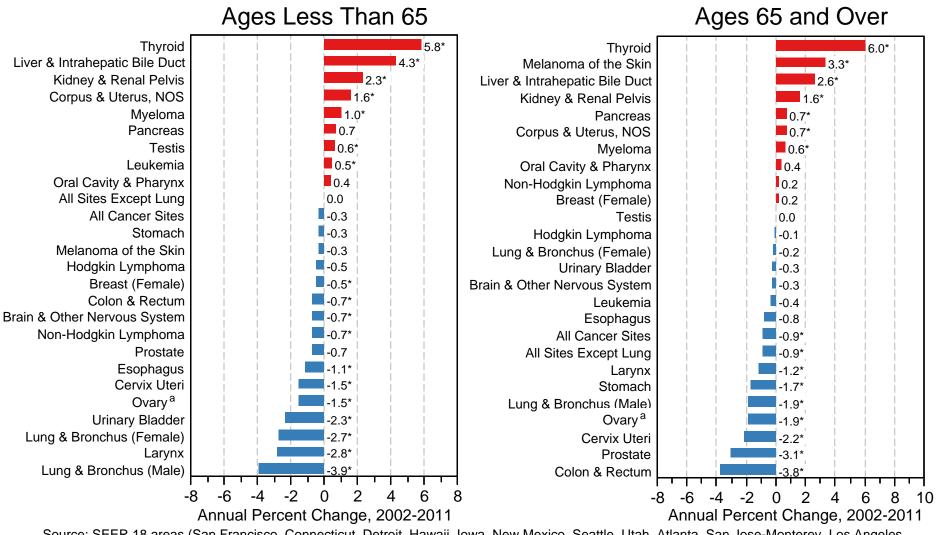
Alaska Native Registry, Rural Georgia, California excluding SF/SJM/LA, Kentucky, Louisiana, New Jersey and Georgia excluding ATL/RG) and US Mortality Files, National Center for Health Statistics, Centers for Disease Control and Prevention. Underlying rates are per 100,000 and age-adjusted to the 2000 US Std Population (19 age groups - Census P25-1103).

For sex-specific cancer sites, the population was limited to the population of the appropriate sex.

* The APC is significantly different from zero (p<.05).

^a Ovary excludes borderline cases or histologies 8442, 8451, 8462, 8472, and 8473.

Trends in SEER Incidence Rates by Age Group and Primary Cancer Site 2002-2011

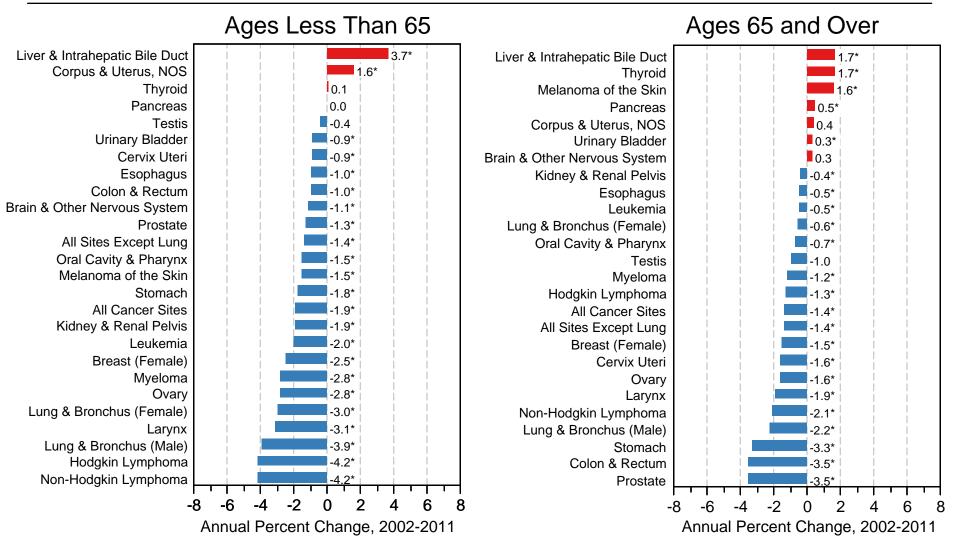


Source: SEER 18 areas (San Francisco, Connecticut, Detroit, Hawaii, Iowa, New Mexico, Seattle, Utah, Atlanta, San Jose-Monterey, Los Angeles, Alaska Native Registry, Rural Georgia, California excluding SF/SJM/LA, Kentucky, Louisiana, New Jersey and Georgia excluding ATL/RG). Underlying rates are per 100,000 and age-adjusted to the 2000 US Std Population (19 age groups - Census P25-1103). For sex-specific cancer sites, the population was limited to the population of the appropriate sex.

* The APC is significantly different from zero (p<.05).

^a Ovary excludes borderline cases or histologies 8442, 8451, 8462, 8472, and 8473.

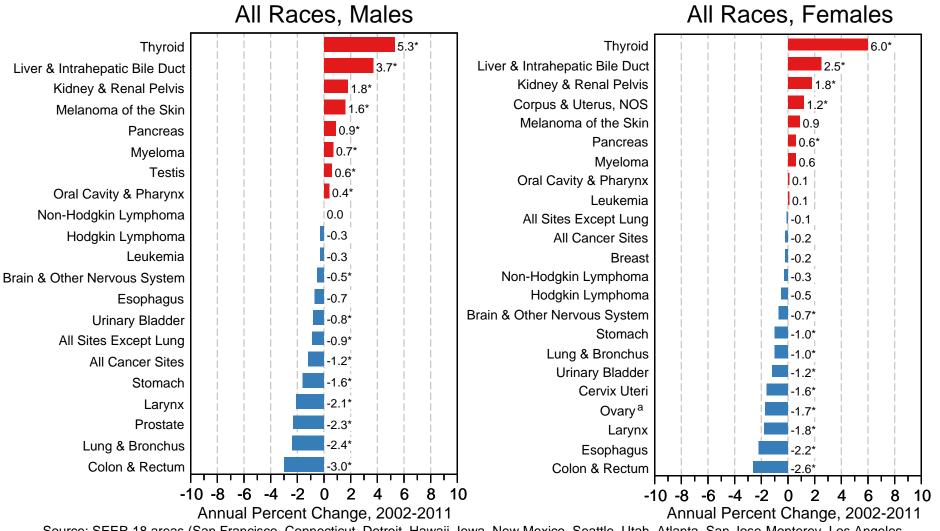
Trends in US Death Rates by Age Group and Primary Cancer Site 2002-2011



