



Editor's choice
Scan to access more
free content

ORIGINAL ARTICLE

Mortality and cancer incidence in a pooled cohort of US firefighters from San Francisco, Chicago and Philadelphia (1950–2009)

Robert D Daniels,¹ Travis L Kubale,¹ James H Yiin,¹ Matthew M Dahm,¹ Thomas R Hales,¹ Dalsu Baris,² Shelia H Zahm,² James J Beaumont,³ Kathleen M Waters,¹ Lynne E Pinkerton¹

► Additional material is published online only. To view please visit the journal online (<http://dx.doi.org/10.1136/oemed-2013-101662>).

¹Division of Surveillance, Hazard Evaluations, and Field Studies, National Institute for Occupational Safety and Health, Cincinnati, Ohio, USA
²Division of Cancer Epidemiology and Genetics, National Cancer Institute, Gaithersburg, Maryland, USA
³UC Davis Department of Public Health Sciences, Davis, Sacramento, California, USA

Correspondence to

Dr Robert D Daniels, Division of Surveillance, Hazard Evaluations, and Field Studies, National Institute for Occupational Safety and Health, 4676 Columbia Parkway, Mailstop R-13, Cincinnati, OH 45226, USA; rdtd2@cdc.gov

Received 12 June 2013
Revised 10 September 2013
Accepted 23 September 2013
Published Online First
14 October 2013



► <http://dx.doi.org/10.1136/oemed-2013-101662>

To cite: Daniels RD, Kubale TL, Yiin JH, et al. *Occup Environ Med* 2014;**71**:388–397.

ABSTRACT

Objectives To examine mortality patterns and cancer incidence in a pooled cohort of 29 993 US career firefighters employed since 1950 and followed through 2009.

Methods Mortality and cancer incidence were evaluated by life table methods with the US population referent. Standardised mortality (SMR) and incidence (SIR) ratios were determined for 92 causes of death and 41 cancer incidence groupings. Analyses focused on 15 outcomes of a priori interest. Sensitivity analyses were conducted to examine the potential for significant bias.

Results Person-years at risk totalled 858 938 and 403 152 for mortality and incidence analyses, respectively. All-cause mortality was at expectation (SMR=0.99, 95% CI 0.97 to 1.01, n=12 028). There was excess cancer mortality (SMR=1.14, 95% CI 1.10 to 1.18, n=3285) and incidence (SIR=1.09, 95% CI 1.06 to 1.12, n=4461) comprised mainly of digestive (SMR=1.26, 95% CI 1.18 to 1.34, n=928; SIR=1.17, 95% CI 1.10 to 1.25, n=930) and respiratory (SMR=1.10, 95% CI 1.04 to 1.17, n=1096; SIR=1.16, 95% CI 1.08 to 1.24, n=813) cancers. Consistent with previous reports, modest elevations were observed in several solid cancers; however, evidence of excess lymphatic or haematopoietic cancers was lacking. This study is the first to report excess malignant mesothelioma (SMR=2.00, 95% CI 1.03 to 3.49, n=12; SIR=2.29, 95% CI 1.60 to 3.19, n=35) among US firefighters. Results appeared robust under differing assumptions and analytic techniques.

Conclusions Our results provide evidence of a relation between firefighting and cancer. The new finding of excess malignant mesothelioma is noteworthy, given that asbestos exposure is a known hazard of firefighting.

INTRODUCTION

There are approximately 1.1 million volunteer and career firefighters in the US.¹ During firefighting activities, these workers may be exposed to many known carcinogens (eg, polycyclic aromatic hydrocarbons (PAHs), formaldehyde, benzene, 1,3-butadiene, asbestos and arsenic) in volatilised combustion and pyrolysis products or debris.² These exposures have raised concerns of increased cancer among firefighters and have prompted a number of exposure assessment and epidemiologic investigations. Some studies have found excess

What this paper adds

- From previous studies, there is limited epidemiological evidence of increased risk of cancer from firefighting.
- We examined cancer in 30 000 career firefighters by pooling information from urban fire departments in three large US cities. The large sample size and long follow-up period improved risk estimates compared with previous studies.
- We report that firefighting may be associated with increased risk of solid cancers. Furthermore, we report a new finding of excess malignant mesothelioma among firefighters, suggesting the presence of an occupational disease from asbestos hazards in the workplace.

cancers of the brain,^{3–8} digestive tract,^{4 5 7–10} genitourinary tract^{5 7 11 12} and lymphohematopoietic organs.^{6 8 13} In a recent meta-analysis of 32 studies, significant excess risk was reported for brain, stomach, colon, rectum, prostate, testes, multiple myeloma and non-Hodgkin lymphoma (NHL).¹⁴ Similarly, the International Agency for Research on Cancer (IARC) reviewed 42 studies and reported significant summary risks for prostatic and testicular cancers and NHL.² Given limited evidence, however, IARC concluded that firefighter exposures were only possibly carcinogenic to humans (Group 2B).

Most studies have examined mortality, but not cancer incidence, among relatively few firefighters recruited from one fire department. The current study examines mortality and cancer incidence in a pooled cohort of firefighters employed in three major US cities. Malignancies of the brain, stomach, oesophagus, intestines, rectum, kidney, bladder, prostate, testes, leukaemia, multiple myeloma and NHL were of a priori interest in the current study, based on possible sites identified in previous reviews.^{2 14} Lung cancer and chronic obstructive pulmonary disease (COPD) were also of interest because inhalation is a major pathway for firefighter exposures, and there is evidence of

chronic and acute inflammatory respiratory effects in firefighters, which may be linked to cancer.² Breast cancer was included as a result of interests shared in researcher discussions with firefighters.

METHODS

Data collection methods

This research was approved by the Institutional Review Boards of the National Institute for Occupational Safety and Health (NIOSH) and the National Cancer Institute (NCI). Personnel records and previous study data were used to assemble the study roster, which comprised male and female career firefighters of all races employed for at least 1 day in fire departments serving San Francisco, Chicago, or Philadelphia, from 1 January 1950, through 31 December 2009. Fire departments were selected based on size, location, work experience, records availability and the willingness of labour and city management to participate. 'Career firefighter' status was determined from job titles categorised by researchers and vetted by each fire department. Selected job titles included general classifications of firefighters, firefighter paramedics, and fire department arson investigators. Persons of known race were mostly Caucasian (81%) and those missing race (2.5%) were hired in earlier periods of lower minority hiring (median year at hire=1955). Therefore, persons missing race were assumed Caucasian and retained in main analyses to maximise study size. Analyses were also conducted excluding persons of unknown race.

Vital status was ascertained from the National Death Index-Plus (NDI-Plus), the Social Security Administration Death Master File (SSA-DMF), personnel and pension board records, and records from the previous studies.^{9 10} Firefighters not found to be deceased were confirmed alive by matches to employment records, Internal Revenue Service (IRS) records, and data accessible through LexisNexis (a private vendor of residential information).

Causes of death were obtained from previous studies,^{9 10} NDI-Plus, and death certificates collected from state vital records and retirement boards. Deaths of Philadelphia firefighters through 1986 were previously determined by Baris *et al.*,⁹ who retrieved and coded death certificates to the ninth revision of the International Classification of Diseases (ICD-9). San Francisco firefighter deaths were determined through 1982 by Beaumont *et al.*¹⁰ In that and the current study, causes of death were coded to the ICD revision in effect at the time of death. The underlying cause of death determined by a trained nosologist was used for all mortality analyses.

Incident cases were defined as all primary invasive cancers, and in situ bladder cancers among firefighters matched to state cancer registries on name, gender, race, date of birth and Social Security number. The last known residence and the state of death were used to narrow inclusion of registries for case ascertainment to 11 states (ie, Arizona, California, Florida, Illinois, Indiana, Michigan, Nevada, New Jersey, Oregon, Pennsylvania and Washington) where nearly 95% of all deaths in known states occurred (see online supplementary table S1). The site and histology of each tumour were used to classify cancers in one of 41 diagnostic groups using the International Classification of Diseases for Oncology, 3rd Edition (ICD-O-3).¹⁵ The conversion from ICD-O-3 to ICD-10 used the Surveillance, Epidemiology and End Results Program (SEER) recodes (dated 27 January 2003) following slight modification to align with mortality groupings and to account for recent classification changes. Diagnosis dates were assigned as of 1 July of the year of diagnosis if only the diagnosis year was

known, and on the 15th of the month of diagnosis if only the diagnosis month and year were known. The death date was used when death preceded the estimated date.

Statistical methods

The NIOSH Life Table Analysis System (LTAS.NET) was used to examine mortality and cancer incidence.¹⁶ Main analyses used the US population as referent. In all analyses, person-years at risk (PYAR) were stratified by gender, race (Caucasian, other races), age (age 15–85+ years in 5-year categories), and calendar year (in 5-year categories). Confidence limits for risk measures were estimated based on a Poisson distribution for the observed outcome, with exact limits for outcomes with 10 or fewer occurrences.

For mortality analyses, PYAR began on the latest of 1 January 1950 or the date of cohort inclusion, and ended the earliest of the date of death (DOD), the date last observed (DLO), or 31 December 2009. US mortality rates (1950–2009) were used to estimate the expected numbers of deaths for all causes, all cancers and 92 categories of underlying cause of death.¹⁷ Additional mortality rates were developed to separately report on cancers of the small intestine, large intestine and testes to coincide with incidence rates; however, these rates were limited to time periods after 1959. In both cases, the subsites of interest (ie, colon and testes) account for the largest proportion of the deaths in the respective aggregate site (ie, intestine or male genital organs excluding prostate); therefore, the aggregate site reasonably approximates the subsite. The standardised mortality ratio (SMR) was calculated as the ratio of the observed to the total number of expected deaths.

Two approaches were used to examine cancer incidence. The main analyses included first and later primary cancers (ie, multiple-cancer approach) occurring within the risk period. PYAR accrued from the date of statewide ascertainment by the respective fire department's state cancer registry (eg, 1 January 1988 for San Francisco firefighters (see online supplementary table S1)) or cohort inclusion, whichever was latest, and ended at the earliest of the DOD, DLO, or 31 December 2009. Secondary analyses were restricted to the first occurrence of invasive cancer (ie, first-cancer approach). In these analyses, PYAR for cases ended on the date of first diagnosis. In both approaches, the standardised incidence ratio (SIR) was calculated as the ratio of observed malignancies to the expected number of cases estimated using US incidence rates (1985–2009) calculated from SEER data.¹⁸ Additional steps required for first-cancer analyses were: selecting the most common cancer when diagnoses included multiple primary tumours on the same day (n=21), excluding firefighters known to have a cancer diagnosis prior to the start of the risk date (n=55), and adjusting US rates for cancer prevalence using methods described by Merrill *et al.*¹⁹

Heterogeneity in fire department-specific SMRs and SIRs was examined using Poisson regression modelling. To control for gender, age, calendar year and race, an offset term was set to the expected number of deaths or cases in each stratum of the classification table. To address differences between fire departments, a mixed model was used that specified a random intercept term. Thus, the model intercept is the log of the pooled SMR, adjusted for heterogeneity among the fire departments. The significance of heterogeneity was assessed by likelihood ratio test (significance level of 0.05).

Several sensitivity analyses were conducted. First, we examined the effects of including prevalent hires (workers employed before 1950) and short-term workers (those employed <1 year)

in mortality analyses. Prevalent hires must be employed long enough to be recruited into the study; thus, these workers may have a survival advantage compared with persons hired during the follow-up period (ie, incident hires).²⁰ Short-term workers include temporary hires and probationary firefighters whose health and lifestyle patterns may differ from those employed one or more years. Short-term workers may also have had substantial occupational histories other than as firefighters, possibly in jobs with hazardous exposures. Second, we examined age effects on risk estimates in two age-at-risk categories (17–64, 65+ years). Testing of an effect across all 5-year age groups was accomplished using mixed models adjusted for age-at-risk groups. Third, we conducted SMR analyses restricting observation to age 84 years or less. Including PYAR for ages 85+ years could bias results from: rates used in analyses that are open-ended, more uncertainty in underlying cause of death at later ages, and subjects who are incorrectly traced as alive having a disproportionate effect in the open-ended age group.²¹ Fourth, we calculated SMRs using California, Illinois and Pennsylvania State populations as referent for firefighters from San Francisco, Chicago and Philadelphia, respectively. Last, SMRs and standardised rate ratios (SRRs) were calculated for categories of employment duration (<10, 10–<20, 20–<30, 30+ years). Trend slopes with Wald-based two-sided p values (significance level of 0.05) were calculated for the change in SRRs with increasing duration.

RESULTS

There were 29 993 firefighters available for study, contributing 858 938 PYAR (table 1). The cohort was largely male (97%), with mean age at first employment and total years employed of 29 and 21 years, respectively. Fewer than 5% of firefighters

were short-term workers and approximately 30% were first employed prior to 1950. A higher percentage of women (9.4%) were short-term workers compared with men (4.3%) (see online supplementary table S2). Prevalent hires, on average, tended to be employed longer (+7.9 years, t test $p < 0.001$) and had a greater attained age (+17.0 years, t test $p < 0.001$) than incident hires. Persons eligible for incidence analyses using the multiple-cancer approach ($n = 24\,453$) contributed 403 152 PYAR. The first-cancer approach included 24 398 persons contributing 383 577 PYAR. There were 4461 malignant tumours distributed among 3903 firefighters with cancer. Among these, 488 reported cancers at multiple primary sites. Mortality and cancer incidence results are summarised in table 2 and in online supplementary tables S3–S5. To aid in comparisons with previous studies, table 2 also shows summary risk estimates (SREs) reported by LeMasters *et al*¹⁴, whose meta-analysis included studies published through 2003.

Mortality

With the US population referent, all-cause mortality was at expectation (SMR=0.99, 95% CI 0.97 to 1.01, $n = 12\,028$). Ischaemic heart disease was the leading cause of death (SMR=1.01, 95% CI 0.98 to 1.04, $n = 3619$). There was significantly decreased mortality in other outcomes that may be related to healthy worker selection and survivor effects (HWE), such as non-malignant respiratory diseases (SMR=0.80, 95% CI 0.74 to 0.86, $n = 796$), cerebrovascular disease (SMR=0.91, 95% CI 0.84 to 0.98, $n = 636$), diabetes mellitus (SMR=0.72, 95% CI 0.62 to 0.83, $n = 175$), nervous system disorders (SMR=0.80, 95% CI 0.69 to 0.93, $n = 187$), and alcoholism (SMR=0.61, 95% CI, 0.41 to 0.86, $n = 31$). In particular, there was a strong decrease in COPD mortality (SMR=0.72, 95% CI

Table 1 Demographic characteristics of the cohort by fire department and combined (1950–2009)

Description	All fire departments	San Francisco	Chicago	Philadelphia
Study cohort:				
Eligible for mortality analysis	29 993	5313	15 185	9495
PYAR	858 938	154 317	419 414	285 207
Years of follow-up; avg. (SD)	29 (16)	29 (16)	28 (16)	30 (16)
Race (%):				
White	24 244 (80.8)	4254 (80.1)	11 736 (77.3)	8254 (86.9)
Other	5008 (16.7)	986 (18.6)	2808 (18.5)	1214 (12.8)
Unknown	741 (2.5)	73 (1.4)	641 (4.2)	27 (<1.0)
Gender (%):				
Male	29 002 (96.7)	5009 (94.3)	14 694 (96.8)	9299 (97.9)
Female	991 (3.3)	304 (5.7)	491 (3.2)	196 (2.1)
Vital status:				
Alive (%)	17 965 (59.9)	3239 (61.0)	9241 (60.9)	5485 (57.8)
Deceased (%)	12 028 (40.1)	2074 (39.0)	5944 (39.1)	4010 (42.2)
Unknown cause of death	144	9	91	44
Attained age*; avg. (SD)	60 (16)	62 (16)	59 (16)	61 (16)
LTFU	175	1	32	142
PYAR potentially LTFU (%)	8809 (1.0)	59 (<1.0)	1483 (<1.0)	7267 (2.5)
Employment:				
Avg. hire year	1968	1967	1970	1965
Age at hire; avg. (SD)	29 (5)	29 (5)	29 (5)	27 (5)
Employment years; avg. (SD)	21 (11)	22 (11)	21 (11)	21 (11)
Hired before 1950 (%)	8085 (27)	1682 (32)	3294 (22)	3109 (33)
Employed <1 year (%)	1328 (4.4)	194 (3.7)	891 (5.9)	243 (2.6)

*Age attained at earliest of the date of death, date LTFU or 31 December 2009. Avg., average; LTFU, lost to follow-up; PYAR, person-years at risk.

