Exhibit 321

Updated Mortality Study of Workers at a Petroleum Refinery in Torrance, California, 1959 to 1997

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The updated cohort consisted of 3328 workers who were employed at the Mobil (now ExxonMobil) Torrance, California, refinery for at least 1 year between 1959 and 1997. The vital status of the cohort was determined through a variety of sources, including company employment or retirement records, the Social Security Administration's Death Master File, and the National Death Index. The updated study covered an observation period of 38 years from 1960 to 1997, with a total of 60,612 person-years of observation. A total of 705 (21.2%) cohort members were identified as having died. Mortality data were analyzed in terms of cause-specific standardized mortality ratios (SMRs) and 95% confidence intervals (95% CIs), with expected deaths based on US national cause-, gender-, race-, year-, and age-specific mortality rates. The overall mortality of the cohort was significantly lower than expected when compared with the US general population (SMR, 81.9; 95% CI, 76.0 to 88.2). Overall cancer mortality was also lower than expected (SMR, 79.8; 95% CI, 67.9 to 93.1). For specific cancer sites, significant mortality deficits were observed for cancer of the digestive system (SMR, 70.9; 95% CI, 49.4 to 98.6) and cancer of the respiratory system (SMR, 74.1; 95% CI, 55.5 to 97.0). No significant increase was reported for any site-specific cancer. For nonmalignant diseases, no significant increase was observed for any cause. In particular, significant mortality deficits were reported for ischemic heart disease (SMR, 87.7; 95% CI, 77.2 to 99.3), chronic endocardial disease and other myocardial insufficiencies (SMR, 8.3; 95% CI, 0.2 to 46.0), all other heart disease (SMR, 64.2; 95 % CI, 43.0 to 92.2), and influenza and pneumonia (SMR, 59.2; 95 % CI, 33.1 to 97.6). Detailed analysis by length of employment did not reveal any significant mortality excess or upward trend. Analyses of male employees by job classification (process and maintenance) were conducted. Among maintenance workers, mortality from cirrhosis of the liver (SMR, 190.1; 95% CI, 101.2 to 325.1) and suicide (SMR, 208.6; 95% CI, 111.1 to 356.7) were significantly elevated. However, these mortality excesses did not seem to be related to employment at the refinery. No other causes of death showed significant increase among maintenance workers. A similar separate analysis was conducted for process workers, and no significant excess was detected for any cause. The findings from the present study are discussed in conjunction with results from previous investigations of employees at the Torrance refinery and with results from other refinery studies. Potential limitations of the study are also discussed. (J Occup Environ Med. 2001;43:1089-1102)

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efinery workers are potentially exposed to a wide range of petroleumderived hydrocarbons and chemical substances used in the manufacturing of petroleum fuels and lubricants. According to the International Agency for Research on Cancer, potential exposures at refineries include aromatic amines (eg, anisidines), arsenic compounds, asbestos, benzene, bitumens, butadiene, carbazole, chlorinated hydrocarbons, chromium and chromium compounds, ethylene dibromide, ethylene dichloride, hydrazine, lead and lead compounds, mineral oils, nickel and nickel compounds, phenylenediamine, polycyclic aromatic compounds, and

The Mobil (now ExxonMobil) Torrance, California, refinery first started operations under General Petroleum Corporation of California in 1929. In 1959, the Torrance refinery became part of Mobil Oil Co. The refinery processes approximately 160,000 barrels of oil each day and produces a variety of petroleum products including gasoline, diesel fuels, jet fuel, liquefied petroleum gases, petroleum coke, and sulfur.

In 1985, Enterline and Henderson completed a retrospective cohort mortality study of white male employees at the Torrance refinery who worked for at least 1 year between January 1, 1959, and December 31, 1978. The vital status of these workers was ascertained through December 31, 1978. The cohort study was subsequently updated and expanded to include all employees at the refin-

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ery who worked for at least a year between January 1, 1959, and December 31, 1987, with vital status follow-up extended to December 31, 1987. However, only analyses for white male employees were presented.