Source: US Mortality Files, National Center for Health Statistics, Centers for Disease Control and Prevention. Underlying rates are per 100,000 and age-adjusted to the 2000 US Std Population (19 age groups - Census P25-1103). For sex-specific cancer sites, the population was limited to the population of the appropriate sex.

* The APC is significantly different from zero (p<.05).

Trends in SEER Incidence Rates by Sex and Primary Cancer Site 2002-2011

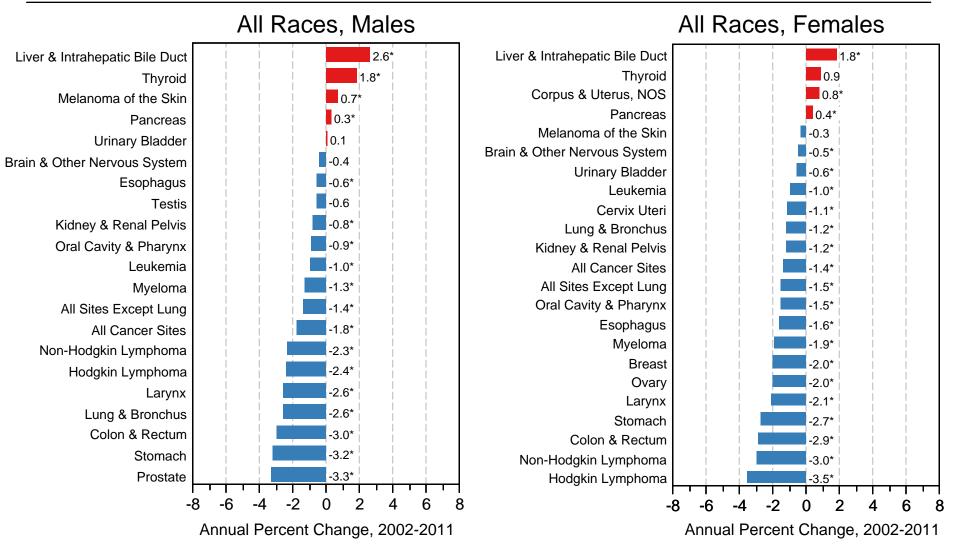


Source: SEER 18 areas (San Francisco, Connecticut, Detroit, Hawaii, Iowa, New Mexico, Seattle, Utah, Atlanta, San Jose-Monterey, Los Angeles, Alaska Native Registry, Rural Georgia, California excluding SF/SJM/LA, Kentucky, Louisiana, New Jersey and Georgia excluding ATL/RG). Underlying rates are per 100,000 and age-adjusted to the 2000 US Std Population (19 age groups - Census P25-1103). For sex-specific cancer sites, the population was limited to the population of the appropriate sex.

The APC is significantly different from zero (p<.05).

^a Ovary excludes borderline cases or histologies 8442, 8451, 8462, 8472, and 8473.

Trends in US Death Rates by Sex and Primary Cancer Site 2002-2011

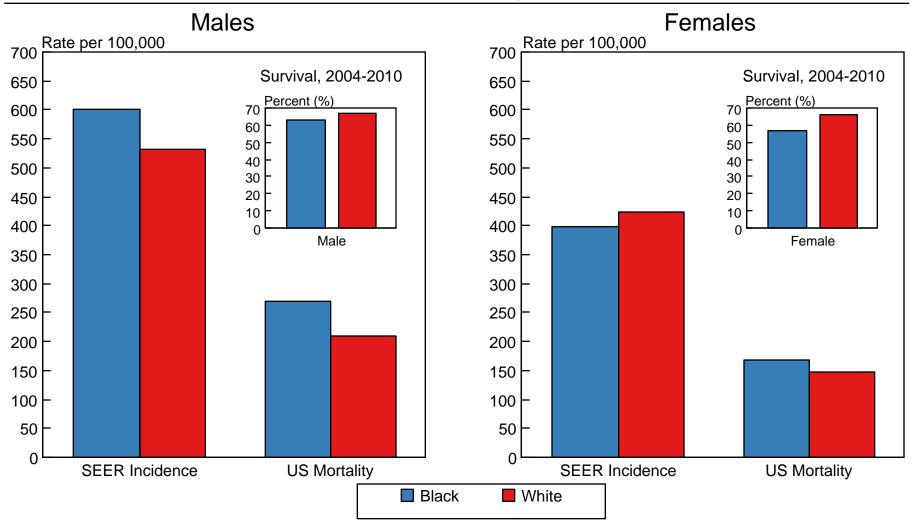


Source: US Mortality Files, National Center for Health Statistics, Centers for Disease Control and Prevention. Underlying rates are per 100.000 and age-adjusted to the 2000 US Std Population (19 age groups - Census P25-1103). For sex-specific cancer sites, the population was limited to the population of the appropriate sex.

* The APC is significantly different from zero (p<.05).