Table 2 Standardised mortality and incidence ratios in firefighters for select outcomes compared to results from a recent meta-analysis

Underlying cause (ICD-10 codes)	Current study results (US population referent)					Meta-analysis of LeMasters <i>et al</i> ^{14*}		
	Mortality (1950–2009)†		Cancer incidence (1985–2009)			Studies	SRE (95% CI), Likelihood rating	
	Obs	SMR (95% CI)	All cancers	First cancer				
			Obs	SIR (95% CI)	Obs	SIR (95% CI)		
All cancers (C00–C97)	3285	1.14 (1.10 to 1.18)	4461	1.09 (1.06 to 1.12)	3890	1.09 (1.06 to 1.12)	25	1.05 (1.00 to 1.09), 3
MN oesophagus (C15)	113	1.39 (1.14 to 1.67)	90	1.62 (1.31 to 2.00)	80	1.71 (1.36 to 2.13)	8	1.16 (0.86 to 1.57), 3
MN stomach (C16)	110	1.10 (0.91 to 1.33)	93	1.15 (0.93 to 1.40)	72	1.02 (0.80 to 1.28)	13	1.22 (1.04 to 1.44), 2
MN intestine (C17–C18)	326	1.30 (1.16 to 1.44)	398	1.21 (1.09 to 1.33)	351	1.29 (1.16 to 1.43)	NA	NA
MN large intestine (C18)	264	1.31 (1.16 to 1.48)	381	1.21 (1.09 to 1.34)	335	1.28 (1.15 to 1.43)	25	1.21 (1.03 to 1.54), 2
MN small intestine (C17)	8	1.66 (0.72 to 3.27)	17	1.15 (0.67 to 1.85)	16	1.43 (0.82 to 2.33)	NA	NA
MN rectum (C19–C21)	89	1.45 (1.16 to 1.78)	166	1.11 (0.95 to 1.30)	140	1.09 (0.91 to 1.28)	13	1.29 (1.10 to 1.51), 2
MN lung (C33–C34)	1046	1.10 (1.04 to 1.17)	716	1.12 (1.04 to 1.21)	602	1.13 (1.04 to 1.22)	19	1.03 (0.97 to 1.08), 3
MN breast (C50)	8	1.39 (0.60 to 2.73)	26	1.26 (0.82 to 1.85)	24	1.32 (0.84 to 1.96)	NA	NA
MN prostate (C61)	282	1.09 (0.96 to 1.22)	1261	1.03 (0.98 to 1.09)	1176	1.03 (0.97 to 1.09)	13	1.28 (1.15 to 1.43), 1
MN other male genital (C60, C62–C63)	<5	0.47 (0.13 to 1.20)	17	0.62 (0.36 to 0.99)	17	0.67 (0.39 to 1.07)	NA	NA
MN testes (C62)	<5	0.73 (0.15 to 2.14)	15	0.75 (0.42 to 1.24)	15	0.79 (0.44 to 1.30)	4	2.02 (1.30 to 3.13), 2
MN kidney (C64–C66)	94	1.29 (1.05 to 1.58)	166	1.27 (1.09 to 1.48)	129	1.24 (1.04 to 1.48)	12	1.07 (0.78 to 1.46), 3
MN bladder (C67–C68)‡	84	0.99 (0.79 to 1.22)	316	1.12 (1.00 to 1.25)	272	1.18 (1.05 to 1.33)	11	1.20 (0.97 to 1.48), 3
MN brain (C47, C70–C72)	73	1.01 (0.79 to 1.27)	51	1.02 (0.76 to 1.34)	48	1.06 (0.78 to 1.41)	19	1.32 (1.12 to 1.54), 2
NHL (C46.3, C82–C85, C88.0, C88.3, C91.4, C96)§	123	1.17 (0.97 to 1.40)	170	0.99 (0.85 to 1.15)	145	0.99 (0.83 to 1.16)	8	1.51 (1.31 to 1.73), 1
Leukaemia (C91.0–C91.3, C91.5–C91.9, C92–C95)	122	1.10 (0.91 to 1.31)	100	0.94 (0.77 to 1.15)	85	0.93 (0.74 to 1.15)	8	1.14 (0.98 to 1.31), 2
Multiple myeloma (C88.7, C88.9, C90)	42	0.89 (0.64 to 1.20)	36	0.72 (0.50 to 0.99)	33	0.75 (0.52 to 1.06)	10	1.53 (1.21 to 1.94), 1
Other cancers:¶								
Mesothelioma (C45)	12	2.00 (1.03 to 3.49)	35	2.29 (1.60 to 3.19)	26	2.00 (1.31 to 2.93)	NA	NA
MN buccal and pharynx (C00–C14)	94	1.40 (1.13 to 1.72)	174	1.39 (1.19 to 1.62)	148	1.41 (1.20 to 1.66)	9	1.23 (0.96 to 1.55), 2

*Results from Table 5 of LeMasters *et al*¹⁴; likelihood of cancer risk by meta-analysis criteria: 1=probable, 2=possible, 3=unlikely.

†SMRs restricted to 1960–2009 for MN large intestine, MN small intestine, and MN testes and 2000–2009 for mesothelioma.

‡Urinary bladder incidence included in situ (D09.0) and invasive cases as per SEER protocol.

§NHL incidence data exclude Kaposi sarcoma (C46.3).

¶Sites not listed among cancers of a priori interest but reporting statistically significant excess mortality and cancer incidence.

ICD-10, International Classification of Diseases, 10th Revision; MN, malignancy; NA, not applicable; NHL, non-Hodgkin lymphoma; Obs, observed; SEER, Surveillance, Epidemiology and End Results; SIR, standardised incidence ratio; SMR, standardised mortality ratio; SRE, summary risk estimate.

0.65 to 0.80, $n=367$). Few non-malignant outcomes were elevated, although statistically significant excess mortality was observed for cirrhosis and other chronic liver disease (SMR=1.26, 95% CI 1.12 to 1.41, $n=299$) and acute glomerulonephritis with renal failure (SMR=1.56, 95% CI 1.07 to 2.20, $n=32$). Deaths from falls (SMR=1.31, 95% CI 1.08 to 1.58, $n=113$) and other accidents (SMR=1.17, 95% CI 1.01 to 1.34, $n=197$) were also elevated.

By contrast with non-malignant outcomes, we observed excess overall cancer mortality (SMR=1.14, 95% CI 1.10 to 1.18, $n=3285$) (table 2). The elevation was largely attributable to excess cancers of the lung (SMR=1.10, 95% CI 1.04 to 1.17, $n=1046$), oesophagus (SMR=1.39, 95% CI 1.14 to 1.67, $n=113$), intestine (SMR=1.30, 95% CI 1.16 to 1.44, $n=326$), rectum (SMR=1.45, 95% CI 1.16 to 1.78, $n=89$) and kidney (SMR=1.29, 95% CI 1.05 to 1.58, $n=94$). There was little evidence of excess mortality from the remaining cancers of a priori interest; however, statistically significant SMRs were apparent for buccal and pharynx cancers (SMR=1.40, 95% CI 1.13 to 1.72, $n=94$), malignancies of the liver, gall bladder and biliary tract (SMR=1.30, 95% CI 1.06 to 1.57, $n=107$), and malignant mesothelioma (SMR=2.00, 95% CI 1.03 to 3.49, $n=12$).

Women and non-Caucasians

All-cause mortality among women was near expectation (SMR=0.91, 95% CI 0.59 to 1.33, $n=26$). Accidental death was the leading cause (SMR=2.79, 95% CI 1.21 to 5.50, $n=8$) resulting in 31% of the total deaths among women. While there was little evidence of excess overall cancer mortality among women (SMR=0.74, 95% CI 0.27 to 1.61, $n=6$), most cancer deaths were from breast cancer (SMR=1.46, 95% CI 0.30 to 4.26, $n<5$). Bladder cancer mortality was statistically significant (SMR=33.51, 95% CI 4.06 to 121.05, $n<5$) based on few cases. Non-Caucasian males were characterised by decreased all-cause mortality (SMR=0.68, 95% CI 0.62 to 0.74, $n=453$) and all-cancers (SMR=0.80, 95% CI 0.65 to 0.97, $n=104$). They had few observed deaths in any a priori outcome, and lung cancer mortality was below expectation (SMR=0.67, 95% CI 0.44 to 0.97, $n=27$). Only prostate cancer mortality showed an excess approaching statistical significance (SMR=1.64, 95% CI 0.95 to 2.63, $n=17$) among non-Caucasian males (table 3).

Cancer incidence

There was little difference in SIRs when comparing analysis approaches; therefore, reporting focused on results from the multiple-cancer approach (table 2). All-cancer incidence was slightly above expectation (SIR=1.09, 95% CI 1.06 to 1.12, $n=4461$). Observed elevations in cancers of a priori interest were generally consistent with mortality data as evidenced by significant excess cancers of the oesophagus (SIR=1.62, 95% CI 1.31 to 2.00, $n=90$); large intestine (SIR=1.21, 95% CI 1.09 to 1.34, $n=381$); kidney (SIR=1.27, 95% CI 1.09 to 1.48, $n=166$) and lung (SIR=1.12, 95% CI 1.04 to 1.21, $n=716$). As in mortality analyses, there were excess buccal and pharynx cancers (SIR=1.39, 95% CI 1.19 to 1.62, $n=174$) and malignant mesothelioma (SIR=2.29, 95% CI 1.60 to 3.19, $n=35$). Of those diagnosed with mesothelioma, 31 (88.6%) were pleural. Excess laryngeal cancer incidence was also observed (SIR=1.50, 95% CI 1.19 to 1.85, $n=84$). The incidence of most remaining cancer sites was near expectation; however, multiple myeloma was significantly decreased (SIR=0.72, 95% CI 0.50 to 0.99, $n=36$).

Women and non-Caucasians

Overall cancer incidence among women was elevated, but not significantly (SIR=1.24, 95% CI 0.89 to 1.69, $n=40$). Consistent with mortality, female bladder cancer incidence was statistically significant but based on few cases (SIR=12.53, 95% CI 3.41 to 32.08, $n<5$). Nearly half of all cases were breast cancer (SIR=1.45, 95% CI 0.86 to 2.29, $n=18$). Nearly all breast cancers were diagnosed prior to the attained age of 55 years, with the highest SIR between the ages of 50 and 54 years (SIR=2.66, 95% CI 0.86 to 6.21, $n=5$). Left-sided disease appeared more frequent (61%, $n=11$). Overall cancer incidence among non-Caucasian male firefighters was near expectation (SIR=0.92, 95% CI 0.81 to 1.05, $n=240$). There was excess prostate cancer (SIR=1.26, 95% CI 1.02 to 1.54, $n=94$) but decreased lung cancer (SIR=0.67, 95% CI 0.43 to 1.00, $n=24$) (tables 3 and 4).

Sensitivity analyses

Except for COPD and cancers of the lung, prostate and brain, there was little evidence of heterogeneity in SMRs (see online supplementary table S6) or SIRs (see online supplementary table S7) across fire departments for outcomes of a priori interest. For mortality, the between-department variance was largely attributable to outlying decreased lung cancer (SMR=0.76, 95% CI 0.64 to 0.89, $n=142$) and COPD (SMR=0.53, 95% CI 0.40 to 0.69, $n=57$) in San Francisco firefighters, and excess cancers of the prostate (SMR=1.28, 95% CI 1.08 to 1.50, $n=152$) and lung (SMR=1.23, 95% CI 1.13 to 1.34, $n=566$) in Chicago firefighters. The between-department variance in mortality persisted when using state populations as referent (see online supplementary table S8). Similarly, heterogeneous lung cancer incidence stemmed from decreased cases among San Francisco firefighters (SIR=0.70, 95% CI 0.56 to 0.87, $n=81$); however, there was outlying excess prostate cancer incidence among San Francisco firefighters (SIR=1.22, 95% CI 1.08 to 1.37, $n=276$). Brain cancer SIRs varied widely across fire departments; excess cancer was observed in San Francisco firefighters (SIR=1.95, 95% CI 1.14 to 3.12, $n=17$), while decreased cancer was reported for Chicago (SIR=0.53, 95% CI 0.28 to 0.91, $n=13$).

Restricting analyses to firefighters with one or more years of employment had negligible effects (see online supplementary table S9). Slight increases in SMRs were observed for most a priori outcomes when restricting the cohort to incident hires, although these differences were not statistically significant. Age-at-risk differences in mortality also lacked statistical significance, but SMRs generally appeared greater at older ages. SMRs for cancers of the breast (SMR=1.42, 95% CI 0.46 to 3.32, $n=5$), oesophagus (SMR=1.41, 95% CI 1.05 to 1.86, $n=51$), and kidney (SMR=1.47, 95% CI 1.09 to 1.95, $n=48$) were highest among workers less than 65 years of age (see online supplementary table S10). Significant age-at-risk differences in SIRs were evident for prostate ($p<0.001$) and bladder ($p=0.002$) cancers (see online supplementary table S11). The heterogeneity was largely attributable to significant increases in prostate (SIR=1.21, 95% CI 1.10 to 1.33, $n=426$) and bladder (SIR=1.33, 95% CI 1.08 to 1.62, $n=97$) cancer risks among firefighter aged 64 years or less. Excess prostate cancer was limited to ages 45–59 years (SIR=1.45, 95% CI 1.28 to 1.64, $n=249$), while the age pattern of excess bladder cancer incidence was unclear. The effects of restricting PYAR to age-at-risk <85 were inconsequential (see online supplementary table S12). Excluding firefighters without race information also had little

Table 3 Standardised mortality and incidence ratios among men compared with the US population for causes of a priori interest

Underlying cause (ICD-10 codes)	Mortality (1950–2009)				Cancer incidence (1985–2009)*			
	Caucasian		Other		Caucasian		Other	
	Obs	SMR (95% CI)	Obs	SMR (95% CI)	Obs	SIR (95% CI)	Obs	SIR (95% CI)
All causes	11 549	1.01 (0.99 to 1.03)	453	0.68 (0.62 to 0.74)	NA	NA	NA	NA
All cancers (C00-C97)	3175	1.16 (1.12 to 1.20)	104	0.80 (0.65 to 0.97)	4181	1.10 (1.07 to 1.13)	240	0.92 (0.81 to 1.05)
MN oesophagus (C15)	110	1.46 (1.20 to 1.75)	<5	0.51 (0.11 to 1.49)	87	1.70 (1.36 to 2.09)	<5	0.73 (0.15 to 2.15)
MN stomach (C16)	105	1.12 (0.92 to 1.36)	5	0.81 (0.26 to 1.89)	87	1.19 (0.96 to 1.47)	6	0.76 (0.28 to 1.66)
MN intestine (C17-C18)	319	1.32 (1.18 to 1.48)	7	0.68 (0.27 to 1.40)	379	1.23 (1.11 to 1.36)	18	0.90 (0.53 to 1.42)
MN rectum (C19-C21)	86	1.46 (1.17 to 1.81)	<5	1.21 (0.25 to 3.53)	159	1.16 (0.99 to 1.36)	7	0.62 (0.25 to 1.28)
MN lung (C33-C34)	1019	1.12 (1.05 to 1.19)	27	0.67 (0.44 to 0.97)	689	1.15 (1.07 to 1.24)	24	0.67 (0.43 to 1.00)
MN breast (C50)	5	1.43 (0.46 to 3.34)	0	NC	6	0.79 (0.29 to 1.72)	<5	3.32 (0.40 to 12.00)
MN prostate (C61)	265	1.06 (0.94 to 1.20)	17	1.64 (0.95 to 2.63)	1167	1.02 (0.96 to 1.08)	94	1.26 (1.02 to 1.54)
MN other male genital (C60, C62-C63)	<5	0.49 (0.13 to 1.26)	0	NC	16	0.64 (0.37 to 1.04)	<5	0.38 (0.01 to 2.13)
MN kidney (C64-C66)	91	1.31 (1.05 to 1.60)	<5	1.05 (0.22 to 3.07)	151	1.26 (1.06 to 1.47)	14	1.46 (0.80 to 2.45)
MN bladder (C67-C68)†	80	0.96 (0.76 to 1.19)	<5	1.19 (0.14 to 4.30)	305	1.11 (0.99 to 1.24)	7	0.92 (0.37 to 1.91)
MN brain (C47, C70-C72)	72	1.03 (0.81 to 1.30)	<5	0.44 (0.01 to 2.47)	49	1.05 (0.78 to 1.39)	<5	0.67 (0.08 to 2.42)
NHL (C46.3, C82-C85, C88.0, C88.3, C91.4, C96)‡	119	1.18 (0.98 to 1.41)	<5	1.01 (0.28 to 2.60)	161	1.02 (0.87 to 1.19)	7	0.56 (0.23 to 1.16)
Leukaemia (C91.0-C91.3, C91.5-C91.9, C92-C95)	117	1.10 (0.91 to 1.32)	5	1.28 (0.41 to 2.98)	88	0.88 (0.71 to 1.09)	11	1.90 (0.95 to 3.40)
Multiple myeloma (C88.7, C88.9, C90)	41	0.92 (0.66 to 1.25)	<5	0.35 (0.01 to 1.97)	35	0.76 (0.53 to 1.06)	<5	0.24 (0.01 to 1.32)
COPD (J40-J44)	362	0.73 (0.65 to 0.81)	5	0.50 (0.16 to 1.16)	NA	NA	NA	NA

*Incidence results based on analysis of all invasive primary cancers (ie, multiple-cancer approach).