The objective of the present investigation was to update and expand the previous cohort study and to continue monitoring the mortality patterns of the Torrance refinery workers. In the present update, enrollment in the study was extended to cover an additional 10 years of employment eligibility (1988 to 1997). Vital status follow-up was extended from December 31, 1987, to December 31, 1997.

Subjects and Methods

The expanded cohort consisted of all Mobil employees who had worked for at least a year at the Torrance refinery between January 1, 1959, and December 31, 1997. The cohort members were identified through a combination of personnel records located on-site and computerized payroll files. Information abstracted from these records included social security number, name, gender, race, date of birth, date of employment, employment status on the closing date of the study, vital status on the closing date of the study, and date of retirement, separation, or death when applicable. Employment histories of individuals in the previous cohort were updated from 1987 to 1997. For each employee, the employment history data consisted of the beginning and ending dates for each job, job titles, and departments.

The vital status of cohort members as of December 31, 1997, was ascertained through several sources, including company employment or retirement records; records at Pension Benefits Information, Inc; the Social Security Administration's Death Master File; and the National Center for Health Statistics' National Death Index (NDI). The Death Master File is a national database of all deaths reported to the Social Security Ad-

ministration since 1939. Vital status of an individual is ascertained by matching the last name and social security number. The NDI, established in 1979, is a national death registry designed to facilitate health investigations. Matching is based on the full name, social security number, birth date, gender, race, and in some cases, father's surname. Vital status information and causes of death of study subjects are provided by NDI through a service known as "NDI Plus." In the present update, causes of death for known decedents were obtained from either NDI Plus or death certificates. The underlying and contributory causes of death were coded or converted to the Eighth Revision of the International Classification of Diseases (ICD-8).

Statistical analyses were based on cause-specific standardized mortality ratios (SMRs). Person-years of observation were classified by age (5year groups), gender, race, and calendar year (5-year groups). The race of approximately 5% cohort members was unknown, and these workers were assumed to be white in the analyses. Expected deaths were calculated by applying the US national age-, cause-, gender-, race-, and year-specific death rates to the corresponding person-years in the cohort. SMRs were computed by expressing the observed deaths as percentages of the expected. The actual calculation was performed through the University of Pittsburgh's OCMAP software program, with standard mortality rates derived from the Mortality and Population Data System.4 Analyses were performed for the entire cohort and for sub-cohorts stratified by gender, length of employment, and time since first employment at the refinery. Trend analyses by length of employment were based on the method described by Breslow and Day. 5 To investigate mortality by job category, male hourly employees at the Torrance refinery were classified into two major job categories: (1) process workers, and (2) maintenance craft workers. This latter subcohort of maintenance craft workers consisted of individuals involved in a broad range of labor and maintenance activities in the mechanical and service divisions of the refinery. Examples of maintenance jobs included pipefitter, auto mechanic, welder, insulator, and painter.

In addition to analyses conducted using the routine OCMAP program, a special analysis for lymphohemopoietic malignancies was also performed. In the United States, because of the way in which mortality rates by ICD-8 categories are tabulated by the National Center for Health Statistics, analyses of non-Hodgkin's lymphoma (NHL) and multiple myeloma (MM) in most occupational cohort studies are generally not reported as such. In the OCMAP program, NHL (ICD-8 200 and 202) appears in two categories: "lymphosarcoma and reticulosarcoma" (ICD-8 200) and "cancer of all other lymphatic and hematopoietic tissue" (ICD-8 202, 203, 208, 209). Thus, part of NHL (ICD-8 202, "other lymphomas") is reported together with MM (ICD-8 203), polycythemia vera (ICD-8 208) and myelofibrosis (ICD-8 209). In the special analysis, NHL and MM were analyzed separately. In the OCMAP program, different types of leukemia were analyzed as a single group. Therefore, the following major types of leukemia were also analyzed separately: acute lymphatic (ICD-8 204.0), chronic lymphatic (ICD-8 204.1), acute myeloid (ICD-8 205.0), and chronic myeloid (ICD-8 205.1) leukemia. US mortality rates for celltype specific leukemias, NHL and MM compiled by the National Cancer Institute based on data derived from the "Surveillance, Epidemiology, and End Results" (SEER) program were used in computing the expected deaths from these causes. 6,7