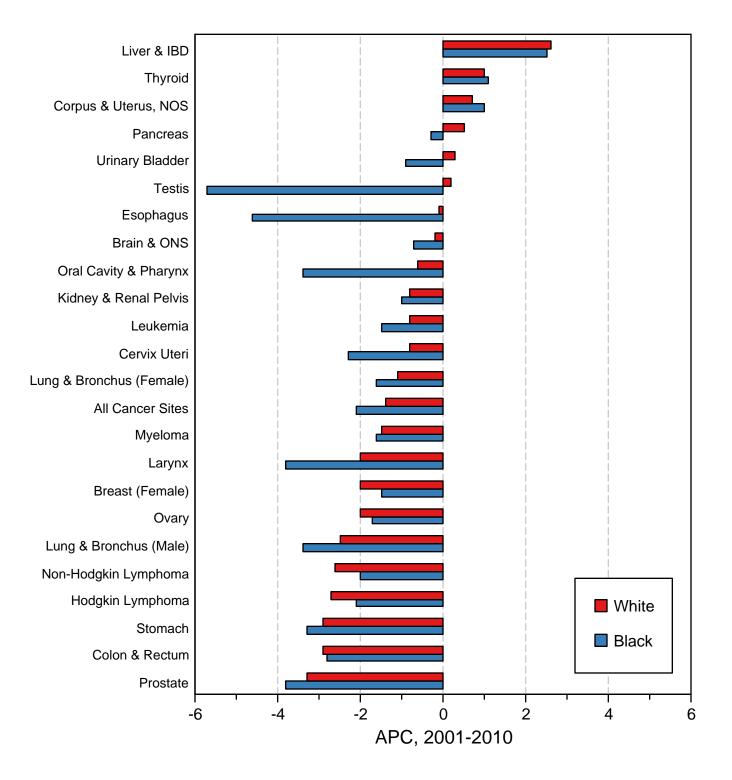
National Cancer Institute

SEER Incidence^a and US Death Rates^b, 2007-2011 5-Year Relative Survival^c, 2004-2010 All Cancer Combined, by Race and Sex



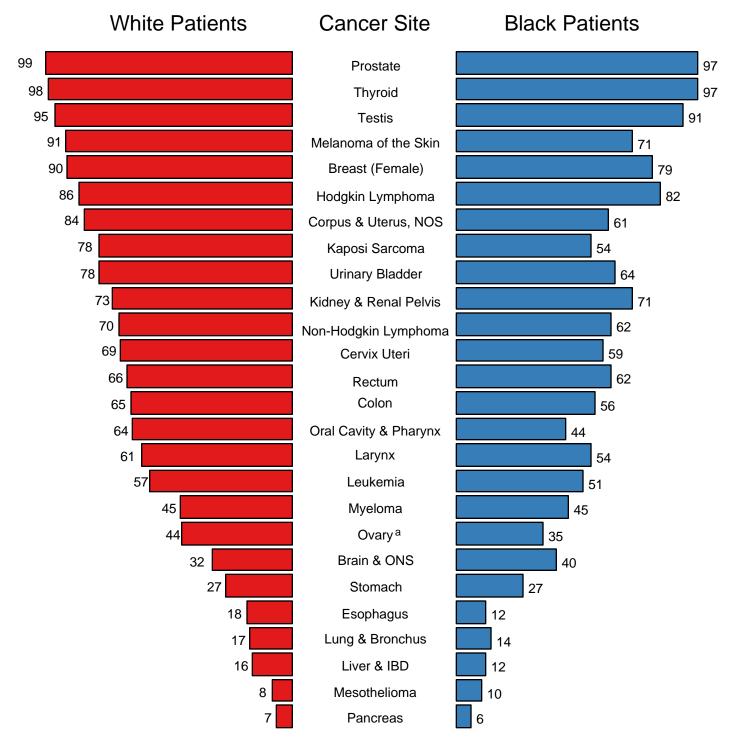
- a Incidence rates are from the SEER 18 areas (San Francisco, Connecticut, Detroit, Hawaii, Iowa, New Mexico, Seattle, Utah, Atlanta, San Jose-Monterey, Los Angeles, Alaska Native Registry, Rural Georgia, California excluding SF/SJM/LA, Kentucky, Louisiana, New Jersey and Georgia excluding ATL/RG) and are age-adjusted to the 2000 US Std Population (19 age groups - Census P25-1103).
- b Death rates are from the US Mortality Files, National Center for Health Statistics, Centers for Disease Control and Prevention and are age-adjusted to the 2000 US Std Population (19 age groups - Census P25-1103).
- ^c Survival rates are from the SEER 18 areas (San Francisco, Connecticut, Detroit, Hawaii, Iowa, New Mexico, Seattle, Utah, Atlanta, San Jose-Monterey, Los Angeles, Alaska Native Registry, Rural Georgia, California excluding SF/SJM/LA, Kentucky, Louisiana, New Jersey and Georgia excluding ATL/RG).

Trends in US Death Rates, 2002-2011 All Ages, by Race and Primary Cancer Site



Source: US Mortality Files, National Center for Health Statistics, Centers for Disease Control and Prevention. The APC is the Annual Percent Change over the time interval. Trends are based on rates age-adjusted to the 2000 US Std Population (19 age groups - Census P25-1130).