†Urinary bladder incidence included in situ (D09.0) and invasive cases as per SEER protocol.

‡NHL incidence data exclude Kaposi sarcoma (C46.3).

COPD, chronic obstructive pulmonary disease; ICD-10, International Classification of Diseases, 10th Revision; MN, malignancy; NA, not applicable; NC, not calculated; NHL, non-Hodgkin lymphoma; Obs, observed; SIR, standardised incidence ratio; SEER, Surveillance, Epidemiology, and End Results; SMR, standardised mortality ratio.

Table 4 Standardised mortality ratios (US population referent) and rate ratios for select outcomes* by employment duration (lagged 10 years)

Underlying cause (ICD-10 codes)	Employment duration (years)								Trend slope†, p Value
	0-<10		10-<20		20-<30		30+		
	Obs	SMR (95% CI) SRR (95% CI)	Obs	SMR (95% CI) SRR (95% CI)	Obs	SMR (95% CI) SRR (95% CI)	Obs	SMR (95% CI) SRR (95% CI)	
MN oesophagus (C15)	13	1.17 (0.62 to 2.00) (Reference)	28	1.72 (1.14 to 2.48) 2.43 (1.07 to 5.50)	53	1.40 (1.05 to 1.83) 1.17 (0.56 to 2.41)	19	1.18 (0.71 to 1.84) 0.60 (0.27 to 1.35)	-2.14×10 ⁻⁶ , 0.141
MN stomach (C16)	12	0.80 (0.41 to 1.40) (Reference)	18	0.92 (0.54 to 1.45) 0.33 (0.08 to 1.43)	47	1.07 (0.79 to 1.43) 0.39 (0.10 to 1.55)	33	1.53 (1.06 to 2.15) 0.40 (0.10 to 1.58)	3.06×10 ⁻⁷ , 0.822
MN intestine (C17-C18)	27	0.86 (0.57 to 1.26) (Reference)	52	1.27 (0.95 to 1.67) 1.16 (0.38 to 3.54)	171	1.42 (1.22 to 1.65) 0.62 (0.27 to 1.44)	76	1.28 (1.01 to 1.60) 0.40 (0.17 to 0.94)	-7.54×10 ⁻⁶ , <0.001
MN rectum (C19-C21)	13	1.48 (0.79 to 2.54) (Reference)	19	1.58 (0.95 to 2.46) 0.99 (0.33 to 2.97)	37	1.35 (0.95 to 1.86) 0.61 (0.24 to 1.52)	20	1.52 (0.93 to 2.34) 0.43 (0.16 to 1.14)	-1.61×10 ⁻⁶ , 0.001
MN lung (C33-C34)	123	1.02 (0.85 to 1.22) (Reference)	184	1.03 (0.88 to 1.19) 1.32 (0.97 to 1.80)	523	1.14 (1.05 to 1.24) 1.24 (0.91 to 1.68)	216	1.12 (0.98 to 1.28) 0.80 (0.59 to 1.08)	-8.83×10 ⁻⁶ , 0.216
MN prostate (C61)	24	1.39 (0.89 to 2.07) (Reference)	23	1.08 (0.68 to 1.62) 0.66 (0.31 to 1.41)	148	1.10 (0.93 to 1.29) 0.84 (0.47 to 1.50)	87	1.01 (0.81 to 1.25) 0.69 (0.39 to 1.22)	-2.03×10 ⁻⁶ , 0.192
MN kidney (C64-C66)	12	1.10 (0.57 to 1.92) (Reference)	18	1.24 (0.73 to 1.95) 0.61 (0.26 to 1.48)	47	1.43 (1.05 to 1.90) 1.25 (0.58 to 2.69)	17	1.19 (0.69 to 1.91) 0.70 (0.29 to 1.67)	-1.05×10 ⁻⁷ , 0.924
MN bladder and other urinary (C67-C68)	8	1.05 (0.45 to 2.08) (Reference)	7	0.65 (0.26 to 1.34) 0.25 (0.08 to 0.79)	46	1.08 (0.79 to 1.45) 1.15 (0.49 to 2.70)	23	0.94 (0.60 to 1.41) 1.03 (0.38 to 2.83)	2.58×10 ⁻⁶ , 0.258
MN brain and other nervous (C47, C70-C72)	12	0.65 (0.34 to 1.13) (Reference)	15	0.88 (0.49 to 1.46) 0.80 (0.30 to 2.19)	32	1.17 (0.80 to 1.65) 1.48 (0.60 to 3.68)	14	1.47 (0.80 to 2.46) 1.52 (0.53 to 4.34)	1.01×10 ⁻⁶ , 0.118
NHL (C46.3, C82-C85, C88.0, C88.3, C91.4, C96)	18	0.98 (0.58 to 1.55) (Reference)	9	0.51 (0.23 to 0.96) 1.18 (0.41 to 3.45)	63	1.35 (1.04 to 1.73) 1.15 (0.60 to 2.22)	33	1.47 (1.01 to 2.06) 1.04 (0.51 to 2.15)	-7.39×10 ⁻⁸ , 0.849
Leukaemia (C91.0-C91.3, C91.5-C91.9, C92-C95)	18	0.91 (0.54 to 1.44) (Reference)	23	1.36 (0.86 to 2.05) 2.24 (0.92 to 5.50)	54	1.11 (0.83 to 1.45) 1.36 (0.65 to 2.87)	27	1.06 (0.70 to 1.54) 1.13 (0.48 to 2.67)	-5.10×10 ⁻⁹ , 0.997
Multiple myeloma (C88.7, C88.9, C90)	5	0.84 (0.27 to 1.96) (Reference)	<5	0.52 (0.14 to 1.34) 0.56 (0.11 to 2.82)	22	0.97 (0.61 to 1.47) 1.59 (0.47 to 5.41)	11	0.99 (0.49 to 1.77) 1.25 (0.33 to 4.75)	5.27×10 ⁻⁷ , 0.407
COPD (J40-J44)	33	0.78 (0.54 to 1.10) (Reference)	38	0.69 (0.49 to 0.94) 1.07 (0.60 to 1.91)	185	0.70 (0.60 to 0.81) 1.03 (0.67 to 1.60)	111	0.75 (0.62 to 0.91) 0.83 (0.53 to 1.31)	-2.80×10 ⁻⁶ , 0.005

*Excluding a priori causes with total observations <20.

†Cause-specific deaths per year of employment-person-year.

COPD, chronic obstructive pulmonary disease; ICD-10, International Classification of Diseases, 10th Revision; MN, malignancy; NHL, non-Hodgkin lymphoma; Obs, observed; SMR, standardised mortality ratio; SRR, standardised rate ratio.

effect on a priori outcomes (results not shown). Finally, there was no apparent trend in increasing risk with employment duration; however, negative trends in COPD and colorectal cancer SRRs were evident (table 4). Subsequent sensitivity analyses revealed that SRRs were largely dependent on selection of cut-points and lag periods (results not shown).

DISCUSSION

This study is among the largest examining cancer risk in career firefighters. The pooled approach and long follow-up period improved risk estimates relative to previous studies. With few exceptions, there was little evidence of significant cancer risk heterogeneity across fire departments or age groups. Furthermore, sensitivity analyses did not suggest the potential for significant bias from including short-term workers, prevalent hires, or person-time in the open-ended age-group (85+ years). Despite notable differences in the analytical approaches, we observed remarkable similarities between mortality and incidence analyses. Additionally, the results of incidence analyses were not significantly affected by the choice of including multiple primaries or only the first cancer diagnosis. The lack of significant differences in results between fire departments, end-points, and analytic techniques suggest that the pooled study findings are robust and generalisable to similar firefighter populations.

We observed decreases in many non-malignant diseases that suggest improved health in these firefighters compared with the general population. This finding is not surprising given health requirements for entering and remaining in the fire service. Nevertheless, there was a modest excess in overall cancer mortality and incidence brought about by excess solid cancers at several sites of a priori interest. With few exceptions, our results are consistent with those previously reported and similar to SREs presented in the meta-analysis by LeMasters *et al.*¹⁴ Nevertheless, we found little evidence of excess cancers of the testes, brain and lymphohematopoietic systems, which is contrary to the synthesis by LeMasters *et al.*¹⁴ and subsequently published studies.^{8 11}

We observed about a twofold increase in malignant mesothelioma mortality and incidence compared with the US population. Malignant mesothelioma is largely attributable to asbestos exposure, with sparse evidence of other causes.²² Excess malignant mesothelioma in US firefighters was not previously described; however, excess incidence was recently observed in Nordic firefighters aged 70+ years,²³ and increased risk of asbestos-induced pulmonary and pleural fibrosis was reported in a study of New York City firefighters.²⁴ Although firefighter exposures to asbestos are known, the absence of previous reports of malignant mesothelioma is not surprising given the rarity and extremely long latency (20–40 years) of the disease. The average time between the date first employed and the date of diagnosis in the current study was 45 years; therefore, firefighting exposure-induced disease may be discernible only after lengthy follow-up. Also, previous studies have been hindered by the lack of specific codes for mesothelioma deaths before ICD-10.

We observed excess digestive cancers, mainly of oesophageal and colorectal sites. Information on occupational causes is sparse, although there is limited evidence suggesting asbestos and diesel exhaust exposures may be weakly associated with gastrointestinal cancers.^{25 26} Still, the relation between these hazardous exposures and digestive cancers appears small compared to the effects of other factors such as diet, obesity, physical activity, tobacco use and alcohol consumption.^{22 27} We also

found increased risk of oral, pharyngeal and laryngeal cancers, compared with the US population. Similar to digestive cancers, important risk factors for these sites are tobacco and alcohol consumption, with lesser evidence that exposures to wood dusts, smoke, asbestos, PAHs and acid mists may also increase risk.^{22 28 29}

Some insight into the degree of a potential bias from the lack of controlling for lifestyle factors can be gained from previous surveillance of firefighter behaviours. For example, the prevalence of smoking among current firefighters appears less than the general population, and is decreasing,^{30–33} a trend that is consistent with observed decreases in non-malignant smoking-related diseases (eg, COPD, stroke) but contradictory to excess digestive, oral and respiratory cancers. As another example, previous studies suggest there is increased obesity among firefighters compared with the general population.^{34–36} Obesity, or a dietary intake that is high in meat, fat, or overall caloric intake could contribute increased gastric or colorectal cancer risk, although concomitant elevations in health outcomes that are more strongly related to these factors (eg, ischaemic heart disease, diabetes mellitus, hypertension and stroke) were not found. Last, information on alcohol consumption within the fire service is sparse and inconsistent.^{37–40} Some studies suggest that firefighter behaviours may differ from the general population, although it is not clear that any perceived behavioural difference is sufficient to explain disparities in alcohol-related health outcomes. In the current study, the information on non-malignant and potentially alcohol-related mortality was at conflict; there was excess mortality from cirrhosis and other chronic liver disease, but fewer than expected alcoholism deaths. Alternate explanations for increased cirrhosis mortality may be exposures to chemical toxins or infectious disease,^{41–43} which may also account for excess acute renal dysfunction, a disease that is more common among those with chronic liver disease.

Fewer than 4% of firefighters in our study were women. There was evidence of excess female bladder and breast cancers; however, only bladder cancer mortality and incidence reached statistical significance. Modest excess bladder cancer has been observed in some occupations involving known or suspected bladder carcinogens (eg, PAHs, and diesel exhaust), yet contrary to our findings, risk patterns by occupation tend not to differ by gender.²² There is little evidence linking female breast cancer to workplace exposures; however, prolonged shift work may be a risk factor (and to a lesser extent a risk factor for prostate, colon and endometrial cancers).² Moreover, similar findings had not been reported previously, although increased risk of Hodgkin lymphoma and cancers of the cervix and thyroid among women firefighters (n=2017) was recently described.¹¹ Given the small sample and the lack of confirmatory results, our findings on female outcomes merit cautious interpretation.

Excess bladder and prostate cancer incidence was found among firefighters less than 65 years of age. Interestingly, the prostate cancer excess was limited to ages between 45 years and 59 years, which was consistent with recent observations in Nordic firefighters.²³ Similar mortality patterns were not observed. These cancers have relatively high survival; therefore, the underlying cause of death may be an inferior risk measure compared to cancer diagnoses. The early onset of these cancers suggests an association with firefighting. Prostate and bladder cancer diagnoses can occur following routine screening.^{44 45} As an alternative explanation, differences in medical screening (eg, prostate-specific antigen tests) among firefighters compared to the general population could have contributed to the observed excess. Data on cancer screening practices are lacking; however,

it is plausible that screening may be more frequent among firefighters with improved healthcare availability and heightened cancer awareness.

There was little evidence of increasing cancer risk with increasing employment; however, there were notable analytical shortcomings that merit discussion. First, rather than specifying cut-points and an exposure lag period specific to each outcome, we applied cut-points (10, 20 and 30 years) used in earlier studies^{5 9 46} and a common exposure lag period (10 years) to all outcomes; these choices were found to be influential in subsequent sensitivity analyses. Second, our methods have limited capability to account for HWE or other sources of bias that may have masked a dose response. Last, employment duration may poorly represent exposure potential given that some jobs are prone to lower exposures compared with others. For these reasons, a detailed exposure assessment is underway to support multivariable regression modelling for improved dose-response analyses.

Death certificates and registry data used in the current study are imperfect measures of cancer risk. In the absence of a national cancer registry, coverage is limited geographically; therefore, cases occurring outside catchment areas would be missed. Cases occurring before the registries attained comprehensive coverage have also been missed. Mortality analyses have the advantage of broader temporal and spatial coverage, but may poorly characterise cancers with relatively high survival (eg, cancers of the breast, bladder, testes and larynx). Finally, there may have been errors in tracing which can also bias study results. Although errors in ascertainment cannot be ruled out, our use of multiple information sources and end points, and the low numbers of participants lost to follow-up or moving out of catchment areas, act to minimise these errors.