In addition, because of the use of asbestos at the refinery in the past, separate analyses for asbestos-related diseases were also performed. The OCMAP program did not pro-

vide a separate analysis for asbestosis. The term asbestosis refers to pulmonary fibrosis caused by exposure to asbestos, or pulmonary fibrosis in persons with a documented asbestos-exposure history. Because certifying physicians might not be aware of the decedents' asbestos exposure (if any), such deaths could have been coded simply as pulmonary fibrosis or pneumoconiosis. Therefore, mortality from "pulmonary fibrosis" or "pneumoconiosis" (ICD-8 515 to 517) was analyzed. Expected deaths were based on US rates for pulmonary fibrosis obtained from the National Center for Health Statistics (unpublished data by special request). For malignant peritoneal or pleural mesothelioma, mortality rates compiled by the National Cancer Institute were used. 8,9

Results

A total of 3328 workers were eligible for inclusion in the updated study (Table 1). Eighty-five percent (85.9%) of the cohort members were white (including Hispanics), and 89.1% were male. Selected employment and demographic characteristics of the cohort are provided in Table 1. Approximately 80% were first employed in their twenties or thirties, and only 10.3% were hired at age 40 or older. Most (81.3%) of the cohort members were hired in or after 1950, and a majority (57.8%) were employed for at least 10 years at the refinery.

The maximum length of vital status follow-up was 38 years (January 1, 1960, to December 31, 1997), and the average duration of follow-up was 18 years. A total of 1356 (40.7%) cohort members had a follow-up of 20 years or more. The increase in person-years from the previous study (white male workers only) was 27,281 or 81.8% to a total of 60,612 (all cohort members). By the end of the follow-up period (December 31, 1997), 1062 (31.9%) cohort members were still employed at the refinery, 1561 (46.9%) had separated or retired, and 705 (21.2%)

TABLE 1
Characteristics of the Torrance, California, Cohort

	n	% of Total
Total employees	3,328	100.0
Total person-years	60,612	100.0
Race		
White	2,859	85.9
Nonwhite	469	14.1
Gender		
Male	2,965	89.1
Female	363	10.9
Year of hire		
<1950	623	18.7
≥1950	2,705	81.3
Duration of employment (yrs)		
<10	1,403	42.2
10-29	1,472	44.2
30+	453	13.6
Age at hire		
<20	317	9.5
20-39	2,669	80.2
40+	342	10.3
Age at death		
<50	64	9.1
50-59	91	12.9
60-69	186	26.4
70+	364	51.6
Vital status (December 31, 1997)		
Alive	2,585	77.7
Dead	705	21.2
Unknown	38	1.1

were known to have died. Information on vital status was 98.9% complete, with 38 (1.1%) separated employees lost to follow-up. Of the 705 cohort members identified to have died between 1959 and 1997, death certificate information (derived from death certificates themselves or from NDI Plus listings) was available for 695 deaths (98.6%). The causes of death of the remaining 10 deaths (1.3%) were unknown, and they were included in the overall SMR (all causes) calculations but not in cause-specific SMR calculations.

The number of observed deaths, SMRs, and 95% confidence limits (95% CI) for selected causes for all workers are presented in Table 2. The total number of observed deaths was 705, compared with 860.43 expected (SMR, 81.9; 95% CI, 76.0 to 88.2), indicating a statistically significant deficit in overall mortality. Mortality from all cancers was also significantly lower than expected.

The number of observed cancer deaths was 161, versus 201.79 expected (SMR, 79.8; 95% CI, 67.9 to 93.1).

For specific cancer sites, significant mortality deficits were observed for both cancer of the digestive system (35 observed vs 49.38 expected deaths; SMR, 70.9; 95% CI, 49.4 to 98.6) and cancer of the respiratory system (53 observed vs 71.50 expected deaths; SMR, 74.1; 95% CI, 55.5 to 97.0). No significant increase was reported for any site-specific cancer.