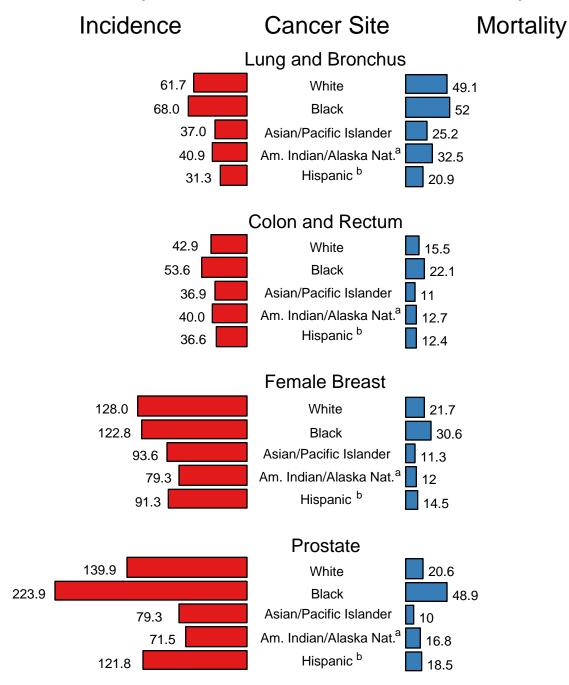
5-Year Relative Survival (%) SEER Program, 2004-2010 Both Sexes, by Race and Cancer Site



Source: SEER 18 areas (San Francisco, Connecticut, Detroit, Hawaii, Iowa, New Mexico, Seattle, Utah, Atlanta, San Jose-Monterey, Los Angeles, Alaska Native Registry, Rural Georgia, California excluding SF/SJM/LA, Kentucky, Louisiana, New Jersey and Georgia excluding ATL/RG).

^a Ovary excludes borderline cases or histologies 8442, 8451, 8462, 8472, and 8473.

SEER Cancer Incidence and US Death Rates, 2007-2011 By Cancer Site and Race/Ethnicity



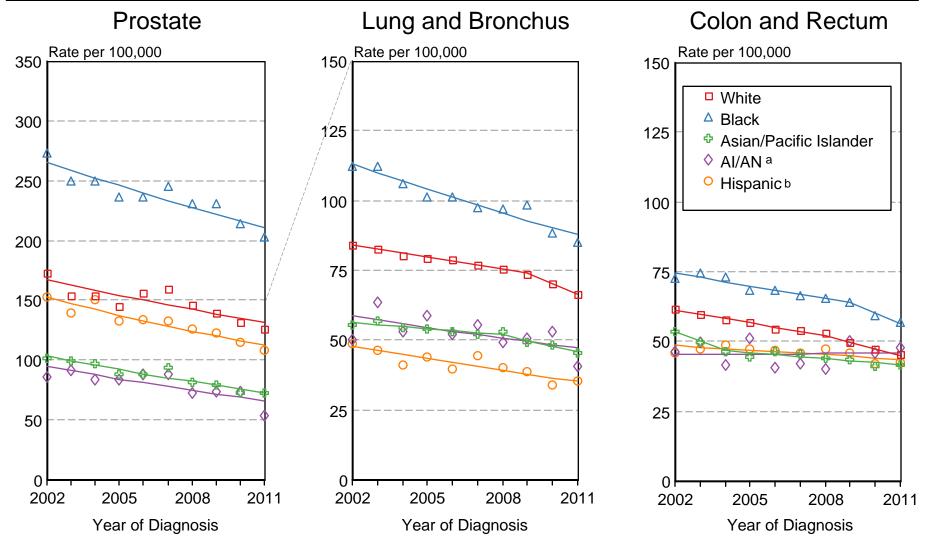
Source: SEER 18 areas (San Francisco, Connecticut, Detroit, Hawaii, Iowa, New Mexico, Seattle, Utah, Atlanta, San Jose-Monterey, Los Angeles, Alaska Native Registry, Rural Georgia, California excluding SF/SJM/LA, Kentucky, Louisiana, New Jersey and Georgia excluding ATL/RG) and US Mortality Files,

- ^a National Center for Health Statistics, Centers for Disease Control and Prevention.
- Rates for American Indian/Alaska Native are based on the CHSDA (Contract Health Service Delivery Area) ^b counties.

Hispanic is not mutually exclusive from whites, blacks, Asian/Pacific Islanders, and American Indians/Alaska Natives.

Incidence data for Hispanics are based on NHIA and exclude cases from the Alaska Native Registry.

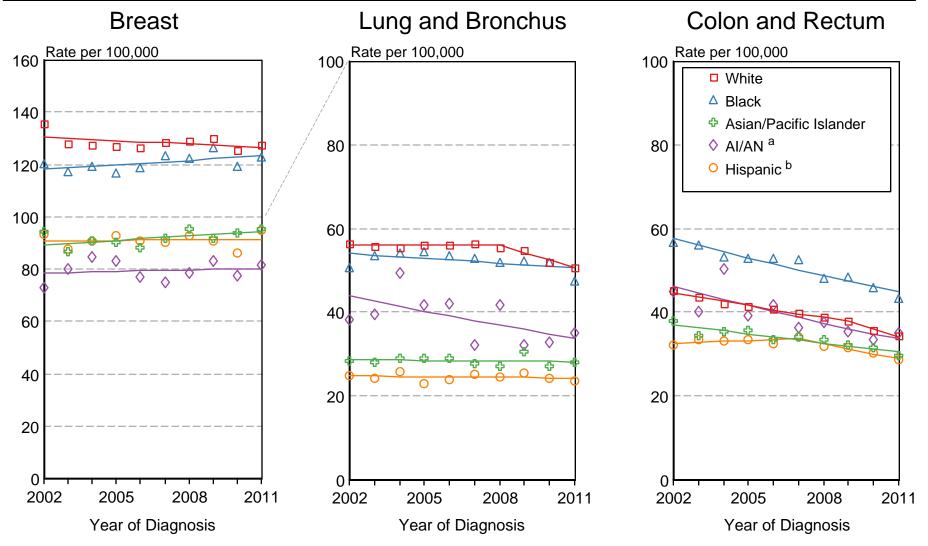
SEER Incidence 2002-2011 Males by Race/Ethnicity



Source: SEER 18 areas (San Francisco, Connecticut, Detroit, Hawaii, Iowa, New Mexico, Seattle, Utah, Atlanta, San Jose-Monterey, Los Angeles, Alaska Native Registry, Rural Georgia, California excluding SF/SJM/LA, Kentucky, Louisiana, New Jersey and Georgia excluding ATL/RG). Rates are age-adjusted to the 2000 US Std Population (19 age groups - Census P25-1103). Regression lines are calculated using the Joinpoint Regression Program Version 4.1.0, April 2014, National Cancer Institute.