CONCLUSION

In this first phase of examining health effects in career firefighters, we report on mortality and cancer incidence among nearly 30 000 career firefighters followed from 1950 through 2009. Compared with the US population, we found small to moderate increases in risk for several cancer sites and for all cancers combined, stemming mostly from excess malignancies of the respiratory, digestive and urinary systems in otherwise healthy individuals. Our findings are consistent with previous studies and strengthen evidence of a relation between firefighters' occupational exposure and cancer. We found a previously unreported twofold excess of malignant mesothelioma among firefighters. Given that asbestos is the only known causal agent for malignant mesothelioma, and firefighter exposures are probable, the excess is likely to be a causal association.

This report provides the foundation for subsequent analyses of firefighter risks, some of which are ongoing. In upcoming research, detailed employment histories (eg, number and types of fire runs) and institutional knowledge (eg, use of respiratory protection and source capture ventilation of diesel exhaust) will be used to derive exposure metrics to more accurately examine dose response. Future regression modelling will also enable examination of temporal effects that are poorly suited to life-table analyses, such as time since first exposure. Expansion and continued follow-up of this cohort would enhance future analyses, particularly among women and non-Caucasian firefighters.

Acknowledgements This study was made possible through the continued cooperation of the men and women serving the Chicago, Philadelphia and San Francisco fire departments. We especially wish to acknowledge those who provided key assistance: Chicago (José Santiago, Tom Ryan, John McNalis, Mark Edingburg, Hugh Russell and Richard Edgeworth); Philadelphia (Lloyd Ayers, Henry

Costo, William Gault and Thomas Garrity); San Francisco (Joanne Hayes-White, Ginny Franklin, Tom O'Connor, Rhab Boughn and Tony Stefani); and within the US Fire Administration (William Troup and Glenn Gains). We acknowledge NIOSH staff and their contractors for contributing to the collection, coding and management of study data. We also wish to thank Grace LeMasters and Paul Demers for their valued counsel during manuscript development.

Contributors RDD participated in design, data collection, analysis and manuscript development. TLK conceived the study and participated in design and data collection. JHY participated in design, data collection and analysis. MMD, TRH, DB, SHZ, JJB and KMW participated in design and data collection. LEP participated in design and critical appraisal. All authors participated in the interpretation and presentation of results and have read and approved the final manuscript.

Funding Research funding was provided by the National Institute for Occupational Safety and Health (NIOSH) by intramural award under the National Occupational Research Agenda (NORA), and by the US Fire Administration (USFA). This research was also supported, in part, by the intramural research programme of the National Institutes of Health (NIH), National Cancer Institute (NCI).

Competing interests None.

Ethics approval This research was approved by the Institutional Review Boards of the National Institute for Occupational Safety and Health (NIOSH) and the National Cancer Institute (NCI). Approvals for cancer registry access were granted by 11 states (ie, Arizona, California, Florida, Illinois, Indiana, Michigan, Nevada, New Jersey, Oregon, Pennsylvania and Washington). Approvals were also granted by vital records centres for death certificates maintained in 25 states (Alaska, Arizona, Arkansas, California, Florida, Georgia, Hawaii, Idaho, Illinois, Indiana, Iowa, Massachusetts, Michigan, Minnesota, Mississippi, New Jersey, New York, Ohio, Oklahoma, Oregon, Pennsylvania, Texas, Virginia, Washington and Wisconsin). The state public health entities provided vital status information in accordance with state policies, and disclaim responsibility for any analyses, interpretations, or conclusions herein.

Provenance and peer review Not commissioned; externally peer reviewed.

Data sharing statement Data were supplied by the Illinois Department of Public Health and the Bureau of Health Statistics and Research, Pennsylvania Department of Health. These public health entities specifically disclaim responsibility for any analyses, interpretations, or conclusions. The Florida cancer incidence data used in this report were collected by the Florida Cancer Data System (FCDS) under contract with the Florida Department of Health (FDOH). The views expressed herein are solely those of the authors and do not necessarily reflect those of the FCDS or FDOH. The collection of cancer incidence data used in this study was also supported by the California Department of Public Health as part of the statewide cancer reporting programme mandated by California Health and Safety Code Section 103885; the National Cancer Institute's Surveillance, Epidemiology and End Results Program under contract N01-PC-35136 awarded to the Northern California Cancer Center, contract N01-PC-35139 awarded to the University of Southern California, and contract N02-PC-15105 awarded to the Public Health Institute; and the Centres for Disease Control and Prevention's National Program of Cancer Registries, under agreement #U55/CCR921930-02 awarded to the Public Health Institute. The ideas and opinions expressed herein are those of the authors, and endorsement by the State of California, Department of Public Health, the National Cancer Institute, and the Centres for Disease Control and Prevention or their contractors and subcontractors is neither intended nor to be inferred.

REFERENCES

- 1 Karter MJ, Stein GP. *U.S. Fire Department Profile Through 2011* NFPA Fire Analysis and Research Division, Quincy, MA: National Fire Protection Association, 2012:1–36.
- 2 International Agency for Research on Cancer. IARC working group on the evaluation of carcinogenic risks to humans. Painting, firefighting, and shiftwork. *IARC Monogr Eval Carcinog Risks Hum* 2010;98:9–764.
- 3 Aronson KJ, Tomlinson GA, Smith L. Mortality among fire fighters in metropolitan Toronto. *Am J Ind Med* 1994;26:89–101.
- 4 Tomling G, Gustavsson P, Hogstedt C. Mortality and cancer incidence in Stockholm fire fighters. *Am J Ind Med* 1994;25:219–28.
- 5 Vena JE, Fiedler RC. Mortality of a municipal-worker cohort: IV. Fire fighters. *Am J Ind Med* 1987;11:671–84.
- 6 Demers PA, Heyer NJ, Rosenstock L. Mortality among firefighters from three northwestern United States cities. *Br J Ind Med* 1992;49:664–70.
- 7 Bates MN. Registry-based case-control study of cancer in California firefighters. *Am J Ind Med* 2007;50:339–44.
- 8 Kang D, Davis LK, Hunt P, et al. Cancer incidence among male Massachusetts firefighters, 1987–2003. *Am J Ind Med* 2008;51:329–35.
- 9 Baris D, Garrity TJ, Telles JL, et al. Cohort mortality study of Philadelphia firefighters. *Am J Ind Med* 2001;39:463–76.
- 10 Beaumont JJ, Chu GS, Jones JR, et al. An epidemiologic study of cancer and other causes of mortality in San Francisco firefighters. *Am J Ind Med* 1991;19:357–72.

- 11 Ma F, Fleming LE, Lee DJ, *et al.* Cancer incidence in Florida professional firefighters, 1981 to 1999. *J Occup Environ Med* 2006;48:883–8.
- 12 Ma F, Fleming LE, Lee DJ, *et al.* Mortality in Florida professional firefighters, 1972 to 1999. *Am J Ind Med* 2005;47:509–17.
- 13 Heyer N, Weiss NS, Demers P, *et al.* Cohort mortality study of Seattle fire fighters: 1945–1983. *Am J Ind Med* 1990;17:493–504.
- 14 LeMasters GK, Genaidy AM, Succop P, *et al.* Cancer risk among firefighters: A review and meta-analysis of 32 studies. *J Occup Environ Med* 2006;48:1189–202.
- 15 Fritz A, Percy C, Jack A, *et al.* eds. *International classification of diseases for oncology*. 3rd edn. (ICD-O-3). Geneva: World Health Organization, 2000.
- 16 Schubauer-Berigan MK, Hein MJ, Raudabaugh WM, *et al.* Update of the NIOSH life table analysis system: a person-years analysis program for the windows computing environment. *Am J Ind Med* 2011;54:915–24.
- 17 Robinson CF, Schnorr TM, Cassinelli RT II, *et al.* Tenth revision U.S. mortality rates for use with the NIOSH Life Table Analysis System. *J Occup Environ Med* 2006;48:662–7.
- 18 Surveillance, Epidemiology, and End Results (SEER) Program (www.seer.cancer.gov) SEER*Stat Database: Incidence –all available registries (1985 to 2009), National Cancer Institute, DCCPS, Surveillance Research Program, Surveillance Systems Branch, released April 2012, based on the November 2011 submission.
- 19 Merrill RM, Sloan A, Novilla LB. Understanding population-based site-specific cancer incidence rates in the USA. *J Cancer Educ* 2012;27:263–8.
- 20 Applebaum KM, Malloy EJ, Eisen EA. Reducing healthy worker survivor bias by restricting date of hire in a cohort study of Vermont granite workers. *Occup Environ Med* 2007;64:681–7.
- 21 Sorahan T. Bladder cancer risks in workers manufacturing chemicals for the rubber industry. *Occup Med (Lond)* 2008;58:496–501.
- 22 Schottenfeld D, Fraumeni JF. eds. *Cancer epidemiology and prevention*. 3rd edn. Oxford; New York, NY: Oxford University Press, 2006.
- 23 Demers PA, Martinsen JI, Kjaerheim K, *et al.* Cancer incidence among Nordic firefighters [abstract]. *Am J Epidemiol* 2011;173:S191.
- 24 Markowitz SB, Garibaldi K, Lillis R, *et al.* Asbestos and fire fighting. *Ann N Y Acad Sci* 1991;643:573–81.
- 25 Benbrahim-Tallaa L, Baan RA, Grosse Y, *et al.* Carcinogenicity of diesel-engine and gasoline-engine exhausts and some nitroarenes. *Lancet Oncol* 2012;13:663–4.
- 26 International Agency for Research on Cancer. IARC Working Group on the Evaluation of Carcinogenic Risks to Humans. Arsenic, metals, fibres, and dusts. Volume 100C. A review of human carcinogens. *IARC Monogr Eval Carcinog Risks Hum* 2012;100:11–465.
- 27 International Agency for Research on Cancer. IARC Working Group on the Evaluation of Carcinogenic Risks to Humans. Personal habits and indoor combustions. Volume 100 E. A review of human carcinogens. *IARC Monogr Eval Carcinog Risks Hum* 2012;100:1–538.
- 28 Paget-Bailly S, Cyr D, Luce D. Occupational exposures to asbestos, polycyclic aromatic hydrocarbons and solvents, and cancers of the oral cavity and pharynx: a quantitative literature review. *Int Arch Occup Environ Health* 2012;85:341–51.
- 29 Paget-Bailly S, Cyr D, Luce D. Occupational exposures and cancer of the larynx-systematic review and meta-analysis. *J Occup Environ Med* 2012; 54:71–84.
- 30 Haddock CK, Jitnarin N, Poston WS, *et al.* Tobacco use among firefighters in the central United States. *Am J Ind Med* 2011;54:697–706.
- 31 Lee DJ, LeBlanc W, Fleming LE, *et al.* Trends in US smoking rates in occupational groups: the National Health Interview Survey 1987–1994. *J Occup Environ Med* 2004;46:538–48.
- 32 Poston WS, Haddock CK, Jitnarin N, *et al.* A national qualitative study of tobacco use among career firefighters and department health personnel. *Nicotine Tob Res* 2012;14:734–41.
- 33 Lee DJ, Fleming LE, Arheart KL, *et al.* Smoking rate trends in U.S. occupational groups: the 1987 to 2004 National Health Interview Survey. *J Occup Environ Med* 2007;49:75–81.
- 34 Munir F, Clemes S, Houdmont J, *et al.* Overweight and obesity in UK firefighters. *Occup Med (Lond)* 2012;62:362–5.
- 35 Poston WS, Jitnarin N, Haddock CK, *et al.* Obesity and injury-related absenteeism in a population-based firefighter cohort. *Obesity (Silver Spring)* 2011; 19:2076–81.
- 36 Tsimenakis AJ, Christophi CA, Burrell JW, *et al.* The obesity epidemic and future emergency responders. *Obesity (Silver Spring)* 2009;17:1648–50.
- 37 Haddock CK, Jahnke SA, Poston WS, *et al.* Alcohol use among firefighters in the Central United States. *Occup Med (Lond)* 2012;62:661–4.
- 38 Carey MG, Al-Zaiti SS, Dean GE, *et al.* Sleep problems, depression, substance use, social bonding, and quality of life in professional firefighters. *J Occup Environ Med* 2011;53:928–33.
- 39 Parker DA, Harford TC. The epidemiology of alcohol consumption and dependence across occupations in the United States. *Alcohol Health Res World* 1992;16:97–105.
- 40 Boxer PA, Wild D. Psychological distress and alcohol use among fire fighters. *Scand J Work Environ Health* 1993;19:121–5.
- 41 Leiss JK, Ratcliffe JM, Lyden JT, *et al.* Blood exposure among paramedics: incidence rates from the national study to prevent blood exposure in paramedics. *Ann Epidemiol* 2006;16:720–5.
- 42 Rischitelli G, Harris J, McCauley L, *et al.* The risk of acquiring hepatitis B or C among public safety workers: a systematic review. *Am J Prev Med* 2001;20:299–306.
- 43 Boal WL, Hales T, Ross CS. Blood-borne pathogens among firefighters and emergency medical technicians. *Prehosp Emerg Care* 2005;9:236–47.
- 44 Potosky AL, Miller BA, Albertsen PC, *et al.* The role of increasing detection in the rising incidence of prostate-cancer. *JAMA* 1995;273:548–52.
- 45 Moyer VA. Screening for bladder cancer: U.S. Preventive Services Task Force recommendation statement. *Ann Intern Med* 2011;155:246–51.
- 46 Demers PA, Checkoway H, Vaughan TL, *et al.* Cancer incidence among firefighters in Seattle and Tacoma, Washington (United States). *Cancer Causes Control* 1994;5:129–35.

Supplementary (online only) data for:

Mortality and cancer incidence in a pooled cohort of U.S. firefighters from San Francisco, Chicago, and Philadelphia (1950-2009)

Robert D. Daniels^{1*}, Travis L. Kubale¹, James H. Yiin¹, Matthew M. Dahm¹, Thomas R. Hales¹, Dalsu Baris², Shelia H. Zahm², James J. Beaumont³, Kathleen M. Waters¹, Lynne E. Pinkerton¹

¹ Division of Surveillance, Hazard Evaluations, and Field Studies, National Institute for Occupational Safety and Health, Cincinnati, Ohio, USA

² Division of Cancer Epidemiology and Genetics, National Cancer Institute, Gaithersburg, Maryland, USA.

³ UC Davis Department of Public Health Sciences, Davis, California, USA

*Corresponding author; NIOSH, DSHEFS, IWSB, 4676 Columbia Parkway, Mailstop R-13, Cincinnati, OH 45226. Email: rtd2@cdc.gov; Phone: (513) 458-7178, Fax: (513) 841-4470

Table S1. Cancer registry information

Cancer Registry	State	NPCR participant	SEER participant	Statewide Data Collection Began	No. of cases (%) [*]	No. deaths (%) [†]
Arizona Cancer Registry	Arizona	Yes	No	1981	105 (2.4)	157 (1.7)
California cancer Registry	California	No	Yes	1988	881 (19.7)	1477 (15.9)
Florida Cancer Data System	Florida	Yes	No	1981	301 (6.7)	532 (5.7)
Illinois State Cancer Registry	Illinois	Yes	No	1986	1783 (40.0)	2760 (29.7)
Indiana Sate Cancer Registry	Indiana	Yes	No	1987	24 (0.5)	117 (1.3)
Michigan Cancer Surveillance Program	Michigan	Yes	Yes	1985	31 (0.7)	60 (0.6)
Nevada Central Cancer Registry	Nevada	Yes	No	1995	35 (0.8)	72 (0.8)
New Jersey State Cancer Registry	New Jersey	No	Yes	1979	216 (4.8)	391 (4.2)
Oregon State Cancer Registry	Oregon	Yes	No	1996	9 (0.2)	21 (0.2)
Pennsylvania Cancer Registry	Pennsylvania	Yes	No	1985	1071 (24.0)	3196 (34.4)
Washington State Cancer Registry	Washington	Yes	Yes	1994	5 (0.1)	19 (0.2)
NA	Others	NA	NA	NA	NA	488 (5.3)

^{*}Includes persons with multiple primary tumors and excludes duplicate reporting ($n=4461$).