For nonmalignant diseases, the majority of SMRs were below 100, and no significant increase was observed for any cause. In particular, significant mortality deficits were reported for ischemic heart disease (SMR, 87.7; 95% CI, 77.2 to 99.3), all other heart disease (SMR, 64.2; 95% CI, 43.0 to 92.2), and influenza and pneumonia (SMR, 59.2; 95% CI, 33.1 to 97.6).

TABLE 2
Cause-Specific Mortality of the Entire Cohort of Employees at the Torrance Refinery^a

95% Confidence Limits OBS Cause of Death EXP SMR Lower Upper All causes of death 705 860.43 81.9** 76.0 88.2 All malignant neoplasms 161 201.79 79.8** 67.9 93.1 Cancer of buccal cavity and pharynx 3 4.97 60.4 12.5 176.4 Cancer of digestive organs and peritoneum 35 49.38 70.9* 49.4 98.6 Cancer of esophagus 3 5.05 59.4 12.3 173.6 Cancer of stomach 5 7.24 69.0 22.4 161.1 Cancer of large intestine 12 18.20 65.9 34.1 115.2 Cancer of rectum 2 4.23 47.3 5.7 170.8 Cancer of biliary passages and liver 1 4.28 9 130.3 23.4 0.6 Cancer of pancreas 12 9.99 120.1 62.1 209.8 Cancer of all other digestive organs 0 1.39 0.0 0.0 265.2 Cancer of respiratory system 53 71.50 74.1* 55.5 97.0 0 2.56 Cancer of larynx 0.0 0.0 144.4 Cancer of bronchus, trachea, lung 52 68.20 76.2* 56.9 100.0 Cancer of all other respiratory 1 0.73 137.4 3.4 765.7 3 Cancer of breast 1.56 192.4 39.7 562.1 0 0.34 All uterine cancers 0.0 0.0 1,073.6 Cancer of cervix uteri 0 0.22 0.0 0.0 1,693.6 Cancer of other female genital organs 0 0.35 0.0 0.0 1,050.0 18 18.98 Cancer of prostate 94.8 56.2 149.9 0 0.54 Cancer of testes and other male genital organs 0.0 0.0 678.7 3 Cancer of kidney 4.79 62.7 12.9 183.2 2 Cancer of bladder and other urinary organs 5.73 . 34.9 4.2 126.0 Malignant melanoma of skin 4 2.73 146.3 39.9 374.7 1 Cancer of eye 0.13 799.6 20.0 4,455.3 Cancer of central nervous system 6 4.91 122.1 44.8 265.8 0 0.55 Cancer of thyroid and other endocrine glands 0.0 0.0 665.9 0 Cancer of bone 0.54 0.0 0.0 689.2 Cancer of all lymphatic, hemopoietic tissue : 19 18.65 101.9 61.3 159.1 Lymphosarcoma and reticulosarcoma 1 2.11 47.4 1.2 264.3 Hodgkins disease 1 1.13 88.4 2.2 492.8 Leukemia & aleukemia 5 7.37 67.8 22.0 158.3 Cancer of all other lymphopoietic tissue 12 8.04 149.2 77.1 260.6 14 15.28 All other malignant neoplasms 91.6 50.1 153.7 0 2.10 Benign neoplasms 0.0 0.0 175.3 Diabetes mellitus 10 14.30 69.9 33.5 128.6 Cerebrovascular disease 40 50.78 78.8 56.3 107.3 All heart disease 289 326.64 88.5* 78.6 99.3 Rheumatic heart disease 5 4.64 107.7 35.0 251.4 249 283.86 Ischemic heart disease 87.7* 77.2 99.3 Chronic endocardial disease; other myocardial insufficiency 1 12.12 8.3** 0.2 46.0 5 6.63 Hypertension with heart disease 75.4 24.5 176.1 All other heart disease 29 45.16 64.2* 43.0 92.2 3 2.30 130.4 Hypertension w/o heart disease 26.9 381.0 Nonmalignant respiratory disease 55 69.56 79.1 59.6 102.9 15 25.35 Influenza & pneumonia 59 2" 33.1 97.6 Bronchitis, emphysema, asthma 12 14.23 84.3 43.6 147.3 **Bronchitis** 3 2.28 131.5 27.1 384.3 9 82.2 10.95 Emphysema 37.6 156.0 Asthma 0 1.02 0.0 0.0 361.7 Other nonmalignant respiratory disease 28 31.37 89.3 59.3 129.0 0 3.35 Ulcer of stomach & duodenum 0.0 0.0 110.2 Cirrhosis of liver 19 17.25 110.2 66.3 172.0 Nephritis and nephrosis 5 4.33 115.5 37.5 269.5 All external causes of death 59 65.15 90.6 68.9 116.8 accidents 37 38.97 94.9 66 8 130.9 Motor vehicle accidents 22 17.65 124.6 78.1 188.7 All other accidents 15 21.47 69.9 39.1 115.2 Suicides 17 15.73 108.1 63.0 173.1 Homicides and other external causes 5 9.65 51.8 120.9