- a Incidence rates for American Indian/Alaska Native (AI/AN) are based on the CHSDA(Contract Health Service Delivery Area) counties.
- ^b Hispanic is not mutually exclusive from whites, blacks, Asian/Pacific Islanders, and American Indians/Alaska Natives. Incidence data for Hispanics are based on NHIA and exclude cases from the Alaska Native Registry.

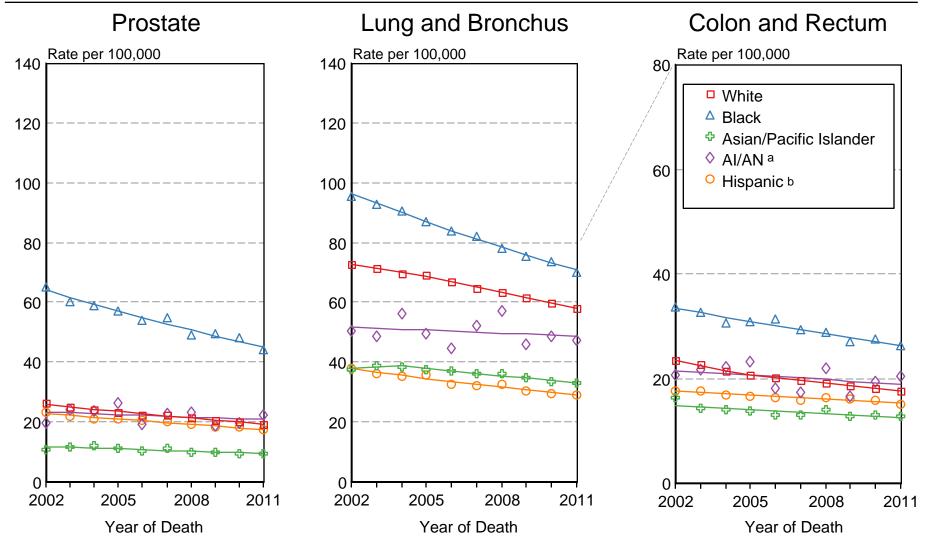
SEER Incidence 2002-2011 Females by Race/Ethnicity



Source: SEER 18 areas (San Francisco, Connecticut, Detroit, Hawaii, Iowa, New Mexico, Seattle, Utah, Atlanta, San Jose-Monterey, Los Angeles, Alaska Native Registry, Rural Georgia, California excluding SF/SJM/LA, Kentucky, Louisiana, New Jersey and Georgia excluding ATL/RG). Rates are age-adjusted to the 2000 US Std Population (19 age groups - Census P25-1103).

- Regression lines are calculated using the Joinpoint Regression Program Version 4.1.0, April 2014, National Cancer Institute. ^a Incidence rates for American Indian/Alaska Native (AI/AN) are based on the CHSDA(Contract Health Service Delivery Area) counties.
- ^b Hispanic is not mutually exclusive from whites, blacks, Asian/Pacific Islanders, and American Indians/Alaska Natives.
 Incidence data for Hispanics are based on NHIA and exclude cases from the Alaska Native Registry.

US Mortality 2001-2010 Males by Race/Ethnicity



Source: US Mortality Files, National Center for Health Statistics, Centers for Disease Control and Prevention.

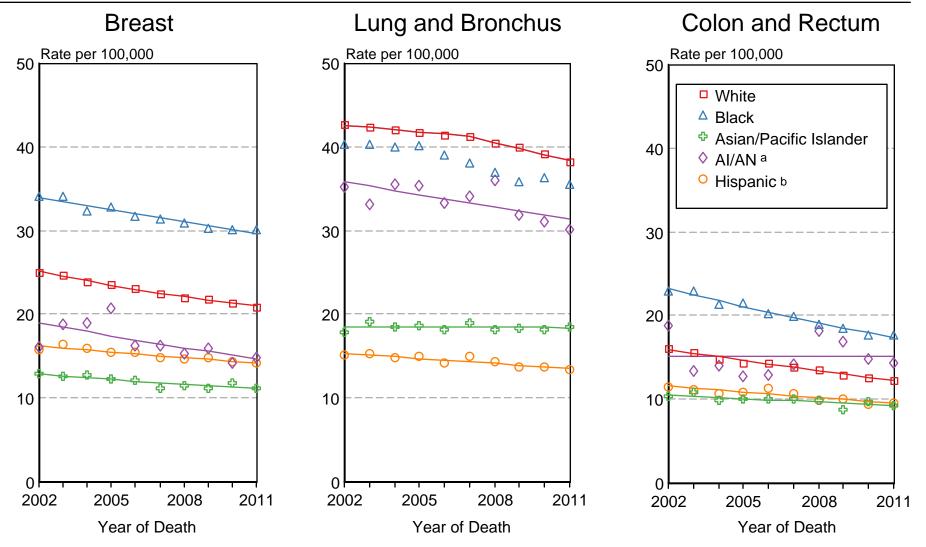
Rates are age-adjusted to the 2000 US Std Population (19 age groups - Census P25-1103).

^a Regression lines are calculated using the Joinpoint Regression Program Version 4.1.0, April 2014, National Cancer Institute.

^b Mortality rates for American Indian/Alaska Native (AI/AN) are based on the CHSDA(Contract Health Service Delivery Area) counties.

Hispanic is not mutually exclusive from whites, blacks, Asian/Pacific Islanders, and American Indians/Alaska Natives.

US Mortality 2001-2010 Females by Race/Ethnicity



Source: US Mortality Files, National Center for Health Statistics, Centers for Disease Control and Prevention.