[†]Number and percentage of deaths among cohort members with known death state ($n=9290$).

NA, not applicable; NPCR, National Program of Cancer Registries; SEER, Surveillance, Epidemiology, and End Results Program

Table S2. Firefighter employment and vital status by gender (1950-2009).

Employment	Men		Women	
	Alive	Deceased	Alive	Deceased
Employed 1+ years	15969	11798	875	23
Employed <1 year	1031	204	90	3
Total	17000	12002	965	26

Table S3. Cause-specific standardized mortality ratios compared to the US population by fire department (1950-2009, $n=29993$, 858938 PYAR)

Minor ID*	Cause	All		San Francisco		Chicago		Philadelphia	
		Obs	SMR (95% CI)	Obs	SMR (95% CI)	Obs	SMR (95% CI)	Obs	SMR (95% CI)
1-92	All causes	12028	0.99 (0.97 to 1.01)	2074	0.84 (0.80 to 0.87)	5944	1.05 (1.02 to 1.08)	4010	1.01 (0.98 to 1.04)
1-2	Tuberculosis	12	0.35 (0.18 to 0.61)	<5	0.30 (0.04 to 1.09)	<5	0.25 (0.07 to 0.65)	6	0.51 (0.19 to 1.10)
1	Respiratory tuberculosis	11	0.35 (0.18 to 0.63)	<5	0.33 (0.04 to 1.19)	<5	0.21 (0.04 to 0.61)	6	0.55 (0.20 to 1.21)
2	Other tuberculosis	<5	0.33 (0.01 to 1.85)	0	NC	<5	0.69 (0.02 to 3.83)	0	NC
3-38	All Cancers	3285	1.14 (1.10 to 1.18)	578	1.00 (0.92 to 1.09)	1670	1.22 (1.16 to 1.28)	1037	1.11 (1.05 to 1.18)
3-6	MN buccal cavity & pharynx	94	1.40 (1.13 to 1.72)	22	1.69 (1.06 to 2.56)	41	1.28 (0.92 to 1.73)	31	1.42 (0.96 to 2.01)
3	MN lip	<5	0.80 (0.02 to 4.44)	<5	3.70 (0.09 to 20.61)	0	NC	0	NC
4	MN tongue	25	1.61 (1.04 to 2.38)	6	1.99 (0.73 to 4.33)	11	1.49 (0.74 to 2.66)	8	1.56 (0.67 to 3.08)
5	MN other buccal	25	1.43 (0.93 to 2.12)	5	1.45 (0.47 to 3.38)	8	0.97 (0.42 to 1.91)	12	2.10 (1.08 to 3.66)
6	MN pharynx	43	1.31 (0.95 to 1.77)	10	1.59 (0.76 to 2.92)	22	1.39 (0.87 to 2.10)	11	1.04 (0.52 to 1.86)
7-13	MN digestive & peritoneum	928	1.26 (1.18 to 1.34)	179	1.21 (1.04 to 1.40)	463	1.33 (1.21 to 1.46)	286	1.19 (1.06 to 1.34)
7	MN esophagus	113	1.39 (1.14 to 1.67)	23	1.45 (0.92 to 2.18)	58	1.47 (1.12 to 1.90)	32	1.22 (0.84 to 1.72)
8	MN stomach	110	1.10 (0.91 to 1.33)	25	1.23 (0.80 to 1.82)	53	1.14 (0.86 to 1.50)	32	0.97 (0.66 to 1.36)
9	MN intestine	326	1.30 (1.16 to 1.44)	56	1.09 (0.82 to 1.41)	157	1.33 (1.13 to 1.55)	113	1.38 (1.13 to 1.65)
10	MN rectum	89	1.45 (1.16 to 1.78)	20	1.59 (0.97 to 2.46)	47	1.65 (1.21 to 2.20)	22	1.08 (0.68 to 1.63)
11	MN biliary, liver, gall bladder	107	1.30 (1.06 to 1.57)	21	1.28 (0.79 to 1.96)	60	1.51 (1.15 to 1.95)	26	0.98 (0.64 to 1.44)
12	MN pancreas	168	1.13 (0.97 to 1.32)	33	1.11 (0.76 to 1.56)	81	1.15 (0.91 to 1.43)	54	1.12 (0.84 to 1.46)
13	MN peritoneum, other & unspecified	<29	1.42 (0.80 to 2.35)	<5	0.47 (0.01 to 2.60)	7	1.42 (0.57 to 2.93)	7	2.01 (0.81 to 4.15)
14-17	MN respiratory system	1096	1.10 (1.04 to 1.17)	147	0.75 (0.63 to 0.88)	594	1.24 (1.14 to 1.34)	355	1.11 (1.00 to 1.24)
14	MN larynx	<46	1.26 (0.91 to 1.69)	<5	0.44 (0.09 to 1.28)	26	1.55 (1.01 to 2.27)	15	1.32 (0.74 to 2.18)
15	MN trachea, bronchus, lung	1046	1.10 (1.04 to 1.17)	142	0.76 (0.64 to 0.89)	566	1.23 (1.13 to 1.34)	338	1.11 (0.99 to 1.23)
16	MN pleura	<5	0.81 (0.10 to 2.93)	0	NC	<5	1.71 (0.21 to 6.19)	0	NC
17	MN other respiratory sites	<5	0.66 (0.18 to 1.69)	<5	1.71 (0.21 to 6.18)	0	NC	<5	1.01 (0.12 to 3.66)
18	MN breast	8	1.39 (0.60 to 2.73)	<5	2.43 (0.50 to 7.11)	<5	1.29 (0.35 to 3.29)	<5	0.70 (0.02 to 3.92)
19-22	MN female genital organs	0	NC	0	NC	0	NC	0	NC
19	MN cervix	0	NC	0	NC	0	NC	0	NC
20	MN other parts of uterus	0	NC	0	NC	0	NC	0	NC
21	MN ovary	0	NC	0	NC	0	NC	0	NC
22	MN other female genital	0	NC	0	NC	0	NC	0	NC
23-24	MN male genital organs	286	1.07 (0.95 to 1.20)	52	0.90 (0.67 to 1.18)	152	1.23 (1.05 to 1.45)	82	0.94 (0.75 to 1.17)
23	MN prostate	282	1.09 (0.96 to 1.22)	51	0.91 (0.68 to 1.19)	152	1.28 (1.08 to 1.50)	79	0.94 (0.74 to 1.17)
24	MN other male genital	<5	0.47 (0.13 to 1.20)	<5	0.63 (0.02 to 3.50)	0	NC	<5	1.02 (0.21 to 2.99)
25-26	MN urinary	178	1.13 (0.97 to 1.31)	29	0.89 (0.60 to 1.28)	96	1.30 (1.06 to 1.59)	53	1.03 (0.77 to 1.34)

Minor ID*	Cause	All		San Francisco		Chicago		Philadelphia	
		Obs	SMR (95% CI)	Obs	SMR (95% CI)	Obs	SMR (95% CI)	Obs	SMR (95% CI)
25	MN kidney	94	1.29 (1.05 to 1.58)	13	0.90 (0.48 to 1.54)	56	1.62 (1.23 to 2.11)	25	1.05 (0.68 to 1.56)
26	MN bladder & other urinary	84	0.99 (0.79 to 1.22)	16	0.88 (0.50 to 1.43)	40	1.02 (0.73 to 1.39)	28	1.00 (0.67 to 1.45)
27-34	MN other & unspecified	397	1.09 (0.98 to 1.20)	89	1.23 (0.99 to 1.51)	181	1.04 (0.89 to 1.20)	127	1.06 (0.89 to 1.27)
27	MN skin	56	0.94 (0.71 to 1.22)	14	1.19 (0.65 to 1.99)	22	0.78 (0.49 to 1.19)	20	1.02 (0.62 to 1.57)
28	Mesothelioma	12	2.00 (1.03 to 3.49)	<5	2.41 (0.50 to 7.05)	8	2.73 (1.18 to 5.38)	<5	0.55 (0.01 to 3.04)
29	MN eye	<5	2.28 (0.62 to 5.84)	0	NC	<5	4.95 (1.35 to 12.67)	0	NC
30	MN brain & other nervous system	73	1.01 (0.79 to 1.27)	16	1.16 (0.66 to 1.89)	34	0.98 (0.68 to 1.37)	23	0.96 (0.61 to 1.44)
31	MN thyroid gland	<5	0.56 (0.11 to 1.62)	<5	0.93 (0.02 to 5.17)	<5	0.39 (0.01 to 2.18)	<5	0.57 (0.01 to 3.15)
32	MN bone	9	1.16 (0.53 to 2.19)	<5	0.65 (0.02 to 3.63)	6	1.66 (0.61 to 3.61)	<5	0.76 (0.09 to 2.74)
33	MN connective tissue	10	0.68 (0.32 to 1.24)	<5	0.70 (0.08 to 2.52)	<5	0.56 (0.15 to 1.44)	<5	0.84 (0.23 to 2.14)
34	MN other & unspecified sites	230	1.16 (1.02 to 1.32)	52	1.31 (0.98 to 1.72)	102	1.08 (0.88 to 1.32)	76	1.18 (0.93 to 1.48)
35-38	MN lymphatic & hematopoietic	298	1.07 (0.95 to 1.20)	57	1.01 (0.77 to 1.31)	139	1.05 (0.88 to 1.24)	102	1.12 (0.92 to 1.36)
35	Non-Hodgkin lymphoma	123	1.17 (0.97 to 1.40)	25	1.19 (0.77 to 1.75)	53	1.06 (0.80 to 1.39)	45	1.32 (0.96 to 1.77)
36	Hodgkin lymphoma	11	0.68 (0.34 to 1.22)	<5	0.66 (0.08 to 2.37)	6	0.79 (0.29 to 1.72)	<5	0.55 (0.11 to 1.60)
37	Leukemia & aleukemia	122	1.10 (0.91 to 1.31)	23	1.02 (0.65 to 1.53)	61	1.17 (0.90 to 1.50)	38	1.05 (0.74 to 1.44)
38	Multiple myeloma	42	0.89 (0.64 to 1.20)	7	0.74 (0.30 to 1.52)	19	0.84 (0.50 to 1.30)	16	1.06 (0.60 to 1.72)
39-41	Benign & unspecified neoplasms	40	1.07 (0.77 to 1.46)	8	1.06 (0.46 to 2.10)	17	0.97 (0.56 to 1.55)	15	1.23 (0.69 to 2.04)
39	Benign eye, brain, other nervous system	<5	0.92 (0.25 to 2.35)	<5	1.16 (0.03 to 6.45)	<5	0.97 (0.12 to 3.52)	<5	0.70 (0.02 to 3.88)
40	Unspecified eye, brain, other nervous system	<15	0.96 (0.52 to 1.61)	5	1.75 (0.57 to 4.09)	5	0.72 (0.23 to 1.68)	<5	0.84 (0.23 to 2.14)
41	Other benign & unspecified	<25	1.20 (0.75 to 1.82)	<5	0.53 (0.06 to 1.90)	10	1.17 (0.56 to 2.15)	10	1.69 (0.81 to 3.10)
42	Diabetes mellitus	175	0.72 (0.62 to 0.83)	17	0.35 (0.20 to 0.55)	89	0.77 (0.62 to 0.94)	69	0.89 (0.69 to 1.12)
43-46	Diseases of blood & blood-forming organs	50	1.11 (0.82 to 1.46)	11	1.16 (0.58 to 2.07)	24	1.13 (0.73 to 1.69)	15	1.04 (0.58 to 1.71)
43	Pernicious Anemia	0	NC	0	NC	0	NC	0	NC
44	Other & unspecified anemia	16	1.01 (0.58 to 1.64)	<5	0.60 (0.07 to 2.15)	10	1.34 (0.64 to 2.47)	<5	0.79 (0.21 to 2.02)
45	Coagulation & hemorrhagic conditions	7	0.77 (0.31 to 1.60)	<5	0.55 (0.01 to 3.07)	<5	0.92 (0.25 to 2.36)	<5	0.69 (0.08 to 2.50)
46	Other diseases of blood-forming organs	27	1.37 (0.91 to 2.00)	8	1.91 (0.83 to 3.77)	10	1.09 (0.52 to 2.01)	9	1.43 (0.65 to 2.72)
47-48	Mental, psychoneurotic, and personality disorders	120	0.81 (0.68 to 0.97)	20	0.65 (0.39 to 1.00)	60	0.87 (0.66 to 1.12)	40	0.85 (0.61 to 1.16)
47	Alcoholism	31	0.61 (0.41 to 0.86)	<5	0.44 (0.12 to 1.13)	19	0.74 (0.45 to 1.16)	8	0.49 (0.21 to 0.96)
48	Other mental disorders	89	0.92 (0.74 to 1.14)	16	0.73 (0.42 to 1.19)	41	0.94 (0.67 to 1.28)	32	1.04 (0.71 to 1.47)

Minor ID*	Cause	All		San Francisco		Chicago		Philadelphia	
		Obs	SMR (95% CI)	Obs	SMR (95% CI)	Obs	SMR (95% CI)	Obs	SMR (95% CI)
49-50	Nervous system disorders	187	0.80 (0.69 to 0.93)	43	0.86 (0.62 to 1.15)	82	0.76 (0.61 to 0.95)	62	0.83 (0.63 to 1.06)
49	Multiple sclerosis	<10	0.57 (0.21 to 1.23)	0	NC	5	0.99 (0.32 to 2.30)	<5	0.28 (0.01 to 1.58)
50	Other nervous system diseases	181	0.82 (0.70 to 0.94)	43	0.89 (0.64 to 1.20)	77	0.75 (0.59 to 0.94)	61	0.86 (0.65 to 1.10)
51-55	Diseases of the heart	4289	0.99 (0.96 to 1.02)	720	0.80 (0.74 to 0.86)	2124	1.06 (1.02 to 1.11)	1445	1.01 (0.95 to 1.06)
51	Rheumatic heart disease	51	0.89 (0.66 to 1.17)	6	0.54 (0.20 to 1.17)	24	0.90 (0.58 to 1.35)	21	1.08 (0.67 to 1.65)
52	Ischemic heart disease	3619	1.01 (0.98 to 1.04)	598	0.80 (0.74 to 0.87)	1812	1.10 (1.05 to 1.15)	1209	1.02 (0.96 to 1.08)
53	Chronic Diseases of endocardium	56	0.99 (0.75 to 1.29)	15	1.22 (0.68 to 2.02)	25	0.97 (0.63 to 1.44)	16	0.87 (0.49 to 1.41)
54	Hypertension w/heart disease	111	0.84 (0.69 to 1.02)	24	0.89 (0.57 to 1.32)	54	0.89 (0.67 to 1.16)	33	0.76 (0.52 to 1.06)
55	Other heart disease	452	0.88 (0.81 to 0.97)	77	0.72 (0.57 to 0.90)	209	0.88 (0.76 to 1.00)	166	1.00 (0.86 to 1.17)
56-58	Other diseases circulatory system	967	0.91 (0.85 to 0.97)	201	0.88 (0.76 to 1.01)	439	0.91 (0.83 to 1.00)	327	0.93 (0.83 to 1.04)
56	Hypertension w/o heart disease	46	0.85 (0.62 to 1.13)	10	0.88 (0.42 to 1.61)	25	0.98 (0.64 to 1.45)	11	0.63 (0.32 to 1.13)
57	Cerebrovascular disease	636	0.91 (0.84 to 0.98)	131	0.86 (0.72 to 1.02)	298	0.95 (0.84 to 1.06)	207	0.89 (0.77 to 1.02)
58	Diseases of the arteries, Veins, and pulmonary circulation	285	0.93 (0.82 to 1.04)	60	0.91 (0.70 to 1.18)	116	0.82 (0.68 to 0.99)	109	1.07 (0.88 to 1.30)
59-64	Diseases of the respiratory system	796	0.80 (0.74 to 0.86)	137	0.65 (0.54 to 0.76)	411	0.89 (0.81 to 0.98)	248	0.77 (0.67 to 0.87)
59	Acute respiratory infections except influenza and pneumonia	0	NC	0	NC	0	NC	0	NC
60	Influenza	<5	0.22 (0.03 to 0.81)	<5	0.51 (0.01 to 2.83)	<5	0.26 (0.01 to 1.43)	0	NC
61	Pneumonia	269	0.90 (0.79 to 1.01)	55	0.84 (0.64 to 1.10)	129	0.95 (0.79 to 1.12)	85	0.86 (0.69 to 1.06)
62	Chronic obstructive pulmonary diseases	367	0.72 (0.65 to 0.80)	57	0.53 (0.40 to 0.69)	206	0.87 (0.75 to 0.99)	104	0.63 (0.52 to 0.77)
63	Asthma	<10	0.36 (0.14 to 0.74)	<5	0.51 (0.06 to 1.84)	5	0.55 (0.18 to 1.27)	0	NC
64	Pneumoconiosis & other respiratory diseases	151	0.97 (0.82 to 1.14)	22	0.67 (0.42 to 1.01)	70	0.97 (0.76 to 1.23)	59	1.18 (0.90 to 1.52)
65-68	Diseases of the digestive system	572	1.13 (1.04 to 1.23)	110	1.11 (0.91 to 1.34)	277	1.16 (1.02 to 1.30)	185	1.12 (0.96 to 1.29)
65	Diseases of the stomach & duodenum	64	1.16 (0.89 to 1.48)	12	1.06 (0.54 to 1.84)	29	1.15 (0.77 to 1.66)	23	1.23 (0.78 to 1.85)
66	Hernia & intestinal obstruction	<23	0.69 (0.41 to 1.07)	<5	0.17 (0.00 to 0.93)	8	0.64 (0.28 to 1.27)	10	1.08 (0.52 to 1.99)
67	Cirrhosis & other chronic liver disease	299	1.26 (1.12 to 1.41)	65	1.47 (1.13 to 1.87)	144	1.25 (1.05 to 1.47)	90	1.15 (0.93 to 1.42)