Number of employees = 3,328; person-years = 60,612. OBS, observed; EXP, expected; SMR, standardized mortality ratio.

^{*} Significant at 5% level.

^{**} Significant at 1% level.

A separate analysis for male employees is presented in Table 3. Because male workers represented approximately 90% of the entire cohort, their mortality experience was quite similar to that of the entire cohort. In particular, no significant increase in mortality from any cause of death was detected in male employees (Table 3). Similar to the total cohort, significant mortality deficits were observed for the following causes: all causes, all cancers, cancer of the digestive system, cancer of the respiratory system, ischemic heart disease, all other heart disease, and influenza and pneumonia.

Among the 363 female employees, only 11 deaths were reported. There were two deaths from breast cancer, one from lung cancer, and one from cancer of the large intestine. For nonmalignant diseases, there were two deaths from cerebrovascular disease, two from ischemic heart disease, one from generalized arteriosclerosis, and one from suicide. The cause of death of the remaining death was unknown. Given the small numbers of reported deaths, we will not present cause-specific SMRs among female employees.

Table 4 shows cause-specific mortality of male employees by length of employment at the refinery. None of the causes of death showed a statistically significant upward trend. Several causes seemed to show an upward trend, but none was statistically significant: all causes (χ^2_{trend} = 1.86, P > 0.05); bronchitis, emphysema, and asthma ($\chi^2_{trend} = 2.74$, P > 0.05); motor vehicle accidents $(\chi^2_{\text{trend}} = 0.77, P > 0.05);$ and suicides ($\chi^2_{\text{trend}} = 2.25, P > 0.05$). None of the SMRs presented in Table 4 was significantly elevated, regardless of length of employment.

Long latent periods (decades) are usually required for chronic diseases to develop. In many instances, it would be more appropriate to examine mortality experience only after a certain lag period has elapsed. Table 5 shows mortality analysis by interval since hire (latency) for male employees. The category "cancer of other lymphopoietic tissue" showed a significantly elevated SMR in the shortest latency group (SMR, 458.6; 95% CI, 148.9 to 1070.1, five observed deaths). This residual category includes non-Hodgkin's lymphoma, multiple myeloma, polycythemia vera, and myelofibrosis. As stated in the previous section, a separate analysis of lymphatic and hemopoietic cancers based on a biologically meaningful classification was conducted, and the results are presented below. Table 5 also shows that male employees with 20 to 39 years of latency experienced a significantly elevated SMR of 195.5 (95% CI, 115.9 to 309.0) for cirrhosis of the liver. On the other hand, several causes of death showed statistically significant mortality deficits by time since first employment, including all causes (<20, 20 to 39 years), all cancers (20 to 39 years), ischemic heart disease (<20 years), and cirrhosis of the liver (<20 years).