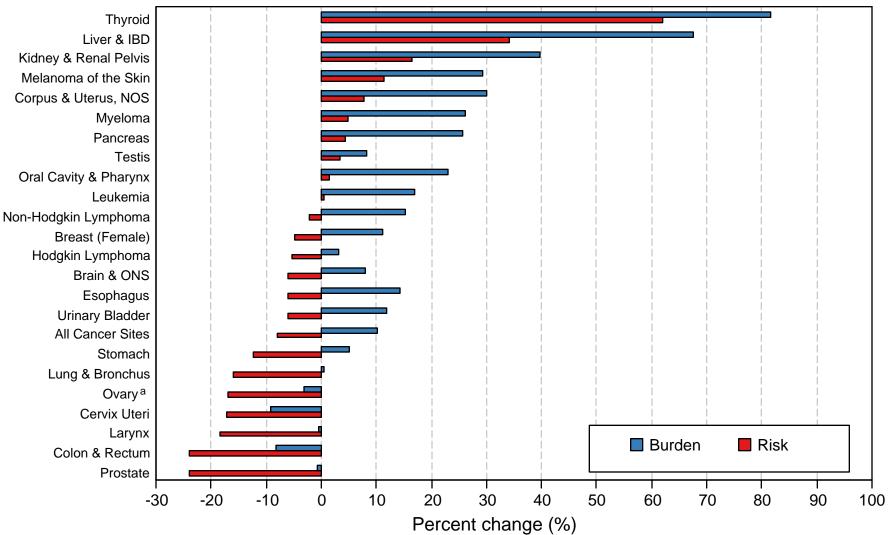
Rates are age-adjusted to the 2000 US Std Population (19 age groups - Census P25-1103).

^a Regression lines are calculated using the Joinpoint Regression Program Version 4.1.0, April 2014, National Cancer Institute.

^b Mortality rates for American Indian/Alaska Native (AI/AN) are based on the CHSDA(Contract Health Service Delivery Area) counties.

Hispanic is not mutually exclusive from whites, blacks, Asian/Pacific Islanders, and American Indians/Alaska Natives.

Incidence Percent Change between 2002 and 2011 Numbers (burden) vs Rates (risk) All Races, All Ages, Both Sexes

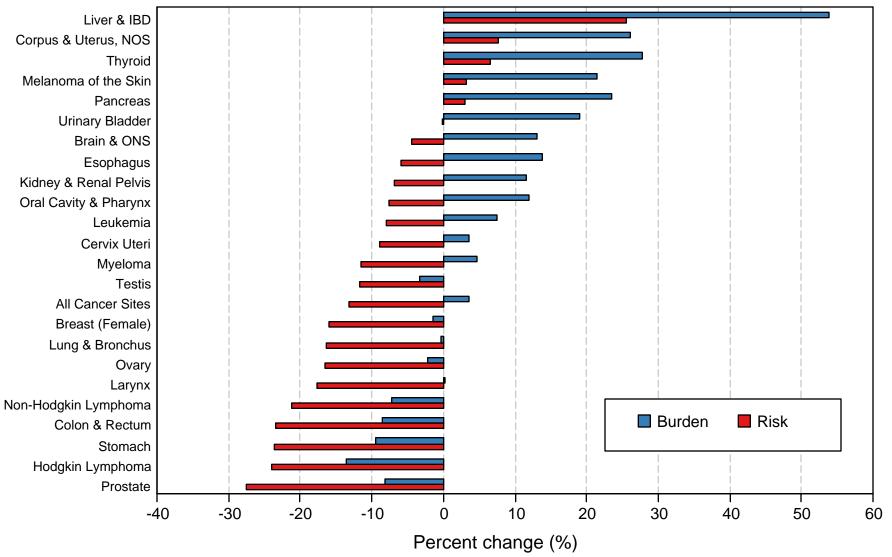


Source: SEER 18 areas (San Francisco, Connecticut, Detroit, Hawaii, Iowa, New Mexico, Seattle, Utah, Atlanta, San Jose-Monterey, Los Angeles, Alaska Native Registry, Rural Georgia, California excluding SF/SJM/LA, Kentucky, Louisiana, New Jersey and Georgia excluding ATL/RG). Burden is the change in the number of incidence cases between 2002 and 2011.

Risk is the change in the cancer incidence rates between 2002 and 2011.

^a Ovary excludes borderline cases or histologies 8442, 8451, 8462, 8472, and 8473.

Mortality Percent Change between 2002 and 2011 Numbers (burden) vs Rates (risk) All Races, All Ages, Both Sexes

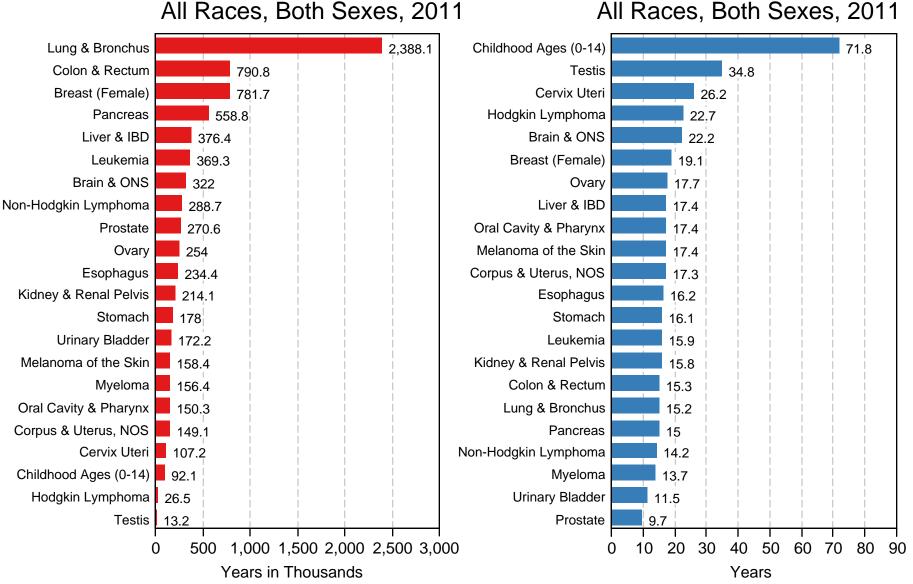


US Mortality estimates based on US age-specific rates applied to US population. Burden is the change in the number of deaths between 2002 and 2011. Risk is the change in the cancer death rates between 2002 and 2011.