Minor ID*	Cause	All		San Francisco		Chicago		Philadelphia	
		Obs	SMR (95% CI)	Obs	SMR (95% CI)	Obs	SMR (95% CI)	Obs	SMR (95% CI)
68	Other diseases digestive system	190	1.04 (0.89 to 1.19)	32	0.85 (0.58 to 1.21)	96	1.11 (0.90 to 1.36)	62	1.04 (0.79 to 1.33)
69-77	Diseases of the genitourinary system	201	1.00 (0.87 to 1.15)	36	0.84 (0.59 to 1.16)	91	0.98 (0.79 to 1.21)	74	1.14 (0.89 to 1.43)
69	Acute glomerulonephritis nephrotic syndrome and acute renal failure	32	1.56 (1.07 to 2.20)	7	1.61 (0.65 to 3.31)	14	1.46 (0.80 to 2.46)	11	1.67 (0.83 to 2.99)
70	Chronic and unspecified nephritis and renal failure & other renal sclerosis	91	0.86 (0.69 to 1.05)	9	0.40 (0.18 to 0.77)	48	0.96 (0.71 to 1.28)	34	1.00 (0.70 to 1.40)
71	Kidney infection	9	0.75 (0.34 to 1.42)	<5	1.14 (0.24 to 3.34)	<5	0.77 (0.21 to 1.96)	<5	0.48 (0.06 to 1.72)
72	Urinary system calculi	<5	1.29 (0.35 to 3.30)	<5	1.51 (0.04 to 8.43)	<5	0.73 (0.02 to 4.06)	<5	1.86 (0.23 to 6.73)
73	Prostate hyperplasia	11	1.27 (0.63 to 2.28)	<5	1.84 (0.50 to 4.71)	5	1.50 (0.49 to 3.51)	<5	0.64 (0.08 to 2.30)
74	Other diseases of male genital organs	<5	0.34 (0.01 to 1.87)	0	NC	0	NC	<5	0.99 (0.03 to 5.53)
75	Diseases of breast	0	NC	0	NC	0	NC	0	NC
76	diseases female genital organs	0	NC	0	NC	0	NC	0	NC
77	Other genitourinary diseases	53	1.12 (0.84 to 1.46)	12	1.18 (0.61 to 2.05)	19	0.87 (0.52 to 1.36)	22	1.44 (0.90 to 2.18)
78-79	diseases of skin & subcutaneous tissue	13	1.18 (0.63 to 2.02)	<5	1.77 (0.48 to 4.53)	<5	0.77 (0.21 to 1.97)	5	1.41 (0.46 to 3.29)
78	Skin & subcutaneous infections	5	1.21 (0.39 to 2.83)	<5	2.35 (0.28 to 8.48)	<5	1.03 (0.12 to 3.71)	<5	0.75 (0.02 to 4.18)
79	Other diseases skin & subcutaneous tissue	8	1.17 (0.50 to 2.30)	<5	1.42 (0.17 to 5.13)	<5	0.62 (0.07 to 2.23)	<5	1.81 (0.49 to 4.63)
80-82	diseases musculoskeletal & connective tissue	15	0.53 (0.30 to 0.88)	<5	0.35 (0.04 to 1.25)	7	0.53 (0.21 to 1.09)	6	0.66 (0.24 to 1.45)
80	Arthritis & spondylitis	<5	0.18 (0.02 to 0.65)	0	NC	0	NC	<5	0.55 (0.07 to 1.98)
81	Osteomyelitis & periostitis	<5	0.91 (0.19 to 2.67)	0	NC	<5	1.27 (0.15 to 4.60)	<5	0.97 (0.02 to 5.41)
82	Other diseases musculoskeletal	10	0.73 (0.35 to 1.35)	<5	0.72 (0.09 to 2.61)	5	0.76 (0.25 to 1.78)	<5	0.69 (0.14 to 2.01)
83	Symptoms & ill-defined conditions	161	1.31 (1.11 to 1.53)	<5	0.08 (0.01 to 0.30)	54	0.92 (0.69 to 1.19)	105	2.62 (2.14 to 3.17)
84-88	Accidents	524	0.86 (0.79 to 0.94)	87	0.78 (0.62 to 0.96)	289	0.99 (0.88 to 1.11)	148	0.73 (0.62 to 0.86)
84	Transportation accidents	160	0.56 (0.48 to 0.65)	29	0.57 (0.38 to 0.82)	89	0.65 (0.52 to 0.80)	42	0.43 (0.31 to 0.58)
85	Accidental poisoning	45	0.88 (0.64 to 1.18)	6	0.70 (0.26 to 1.51)	26	0.98 (0.64 to 1.43)	13	0.82 (0.43 to 1.39)
86	Accidental falls	113	1.31 (1.08 to 1.58)	27	1.49 (0.98 to 2.17)	62	1.57 (1.20 to 2.01)	24	0.84 (0.54 to 1.26)
87	Other accidents	197	1.17 (1.01 to 1.34)	24	0.77 (0.50 to 1.15)	109	1.34 (1.10 to 1.61)	64	1.14 (0.88 to 1.46)
88	Medical causes	9	0.63 (0.29 to 1.20)	<5	0.35 (0.01 to 1.95)	<5	0.44 (0.09 to 1.29)	5	1.08 (0.35 to 2.52)

Minor ID*	Cause	All		San Francisco		Chicago		Philadelphia	
		Obs	SMR (95% CI)	Obs	SMR (95% CI)	Obs	SMR (95% CI)	Obs	SMR (95% CI)
89-91	Violence	251	0.78 (0.69 to 0.89)	57	1.00 (0.76 to 1.30)	120	0.75 (0.63 to 0.90)	74	0.71 (0.55 to 0.89)
89	Suicide	193	0.87 (0.75 to 1.00)	50	1.24 (0.92 to 1.63)	86	0.81 (0.65 to 1.00)	57	0.76 (0.58 to 0.99)
90	Homicide	58	0.59 (0.45 to 0.76)	7	0.43 (0.17 to 0.88)	34	0.65 (0.45 to 0.91)	17	0.57 (0.33 to 0.91)
91	Terrorism	0	NC	0	NC	0	NC	0	NC
92	Other & unspecified causes	370	1.15 (1.04 to 1.27)	39	0.63 (0.45 to 0.86)	182	1.14 (0.98 to 1.32)	149	1.49 (1.26 to 1.75)

*Minor cause-of-death category in NIOSH Life Table Analysis System (LTAS) 92-cause rate file: <http://www.cdc.gov/niosh/ltras/rates.html>.

MN, malignancy; NC, not calculated; Obs, observed; SMR, standardized mortality ratio.

Table S4. Cancer-specific standardized incidence ratios compared to the US population by fire department (1985-2009, $n=24453$, 403152 PYAR)*

Minor ID [†]	Cause	All		San Francisco		Chicago		Philadelphia	
		Obs	SIR (95% CI)	Obs	SIR (95% CI)	Obs	SIR (95% CI)	Obs	SIR (95% CI)
1-41	All Cancers	4461	1.09 (1.06 to 1.12)	855	1.14 (1.06 to 1.22)	2186	1.09 (1.04 to 1.13)	1420	1.07 (1.01 to 1.12)
1-4	MN buccal cavity & pharynx	174	1.39 (1.19 to 1.62)	43	1.96 (1.42 to 2.64)	81	1.32 (1.05 to 1.64)	50	1.20 (0.89 to 1.58)
1	MN lip	<20	1.11 (0.65 to 1.78)	10	3.71 (1.78 to 6.82)	5	0.68 (0.22 to 1.58)	<5	0.38 (0.05 to 1.38)
2	MN tongue	52	1.74 (1.30 to 2.28)	9	1.70 (0.78 to 3.23)	23	1.58 (1.00 to 2.38)	20	1.97 (1.20 to 3.04)
3	MN other buccal	46	1.24 (0.91 to 1.65)	11	1.67 (0.83 to 2.99)	25	1.37 (0.89 to 2.03)	10	0.81 (0.39 to 1.49)
4	MN pharynx	59	1.39 (1.06 to 1.79)	13	1.77 (0.94 to 3.02)	28	1.33 (0.88 to 1.92)	18	1.29 (0.76 to 2.03)
5-6	MN colon & rectum	537	1.18 (1.08 to 1.28)	100	1.20 (0.98 to 1.46)	267	1.20 (1.06 to 1.35)	170	1.14 (0.97 to 1.32)
5	MN large intestine	381	1.21 (1.09 to 1.34)	72	1.24 (0.97 to 1.56)	186	1.21 (1.04 to 1.40)	123	1.19 (0.99 to 1.42)
6	MN rectum	156	1.11 (0.94 to 1.30)	28	1.12 (0.74 to 1.61)	81	1.17 (0.93 to 1.46)	47	1.02 (0.75 to 1.35)
7-12	MN digestive other & peritoneum	393	1.16 (1.05 to 1.28)	69	1.11 (0.87 to 1.41)	200	1.20 (1.04 to 1.37)	124	1.13 (0.94 to 1.35)
7	MN esophagus	90	1.62 (1.31 to 2.00)	15	1.49 (0.83 to 2.45)	49	1.79 (1.33 to 2.37)	26	1.44 (0.94 to 2.12)
8	MN stomach	93	1.15 (0.93 to 1.40)	12	0.82 (0.42 to 1.43)	50	1.24 (0.92 to 1.64)	31	1.18 (0.80 to 1.68)
9	MN small intestine	<18	1.15 (0.67 to 1.85)	<5	1.49 (0.41 to 3.82)	5	0.69 (0.22 to 1.61)	8	1.66 (0.72 to 3.28)
10	MN biliary, liver, gall bladder	85	1.10 (0.88 to 1.36)	16	1.12 (0.64 to 1.82)	47	1.22 (0.89 to 1.62)	22	0.89 (0.56 to 1.35)
11	MN pancreas	90	0.96 (0.77 to 1.18)	19	1.10 (0.66 to 1.72)	43	0.94 (0.68 to 1.27)	28	0.92 (0.61 to 1.32)
12	MN peritoneum, other & unspecified	<20	1.10 (0.65 to 1.74)	<5	1.01 (0.21 to 2.96)	6	0.74 (0.27 to 1.62)	9	1.70 (0.78 to 3.23)
13-16	MN respiratory & Intrathoracic	813	1.16 (1.08 to 1.24)	91	0.72 (0.58 to 0.88)	463	1.33 (1.21 to 1.46)	259	1.14 (1.00 to 1.29)
13	MN larynx	84	1.50 (1.19 to 1.85)	10	1.02 (0.49 to 1.88)	42	1.51 (1.08 to 2.03)	32	1.73 (1.18 to 2.44)
14	MN trachea, bronchus, lung	716	1.12 (1.04 to 1.21)	81	0.70 (0.56 to 0.87)	409	1.30 (1.17 to 1.43)	226	1.09 (0.96 to 1.25)
15	MN pleura	0	NC	0	NC	0	NC	0	NC
16	MN other respiratory & intrathoracic	<17	1.37 (0.73 to 2.34)	0	NC	12	2.55 (1.32 to 4.45)	<5	0.32 (0.01 to 1.80)
17	MN breast	26	1.26 (0.82 to 1.85)	6	1.23 (0.45 to 2.67)	14	1.19 (0.65 to 1.99)	6	1.53 (0.56 to 3.32)
18-21	MN female genital organs	<5	0.62 (0.13 to 1.81)	<5	0.77 (0.02 to 4.28)	<5	0.66 (0.08 to 2.40)	0	NC
18	MN cervix uteri	<5	1.20 (0.15 to 4.33)	<5	2.06 (0.05 to 11.49)	<5	1.02 (0.03 to 5.67)	0	NC
19	MN other & unspecified parts of uterus	<5	0.56 (0.01 to 3.11)	0	NC	<5	0.86 (0.02 to 4.81)	0	NC
20	MN ovary, fallopian tube, & broad ligament	0	NC	0	NC	0	NC	0	NC
21	MN other female genital organs	0	NC	0	NC	0	NC	0	NC
22-24	MN male genital organs	1278	1.02 (0.97 to 1.08)	278	1.21 (1.07 to 1.36)	602	0.98 (0.91 to 1.07)	398	0.98 (0.89 to 1.09)
22	MN prostate	1261	1.03 (0.98 to 1.09)	276	1.22 (1.08 to 1.37)	592	0.99 (0.91 to 1.07)	393	0.99 (0.90 to 1.10)
23	MN testes	<18	0.75 (0.42 to 1.24)	<5	0.74 (0.09 to 2.67)	8	0.76 (0.33 to 1.50)	5	0.75 (0.24 to 1.75)
24	MN other & unspecified male genital organs	<5	0.26 (0.03 to 0.93)	0	NC	<5	0.53 (0.06 to 1.92)	0	NC
25-26	MN urinary organs	482	1.17 (1.06 to 1.27)	89	1.15 (0.93 to 1.42)	234	1.17 (1.02 to 1.32)	159	1.17 (1.00 to 1.37)
25	MN kidney	166	1.27 (1.09 to 1.48)	26	1.10 (0.72 to 1.61)	83	1.30 (1.04 to 1.61)	57	1.33 (1.00 to 1.72)