Person-Years of Life Lost

Due to Cancer

Average Years of Life Lost Per Person Dying of Cancer All Races, Both Sexes, 2011



Person-Years of Life Lost Due to Major Causes of Death in US All Races, Both Sexes, 2011

Average Years of Life Lost Per Person Due to Major Causes of Death in US All Races, Both Sexes, 2011

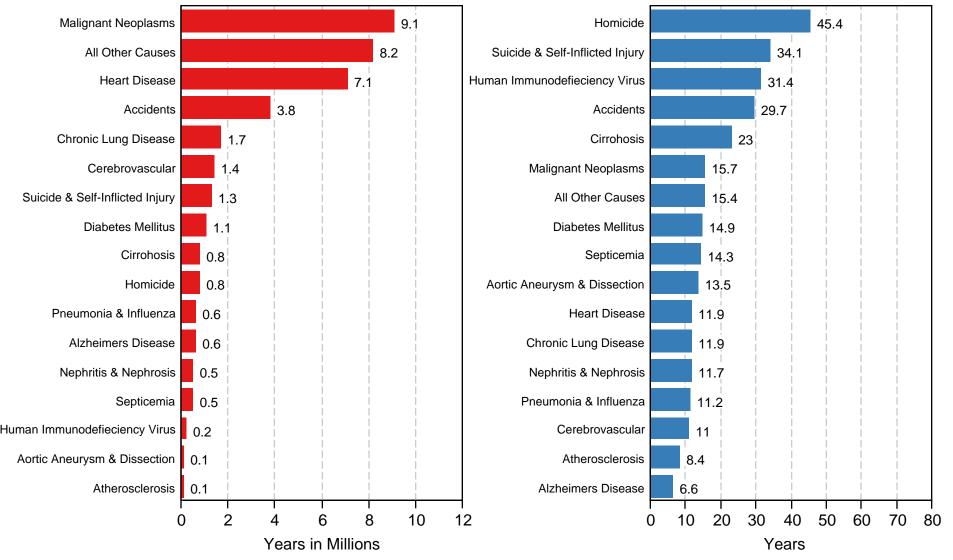
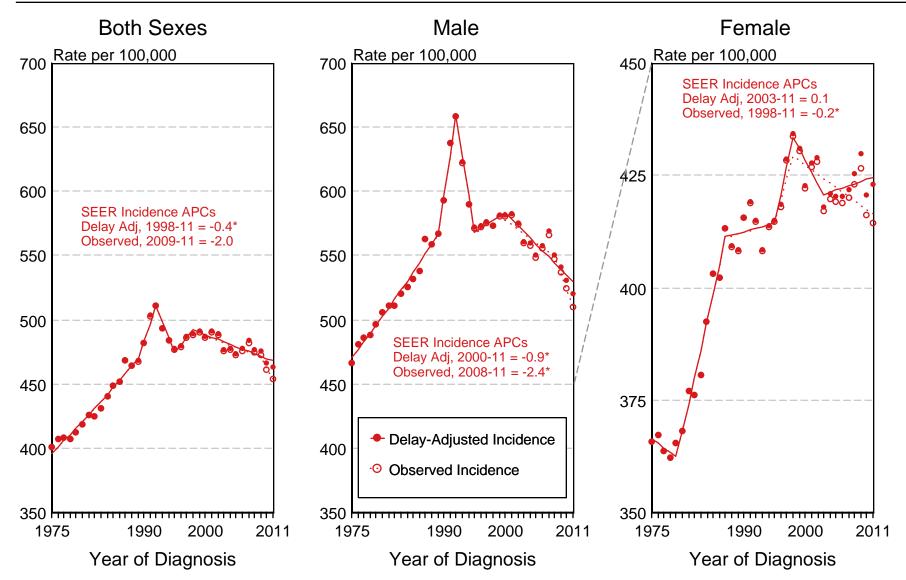
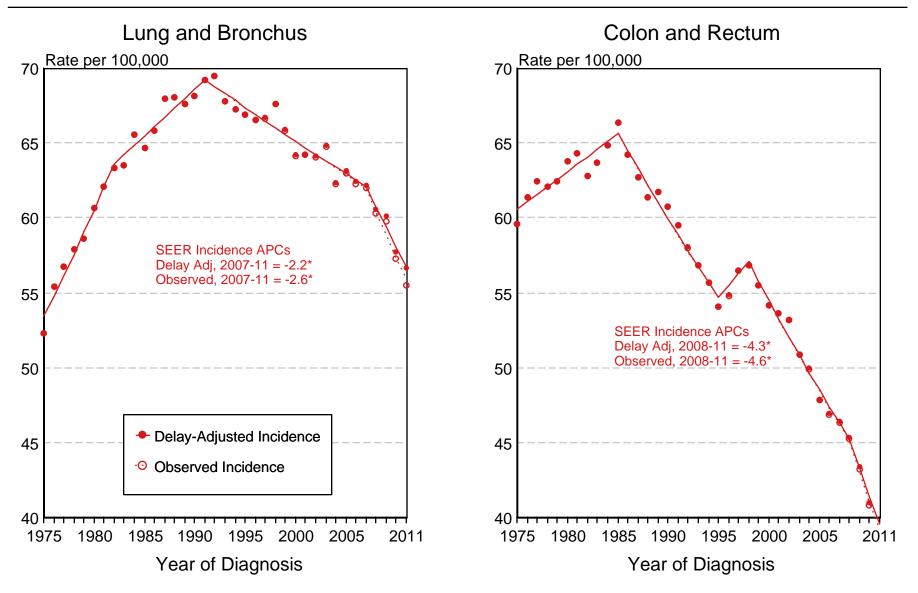


Figure 1.21 SEER Observed Incidence and Delay Adjusted Incidence Rates^a All Cancer Sites, By Sex



^a Source: SEER 9 areas. Rates are age-adjusted to the 2000 US Std Population (19 age groups - Census P25-1103). Regression lines and APCs are calculated using the Joinpoint Regression Program Version 4.1.0, April 2014, National Cancer Institute. The APC is the Annual Percent Change for the regression line segments. The APC shown on the graph is for the most recent trend. * The APC is significantly different from zero (p < 0.05).

Figure 1.22 SEER Observed Incidence and Delay Adjusted Incidence Rates^a **Both Sexes**

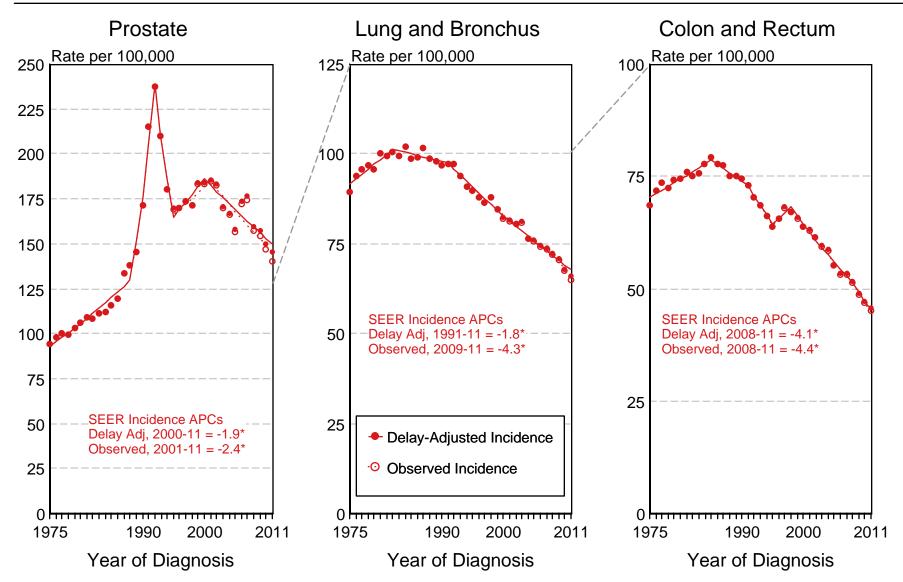


^a Source: SEER 9 areas. Rates are age-adjusted to the 2000 US Std Population (19 age groups - Census P25-1103). Regression lines and APCs are calculated using the Joinpoint Regression Program Version 4.1.0, April 2014, National Cancer Institute. The APC is the Annual Percent Change for the regression line segments. The APC shown on the graph is for the most recent trend.

* The APC is significantly different from zero (p < 0.05).

SEER Observed Incidence and Delay Adjusted Incidence Rates^a Males

Figure 1.23

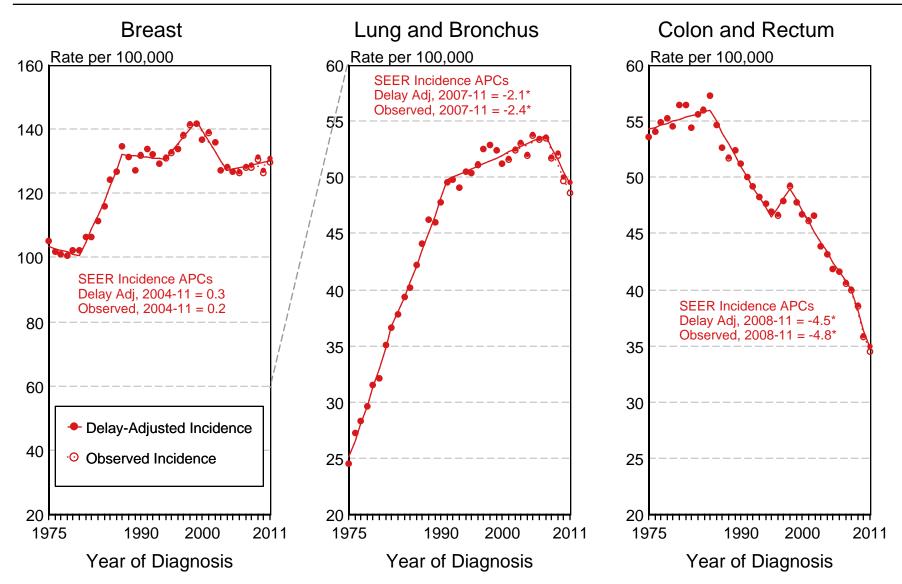


^a Source: SEER 9 areas. Rates are age-adjusted to the 2000 US Std Population (19 age groups - Census P25-1103). Regression lines and APCs are calculated using the Joinpoint Regression Program Version 4.1.0, April 2014, National Cancer Institute. The APC is the Annual Percent Change for the regression line segments. The APC shown on the graph is for the most recent trend.

* The APC is significantly different from zero (p < 0.05).

SEER Observed Incidence and Delay Adjusted Incidence Rates^a Females

Figure 1.24



^a Source: SEER 9 areas. Rates are age-adjusted to the 2000 US Std Population (19 age groups - Census P25-1103). Regression lines and APCs are calculated using the Joinpoint Regression Program Version 4.1.0, April 2014, National Cancer Institute. The APC is the Annual Percent Change for the regression line segments. The APC shown on the graph is for the most recent trend.

* The APC is significantly different from zero (p < 0.05).