Minor ID [†]	Cause	All		San Francisco		Chicago		Philadelphia	
		Obs	SIR (95% CI)	Obs	SIR (95% CI)	Obs	SIR (95% CI)	Obs	SIR (95% CI)
26	MN bladder & other urinary organs	316	1.12 (1.00 to 1.25)	63	1.18 (0.91 to 1.51)	151	1.10 (0.93 to 1.29)	102	1.10 (0.90 to 1.33)
27-28	MN thyroid & other endocrine glands	28	0.91 (0.60 to 1.31)	<5	0.72 (0.20 to 1.84)	15	0.98 (0.55 to 1.61)	9	0.91 (0.42 to 1.72)
27	MN thyroid gland	25	0.87 (0.56 to 1.28)	<5	0.57 (0.12 to 1.68)	13	0.90 (0.48 to 1.55)	9	0.97 (0.44 to 1.85)
28	MN other endocrine glands	<5	1.50 (0.31 to 4.39)	<5	2.90 (0.07 to 16.18)	<5	2.03 (0.25 to 7.32)	0	NC
29-35	MN other solid cancers	275	0.97 (0.86 to 1.09)	85	1.69 (1.35 to 2.09)	104	0.76 (0.62 to 0.92)	86	0.90 (0.72 to 1.11)
29	MN bone	<13	2.62 (1.35 to 4.57)	0	NC	8	3.49 (1.51 to 6.87)	<5	2.66 (0.72 to 6.81)
30	MN melanoma (skin)	141	0.87 (0.73 to 1.03)	56	1.89 (1.43 to 2.46)	44	0.56 (0.41 to 0.76)	41	0.75 (0.54 to 1.02)
31	Kaposi sarcoma	<5	0.17 (0.05 to 0.43)	<5	0.30 (0.01 to 1.69)	<5	0.08 (0.00 to 0.47)	<5	0.23 (0.03 to 0.83)
32	MN mesothelioma	35	2.29 (1.60 to 3.19)	6	2.05 (0.75 to 4.47)	20	2.71 (1.65 to 4.18)	9	1.82 (0.83 to 3.46)
33	MN connective tissue	22	1.07 (0.67 to 1.62)	<5	1.06 (0.29 to 2.72)	10	0.99 (0.47 to 1.82)	8	1.19 (0.51 to 2.34)
34	MN brain & other nervous system	51	1.02 (0.76 to 1.34)	17	1.95 (1.14 to 3.12)	13	0.53 (0.28 to 0.91)	21	1.25 (0.77 to 1.91)
35	MN eye	<18	1.45 (0.69 to 2.66)	<5	0.81 (0.02 to 4.53)	8	2.44 (1.05 to 4.80)	<5	0.44 (0.01 to 2.44)
36-40	MN lymphohematopoietic tissue	345	0.94 (0.84 to 1.04)	68	1.00 (0.78 to 1.27)	154	0.85 (0.72 to 1.00)	123	1.02 (0.85 to 1.22)
36	Hodgkin lymphoma	<17	0.96 (0.54 to 1.59)	6	2.36 (0.87 to 5.15)	6	0.76 (0.28 to 1.65)	<5	0.58 (0.12 to 1.70)
37	Non-Hodgkin lymphoma	169	0.99 (0.84 to 1.15)	28	0.90 (0.60 to 1.30)	79	0.94 (0.75 to 1.18)	62	1.10 (0.84 to 1.41)
38	Multiple myeloma	36	0.72 (0.50 to 0.99)	9	0.97 (0.44 to 1.84)	16	0.65 (0.37 to 1.05)	11	0.68 (0.34 to 1.21)
39	Leukemia & aleukemia	100	0.94 (0.77 to 1.15)	22	1.12 (0.70 to 1.69)	43	0.83 (0.60 to 1.12)	35	1.01 (0.70 to 1.40)
40	Other lymphohematopoietic neoplasms	<27	0.97 (0.63 to 1.43)	<5	0.56 (0.12 to 1.64)	10	0.79 (0.38 to 1.45)	12	1.54 (0.79 to 2.68)
41	MN III-specified & residual	107	1.04 (0.86 to 1.26)	21	1.10 (0.68 to 1.68)	50	1.00 (0.75 to 1.32)	36	1.07 (0.75 to 1.48)

*Analysis of all occurrences of invasive cancer (i.e., multiple-cancer approach).

[†] Minor cause-of-death category in Table S4 of this appendix.

MN, malignancy; NC, not calculated; Obs, observed; SIR, standardized incidence ratio.

Table S5. Recode from ICD-O-3 codes reported by cancer incidence registries to diagnostic minor codes used in NIOSH LTAS*.

Major Category	Minor ID	Minor Category	ICD-10 Codes	ICD-O-3 Site Codes	ICD-O-3 Histology Codes
MN of buccal cavity and pharynx	1	MN of lip	C00	C000-C009	All excluding 9140, 9050-9055, and 9590-9989
	2	MN of tongue	C01, C02	C019-C029	
	3	MN of other buccal cavity	C03-C08	C039-C069, C079-C089	
	4	MN of pharynx	C09-C14	C090- C119, C129-C148	
MN of colon and rectum	5	MN of colon	C18	C180-C189	
	6	MN of rectum	C19, C20	C199, C209	
MN of other digestive organs and peritoneum	7	MN of esophagus	C15	C150- C159	
	8	MN of stomach	C16	C160-C169	
	9	MN of small intestine	C17	C170-C179	
	10	MN of biliary, liver, gall bladder	C22-C24	C220, C221, C239-C249	
	11	MN of pancreas	C25	C250-C259	
	12	MN of anus, peritoneum, other, and unspecified digestive	C21, C26, C48	C210-C212, C218, C260, C268, C269, C422, C480-C482, C488	
MN of respiratory and Intrathoracic organs	13	MN of larynx	C32	C320-C329	
	14	MN of trachea, bronchus, and lung	C33, C34	C339-C349	
	15	MN of pleura	C38.4	C384	
	16	MN of other respiratory and intrathoracic organs	C30, C31, C37, C38.0-C38.3, C38.8, C39	C300,C301, C310-C319, C379, C380-C383, C388, C390, C398, C399	
MN of breast	17	MN of breast	C50	C500-C509	
MN of female genital organs	18	MN of cervix uteri	C53	C530-C539	
	19	MN of other and unspecified parts of uterus	C54, C55, C58	C540-C549, C559, C589	
	20	MN of ovary, fallopian tube, and broad ligament	C56, 57.0-C57.4, C57.8	C569-C574, C578	

Major Category	Minor ID	Minor Category	ICD-10 Codes	ICD-O-3 Site Codes	ICD-O-3 Histology Codes
	21	MN of other and unspecified female genital organs	C51, C52, C57.7, C57.9	C510-C519, C529, C577, C579	
MN of male genital organs	22	MN of prostate	C61	C619	
	23	MN of testes	C62	C620-C629	
	24	MN of other and unspecified male genital organs	C60, C63	C600-C609, C630-C639	
MN of urinary organs	25	MN of kidney	C64-C66	C649, C659, C669	
	26	MN of bladder and other urinary organs	C67, C68, D09.0 [†]	C670-C689	
MN of thyroid and other endocrine glands	27	MN of thyroid gland	C73	C739	
	28	MN of other endocrine glands	C74, C75	C740-C749, C750-C759	
MN of other solid cancers	29	MN of bone	C40, C41	C400-C419	
	30	Malignant melanoma of skin	C43	C440-C449	8720-8790
	31	Kaposi sarcoma	C46	Not used	9140
	32	Mesothelioma	C45	Not used	9050-9055
	33	MN of connective tissue	C49	C490-C499	All excluding 9140, 9050-9055, and 9590-9989
	34	MN brain and other parts of nervous system	C47, C70-C72	C470-C479, C700-C729	
	35	MN eye	C69	C690-C699	
	Malignant neoplasms of lymphatic and hematopoietic tissue	36	Hodgkin lymphoma	C81	Not used
37		Non-Hodgkin lymphoma	C82-C85, C88.0, C88.3, C91.4, C96.0-C96.3, C96.7	Not used	9590, 9591, 9596, 9670, 9671, 9673, 9675, 9678-9680, 9684, 9687, 9688, 9689-9691, 9695, 9698-9702, 9705, 9708, 9709, 9712, 9714-9719, 9724-9729, 9735, 9737, 9738, 9740, 9750, 9754-9759, 9761, 9764, 9940
38		Multiple myeloma	C90	Not used	9731-9734
39		Leukemia and aleukemia	C91.0-C91.3, C91.5, C91.7, C91.9, C92-C95	Not used	9742, 9800, 9801, 9805, 9820, 9823, 9826, 9827, 9831-9837, 9840, 9860, 9861, 9863, 9866, 9867, 9870-9876, 9891, 9895-9897, 9910, 9920, 9930, 9931, 9945, 9946, 9948, 9963

Major Category	Minor ID	Minor Category	ICD-10 Codes	ICD-O-3 Site Codes	ICD-O-3 Histology Codes
	40	Other lymphatic and hematopoietic neoplasms	C88.2, C88.7, C88.9, C96.9, D45, D46.1-D46.4, D46.7, D46.9, D47.1, D47.3, D47.7	Not used	9751, 9760, 9762, 9950, 9960-9962, 9970, 9975, 9980, 9982-9987, 9989
Ill-specified and residual	41	MN of Ill-specified and residual sites	C44, C76, C77, C80, C97	C440-C449	All excluding 8720-8790, 9140, 9050-9055, and 9590-9989
				C760-C768, C809, C420-C424, C770-C779	All excluding 9140, 9050-9055, and 9590-9989

*Results in Table 2 of the main manuscript differ slightly from Table S3 due to classification adjustment made to better align cancer incidence diagnoses with mortality results in Table 2. Specifically, Table 2 differs from this recode by including ICD-10 codes C21 with MN rectum, C96.9 with NHL, and C88.7 and C88.9 with multiple myeloma.

†Urinary bladder incidence cases originally coded *in situ* (Behavior=2) were recoded to invasive (Behavior=3) per SEER protocol.

ICD-O-3, International Classification of Diseases for Oncology, 3rd Edition; ICD-10, International Classification of Diseases, 10th Revision; MN, malignancy; SEER, Surveillance, Epidemiology, and End Results Program

Table S6. Heterogeneity in standardized mortality ratios across fire departments with U.S. population referent (1950-2009).

Minor ID*	Underlying Cause	All fire departments [†]		San Francisco		Chicago		Philadelphia		Heterogeneity [‡] LRT p-value
		Obs	SMR (95% CI)	Obs	SMR (95% CI)	Obs	SMR (95% CI)	Obs	SMR (95% CI)	
7	MN esophagus	113	1.39 (1.15 to 1.67)	23	1.45 (0.92 to 2.18)	58	1.47 (1.12 to 1.90)	32	1.22 (0.84 to 1.72)	1.00
8	MN stomach	110	1.10 (0.91 to 1.33)	25	1.23 (0.80 to 1.82)	53	1.14 (0.86 to 1.50)	32	0.97 (0.66 to 1.36)	1.00
9	MN intestine	326	1.30 (1.16 to 1.44)	56	1.09 (0.82 to 1.41)	157	1.33 (1.13 to 1.55)	113	1.38 (1.13 to 1.65)	1.00
10	MN rectum	89	1.45 (1.16 to 1.80)	20	1.59 (0.97 to 2.46)	47	1.65 (1.21 to 2.20)	22	1.08 (0.68 to 1.63)	0.48
15	MN lung	1046	1.02 (0.81 to 1.29)	142	0.76 (0.64 to 0.89)	566	1.23 (1.13 to 1.34)	338	1.11 (0.99 to 1.23)	<0.01
23	MN prostate	282	1.05 (0.87 to 1.27)	51	0.91 (0.68 to 1.19)	152	1.28 (1.08 to 1.50)	79	0.94 (0.74 to 1.17)	0.08
25	MN kidney	94	1.23 (0.90 to 1.67)	13	0.90 (0.48 to 1.54)	56	1.62 (1.23 to 2.11)	25	1.05 (0.68 to 1.56)	0.17
26	MN bladder	84	0.99 (0.80 to 1.22)	16	0.88 (0.50 to 1.43)	40	1.02 (0.73 to 1.39)	28	1.00 (0.67 to 1.45)	1.00
30	MN brain	73	1.01 (0.80 to 1.27)	16	1.16 (0.66 to 1.89)	34	0.98 (0.68 to 1.37)	23	0.96 (0.61 to 1.44)	1.00
35	NHL	123	1.17 (0.98 to 1.40)	25	1.19 (0.77 to 1.75)	53	1.06 (0.80 to 1.39)	45	1.32 (0.96 to 1.77)	1.00
37	Leukemia	122	1.10 (0.92 to 1.32)	23	1.02 (0.65 to 1.53)	61	1.17 (0.90 to 1.50)	38	1.05 (0.74 to 1.44)	1.00
38	Multiple myeloma	42	0.89 (0.66 to 1.20)	7	0.74 (0.30 to 1.52)	19	0.84 (0.50 to 1.30)	16	1.06 (0.60 to 1.72)	1.00
62	COPD	367	0.68 (0.53 to 0.85)	57	0.53 (0.40 to 0.69)	206	0.87 (0.75 to 0.99)	104	0.63 (0.52 to 0.77)	<0.01

* Minor cause-of-death category in NIOSH Life Table Analysis System (LTAS) 92-cause rate file: <http://www.cdc.gov/niosh/ltras/rates.html>. Reporting restricted to 20 or more total cases.

[†] Results from Poisson mixed model that adjusts for fire department-specific effects as a random variable.

[‡] Results of testing against the null model (i.e., no between-department variance).

COPD, chronic obstructive pulmonary disease; ICD-10, International Classification of Diseases, 10th Revision; LRT, likelihood ratio test; MN, malignancy; NC. Not calculated; NHL, non-Hodgkin lymphoma; Obs, observed; SMR, standardized mortality ratio.

Table S7. Heterogeneity in standardized incidence ratios by department compared to the US population for cancers of *a priori* interest (1985-2009).

Minor ID*	Cause	All departments [†]		San Francisco		Chicago		Philadelphia		Heterogeneity [‡] LRT p-value
		Obs	SIR (95% CI)	Obs	SIR (95% CI)	Obs	SIR (95% CI)	Obs	SIR (95% CI)	
5	MN large intestine	381	1.21 (1.09 to 1.34)	72	1.24 (0.97 to 1.56)	186	1.21 (1.04 to 1.40)	123	1.19 (0.99 to 1.42)	1.00
6	MN rectum	156	1.11 (0.95 to 1.30)	28	1.12 (0.74 to 1.61)	81	1.17 (0.93 to 1.46)	47	1.02 (0.75 to 1.35)	1.00
7	MN esophagus	90	1.62 (1.32 to 1.99)	15	1.49 (0.83 to 2.45)	49	1.79 (1.33 to 2.37)	26	1.44 (0.94 to 2.12)	1.00
8	MN stomach	93	1.15 (0.94 to 1.40)	12	0.82 (0.42 to 1.43)	50	1.24 (0.92 to 1.64)	31	1.18 (0.80 to 1.68)	1.00
14	MN lung	716	1.01 (0.76 to 1.35)	81	0.70 (0.56 to 0.87)	409	1.30 (1.17 to 1.43)	226	1.10 (0.96 to 1.25)	<0.001
17	MN breast	26	1.26 (0.86 to 1.85)	6	1.23 (0.45 to 2.67)	14	1.19 (0.65 to 1.99)	6	1.53 (0.56 to 3.32)	1.00
22	MN prostate	1261	1.05 (0.95 to 1.17)	276	1.22 (1.08 to 1.37)	592	0.99 (0.91 to 1.07)	393	0.99 (0.90 to 1.10)	0.078
25	MN kidney	166	1.27 (1.09 to 1.48)	26	1.10 (0.72 to 1.61)	83	1.30 (1.04 to 1.61)	57	1.33 (1.00 to 1.72)	1.00
26	MN bladder	316	1.11 (1.00 to 1.25)	63	1.18 (0.91 to 1.51)	151	1.10 (0.93 to 1.29)	102	1.10 (0.90 to 1.33)	1.00
34	MN brain	51	1.07 (0.59 to 1.95)	17	1.95 (1.14 to 3.12)	13	0.53 (0.28 to 0.91)	21	1.25 (0.77 to 1.91)	0.007
37	NHL	169	0.99 (0.85 to 1.15)	28	0.90 (0.60 to 1.30)	79	0.94 (0.75 to 1.18)	62	1.10 (0.84 to 1.41)	1.00
38	Multiple myeloma	36	0.72 (0.52 to 0.99)	9	0.97 (0.44 to 1.84)	16	0.65 (0.37 to 1.05)	11	0.68 (0.34 to 1.21)	1.00
39	Leukemia	100	0.94 (0.78 to 1.15)	22	1.12 (0.70 to 1.69)	43	0.83 (0.60 to 1.12)	35	1.01 (0.70 to 1.40)	1.00

* Minor cause shown in Table S4 of this appendix. Reporting restricted to 20 or more total cases.

[†] Results from Poisson mixed model that adjusts for department-specific effects as a random variable.

[‡] Results of testing against the null model (i.e., no between-department variance).

COPD, chronic obstructive pulmonary disease; LRT, likelihood ratio test; MN, malignancy; NHL, non-Hodgkin lymphoma; Obs, observed; SMR, standardized mortality ratio.

Table S8. Standardized mortality ratios using State mortality rates for causes of death of *a priori* interest (1950-2009).

Minor ID*	Underlying Cause	All departments [†]		San Francisco (California rates)		Chicago (Illinois rates)		Philadelphia (Pennsylvania rates)		Heterogeneity [‡] LRT p-value
		Obs	SMR (95% CI)	Obs	SMR (95% CI)	Obs	SMR (95% CI)	Obs	SMR (95% CI)	
7	MN esophagus	113	1.28 (1.06 to 1.54)	23	1.57 (1.00 to 2.36)	58	1.32 (1.00 to 1.71)	32	1.07 (0.73 to 1.52)	1.00
8	MN stomach	110	1.05 (0.87 to 1.26)	25	1.20 (0.78 to 1.77)	53	1.10 (0.82 to 1.43)	32	0.90 (0.61 to 1.27)	1.00
9	MN intestine	326	1.19 (1.07 to 1.33)	56	1.22 (0.92 to 1.59)	157	1.17 (1.00 to 1.37)	113	1.19 (0.98 to 1.44)	1.00
10	MN rectum	89	1.30 (1.01 to 1.68)	20	1.67 (1.02 to 2.59)	47	1.45 (1.07 to 1.93)	22	0.92 (0.58 to 1.40)	0.32
15	MN lung	1046	1.06 (0.92 to 1.23)	142	0.87 (0.73 to 1.02)	566	1.20 (1.10 to 1.30)	338	1.10 (0.99 to 1.22)	0.02
23	MN prostate	282	1.04 (0.86 to 1.26)	51	0.90 (0.67 to 1.19)	152	1.26 (1.07 to 1.48)	79	0.92 (0.73 to 1.15)	0.08
25	MN kidney	94	1.24 (0.96 to 1.61)	13	0.97 (0.51 to 1.66)	56	1.51 (1.14 to 1.96)	25	1.06 (0.68 to 1.56)	0.37
26	MN bladder	84	0.94 (0.76 to 1.17)	16	0.91 (0.52 to 1.47)	40	0.98 (0.70 to 1.33)	28	0.92 (0.61 to 1.33)	1.00
30	MN brain	73	1.05 (0.83 to 1.32)	16	1.13 (0.64 to 1.83)	34	1.01 (0.70 to 1.42)	23	1.05 (0.66 to 1.57)	1.00
35	NHL	123	1.11 (0.93 to 1.33)	25	1.17 (0.76 to 1.73)	53	0.99 (0.74 to 1.30)	45	1.25 (0.91 to 1.68)	1.00
37	Leukemia	122	1.07 (0.90 to 1.28)	23	1.07 (0.68 to 1.60)	61	1.10 (0.84 to 1.42)	38	1.03 (0.73 to 1.41)	1.00
38	Multiple myeloma	42	0.91 (0.67 to 1.23)	7	0.75 (0.30 to 1.55)	19	0.86 (0.52 to 1.34)	16	1.07 (0.61 to 1.74)	1.00
62	COPD	367	0.71 (0.55 to 0.93)	57	0.54 (0.41 to 0.69)	206	0.93 (0.81 to 1.07)	104	0.69 (0.56 to 0.84)	<0.01

* Minor cause-of-death category in NIOSH Life Table Analysis System (LTAS) 92-cause rate file: <http://www.cdc.gov/niosh/ltras/rates.html>. Reporting restricted to 20 or more total cases.

[†] Results from Poisson mixed model that adjusts for department-specific effects as a random variable.

[‡] Results of testing against the null model (i.e., no between-department variance).

COPD, chronic obstructive pulmonary disease; LRT, likelihood ratio test; MN, malignancy; NC, not calculated; NHL, non-Hodgkin lymphoma; Obs, observed; SMR, standardized mortality ratio.

Table S9. Standardized mortality ratios by hire type, with U.S. population referent (1950-2009).

Minor ID*	Underlying Cause	All hires		Hired on or after 1950 (incident hires only)			
		employed 1+ years		All		Employed 1+ years	
		Obs	SMR (95% CI)	Obs	SMR (95% CI)	Obs	SMR (95% CI)
1-92	All causes	11821	0.99 (0.97 to 1.01)	4441	0.91 (0.89 to 0.94)	4236	0.90 (0.88 to 0.93)
3-38	All cancers	3231	1.14 (1.10 to 1.18)	1454	1.12 (1.07 to 1.18)	1400	1.12 (1.06 to 1.18)
7	MN esophagus	111	1.39 (1.14 to 1.67)	67	1.51 (1.17 to 1.92)	65	1.52 (1.17 to 1.94)
8	MN stomach	109	1.11 (0.91 to 1.34)	45	1.29 (0.94 to 1.73)	44	1.31 (0.95 to 1.76)
9	MN intestine	322	1.30 (1.16 to 1.45)	120	1.18 (0.98 to 1.41)	116	1.18 (0.98 to 1.42)
10	MN rectum	88	1.45 (1.17 to 1.79)	36	1.56 (1.09 to 2.15)	35	1.57 (1.09 to 2.19)
15	MN lung	1028	1.10 (1.03 to 1.17)	476	1.05 (0.96 to 1.15)	458	1.05 (0.95 to 1.15)
18	MN breast	7	1.25 (0.50 to 2.57)	6	1.68 (0.62 to 3.67)	5	1.47 (0.48 to 3.43)
23	MN prostate	278	1.09 (0.96 to 1.22)	95	1.18 (0.96 to 1.45)	91	1.18 (0.95 to 1.45)
24	MN other male genital organs	<5	0.48 (0.13 to 1.23)	<5	0.42 (0.05 to 1.52)	<5	0.43 (0.05 to 1.57)
25	MN kidney	91	1.28 (1.03 to 1.57)	53	1.46 (1.10 to 1.92)	50	1.43 (1.06 to 1.89)
26	MN bladder& other urinary	84	1.00 (0.80 to 1.24)	27	0.89 (0.59 to 1.30)	27	0.93 (0.61 to 1.35)
30	MN brain & other nervous	73	1.03 (0.81 to 1.29)	36	0.90 (0.63 to 1.25)	36	0.94 (0.66 to 1.30)
35	NHL	120	1.16 (0.96 to 1.39)	56	1.06 (0.80 to 1.38)	53	1.04 (0.78 to 1.37)
37	Leukemia	121	1.11 (0.92 to 1.33)	55	1.10 (0.83 to 1.43)	54	1.12 (0.84 to 1.46)
38	Multiple myeloma	42	0.90 (0.65 to 1.22)	17	0.76 (0.44 to 1.22)	17	0.79 (0.46 to 1.26)
62	COPD	361	0.72 (0.65 to 0.80)	145	0.75 (0.63 to 0.88)	139	0.75 (0.63 to 0.88)

* Minor cause-of-death category in NIOSH Life Table Analysis System (LTAS) 92-cause rate file: <http://www.cdc.gov/niosh/ltras/rates.html>.

COPD, chronic obstructive pulmonary disease; ICD-10, International Classification of Diseases, 10th Revision; MN, malignancy; NHL, non-Hodgkin lymphoma; Obs, observed; SMR, standardized mortality ratio.

Table S10. Heterogeneity in standardized mortality ratios by age compared to the US population for causes of death of *a priori* interest (1950-2009).

Minor ID*	Underlying Cause	All Ages [†]		Age 65-85+		Age 17-64		Heterogeneity [‡] LRT p-value
		Obs	SMR (95% CI)	Obs	SMR (95% CI)	Obs	SMR (95% CI)	
7	MN esophagus	113	1.39 (1.15 to 1.67)	62	1.36 (1.05 to 1.75)	51	1.41 (1.05 to 1.86)	1.00
8	MN stomach	110	1.02 (0.76 to 1.37)	74	1.33 (1.05 to 1.68)	36	0.81 (0.57 to 1.12)	0.07
9	MN intestine	326	1.23 (1.03 to 1.48)	231	1.42 (1.24 to 1.61)	95	1.07 (0.87 to 1.31)	0.08
10	MN rectum	89	1.45 (1.18 to 1.78)	51	1.49 (1.11 to 1.96)	38	1.40 (0.99 to 1.92)	1.00
15	MN lung	1046	1.09 (1.01 to 1.18)	662	1.17 (1.09 to 1.27)	384	0.99 (0.90 to 1.10)	0.29
23	MN prostate	282	1.09 (0.97 to 1.22)	245	1.08 (0.95 to 1.23)	37	1.10 (0.77 to 1.51)	1.00
25	MN kidney	94	1.29 (1.06 to 1.58)	46	1.15 (0.84 to 1.53)	48	1.47 (1.09 to 1.95)	1.00
26	MN bladder	84	0.97 (0.74 to 1.26)	65	1.01 (0.78 to 1.29)	19	0.90 (0.54 to 1.41)	0.35
30	MN brain	73	1.01 (0.80 to 1.27)	34	1.20 (0.83 to 1.67)	39	0.89 (0.63 to 1.21)	1.00
35	NHL	123	1.15 (0.90 to 1.46)	86	1.37 (1.09 to 1.69)	37	0.88 (0.62 to 1.21)	0.39
37	Leukemia	122	1.10 (0.92 to 1.31)	81	1.17 (0.93 to 1.46)	41	0.98 (0.70 to 1.33)	1.00
38	Multiple myeloma	42	0.89 (0.66 to 1.20)	29	0.93 (0.63 to 1.34)	13	0.80 (0.42 to 1.37)	1.00
62	COPD	367	0.70 (0.59 to 0.82)	313	0.77 (0.69 to 0.86)	54	0.53 (0.40 to 0.70)	0.06

* Minor cause-of-death category in NIOSH Life Table Analysis System (LTAS) 92-cause rate file: <http://www.cdc.gov/niosh/ltras/rates.html>. Reporting restricted to 20 or more total cases.

[†]Results from Poisson mixed model that adjusts for age-specific effects as a random variable.

[‡]Results of testing against the null model (i.e., no between-age group variance). Age defined as all ages in 5-year periods.

COPD, chronic obstructive pulmonary disease; LRT, likelihood ratio test; NC, not calculated; MN, malignancy; NHL, non-Hodgkin lymphoma; Obs, observed; SMR, standardized mortality ratio.

Table S11. Heterogeneity in standardized incidence ratios by age compared to the US population for cancers of *a priori* interest (1985-2009).

Minor ID*	Underlying Cause	All Ages [†]		Age 65-85+		Age 17-64		Heterogeneity [‡] LRT p-value
		Obs	SIR (95% CI)	Obs	SIR (95% CI)	Obs	SIR (95% CI)	
5	MN large intestine	381	1.21 (1.09 to 1.34)	269	1.20 (1.06 to 1.35)	112	1.23 (1.01 to 1.48)	1.00
6	MN rectum	156	1.11 (0.95 to 1.30)	97	1.12 (0.91 to 1.37)	59	1.10 (0.83 to 1.41)	1.00
7	MN esophagus	90	1.60 (1.27 to 2.03)	55	1.59 (1.20 to 2.07)	35	1.68 (1.17 to 2.34)	0.40
8	MN stomach	93	1.15 (0.94 to 1.40)	62	1.15 (0.88 to 1.48)	31	1.14 (0.77 to 1.61)	1.00
14	MN lung	716	1.12 (1.05 to 1.21)	494	1.13 (1.03 to 1.23)	222	1.12 (0.98 to 1.28)	1.00
17	MN breast	26	1.26 (0.86 to 1.85)	<5	0.53 (0.11 to 1.56)	23	1.53 (0.97 to 2.30)	1.00
22	MN prostate	1261	1.10 (0.95 to 1.28)	835	0.96 (0.90 to 1.03)	426	1.21 (1.10 to 1.33)	<0.001
25	MN kidney	166	1.27 (1.09 to 1.48)	87	1.17 (0.94 to 1.44)	79	1.41 (1.12 to 1.76)	1.00
26	MN bladder	316	1.11 (0.90 to 1.36)	219	1.04 (0.91 to 1.19)	97	1.33 (1.08 to 1.62)	0.002
34	MN brain	51	1.02 (0.77 to 1.34)	25	1.04 (0.67 to 1.54)	26	1.00 (0.65 to 1.46)	1.00
37	NHL	169	0.98 (0.82 to 1.17)	107	1.10 (0.90 to 1.33)	62	0.84 (0.64 to 1.07)	0.351
38	Multiple myeloma	36	0.72 (0.52 to 0.99)	26	0.77 (0.50 to 1.13)	10	0.60 (0.29 to 1.11)	1.00
39	Leukemia	100	0.94 (0.78 to 1.15)	61	0.86 (0.66 to 1.10)	39	1.12 (0.80 to 1.53)	1.00

* Minor cause shown in Table S4 of this appendix. Reporting restricted to 20 or more total cases.

[†] Results from Poisson mixed model that adjusts for age-specific effects as a random variable.

[‡] Results of testing against the null model (i.e., no between-age group variance). Age defined as all ages in 5-year periods.

LRT, likelihood ratio test; MN, malignancy; NHL, non-Hodgkin lymphoma; Obs, observed; SIR, standardized incidence ratio.

Table S12. Standardized mortality ratios with censoring at age ≥ 85 for causes of death of *a priori* interest (1950-2009), US population referent.

Minor ID*	Underlying Cause	All ages		Age <85	
		Obs	SMR (95% CI)	Obs	SMR (95% CI)
7	MN esophagus	113	1.39 (1.14 to 1.67)	108	1.38 (1.13 to 1.66)
8	MN stomach	110	1.10 (0.91 to 1.33)	102	1.07 (0.87 to 1.30)
9	MN intestine	326	1.30 (1.16 to 1.44)	301	1.29 (1.15 to 1.44)
10	MN rectum)	89	1.45 (1.16 to 1.78)	82	1.40 (1.12 to 1.74)
15	MN lung	1046	1.10 (1.04 to 1.17)	1001	1.09 (1.03 to 1.16)
18	MN breast	8	1.39 (0.60 to 2.73)	7	1.27 (0.51 to 2.62)
23	MN prostate	282	1.09 (0.96 to 1.22)	235	1.08 (0.95 to 1.23)
24	MN other male genital	<5	0.47 (0.13 to 1.20)	<5	0.48 (0.13 to 1.24)
25	MN kidney	94	1.29 (1.05 to 1.58)	91	1.32 (1.06 to 1.62)
26	MN bladder	84	0.99 (0.79 to 1.22)	75	1.00 (0.78 to 1.25)
30	MN brain	73	1.01 (0.79 to 1.27)	71	1.00 (0.78 to 1.26)
35	NHL	123	1.17 (0.97 to 1.40)	113	1.15 (0.95 to 1.39)
37	Leukemia	122	1.10 (0.91 to 1.31)	116	1.13 (0.93 to 1.36)
38	Multiple myeloma	42	0.89 (0.64 to 1.20)	39	0.88 (0.63 to 1.20)
62	COPD	367	0.72 (0.65 to 0.80)	321	0.71 (0.64 to 0.79)

* Minor cause-of-death category in NIOSH Life Table Analysis System (LTAS) 92-cause rate file:

<http://www.cdc.gov/niosh/ltas/rates.html>.

COPD, chronic obstructive pulmonary disease; LRT, likelihood ratio test; MN, malignancy; NHL, non-Hodgkin lymphoma; Obs, observed; SMR, standardized mortality ratio.