Workers were classified into broad categories of jobs (at least 6 months in the category). Separate mortality analyses were conducted for maintenance and process workers. Some workers were in both categories. Table 6 shows cause-specific mortality of male maintenance employees. Two causes of death showed significant increases: cirrhosis of the liver (SMR, 190.1; 95% CI, 101.2 to 325.1) and suicide (SMR, 208.6; 95% CI, 111.1 to 356.7). For cirrhosis of the liver, the increase seemed to concentrate in workers with 10 to 29 years of employment in maintenance jobs (SMR, 249.5; 95% CI, 107.7 to 491.6, 8 deaths); and for suicide, among those with 30 years or longer in maintenance jobs (SMR, 1130.5; 95% CI, 136.8 to 4084.1, 2 deaths) (not shown in Table 6). No other SMRs were significantly elevated, and no upward trend was reported for any cause of death among male maintenance workers.

A similar analysis was performed for male process employees (Table 7). The overall mortality was significantly less than expected (SMR, 81.3; 95% CI, 70.9 to 92.8). Lung cancer also showed a significant deficit (SMR, 46.7; 95% CI, 22.4 to 85.8). Similar to maintenance workers, process workers also showed an elevated mortality from cirrhosis of the liver, but the excess was not statistically significant (SMR, 140.1; 95% CI, 60.5 to 276.1).

The specific categories of lymphatic and hemopoietic cancers used in the OCMAP program are based on the statistical classifications compiled by National Center for Health Statistics and do not permit a specific analysis for major cell-type specific leukemias, NHL, or MM, which are more appropriate classifications from the biological point of view. 10-13 Therefore, analyses for these specific subcategories of lymphatic and hemopoietic cancers among male employees were performed separately.

Table 8 shows that there was only one observed acute myeloid leukemia death among male employees, compared with 2.20 expected (SMR, 45.4; 95% CI, 1.1 to 252.8). For chronic myeloid leukemia, there were two observed deaths, approximately one more than the 1.02 expected (SMR, 195.6; 95% CI, 23.7 to 706.6). No death from acute lymphatic leukemia was reported. For chronic lymphatic leukemia, one death was observed, comparable with the 1.34 expected. There were 4 MM deaths, similar to the expected 3.28 (SMR, 122.1; 95% CI, 33.3 to 312.5). Similarly, there were 9 NHL deaths, comparable with the 7.42 expected (SMR, 121.4; 95% CI, 55.5 to 230.4). Table 8 also presents analysis by job category. No significant mortality increase in any disease subcategory was reported by job category.

Table 9 shows the mortality from lymphohemopoietic cancers of male employees by length of employment. There was no upward trend for any of the specific categories of lymphohemopoietic cancer. No significant increase was found for any length of employment. Analysis by time since first employment is also presented in

TABLE 3
Cause-Specific Mortality of Male Employees at the Torrance Refinery^a

95% Confidence Limits Cause of Death OBS EXP SMR Lower Upper 82.3** All causes of death 694 843.36 76.3 88.6 All malignant neoplasms 157 196.22 80.0** 68.0 93.6 4.90 Cancer of buccal cavity and pharynx 3 61.2 12.6 178.9 Cancer of digestive organs and peritoneum 34 48.34 70.3* 48.7 98.3 Cancer of esophagus 3 4.99 60.1 12.4 175.5 7.13 5 70.1 22.8 Cancer of stomach 163.7 Cancer of large intestine 11 17.76 61.9 30.9 110.8 2 4.15 48.2 5.8 Cancer of rectum 174.0 Cancer of biliary passages and liver 1 4.16 24.0 0.6 133.8 Cancer of pancreas 12 9.76 122.9 63.5 214.8 0 1.36 0.0 0.0 Cancer of all other digestive organs 271.2 52 70.36 73.9* Cancer of respiratory system 55.2 96.9 0 2.53 0.0 0.0 145.6 Cancer of larvnx 67.10 76.0* Cancer of bronchus, trachea, lung 51 56.6 99.9 Cancer of all other respiratory 1 0.71 139.9 3.5 779.5 0.24 418.7 Cancer of breast 4 10.5 2.332.9 Cancer of prostate 18 18.98 94.8 56.2 149.9 0 0.54 0.0 Cancer of testes and other male genital organs 0.0 678.7 3 4.70 63.8 13.2 186.5 Cancer of kidney 2 5.69 35.2 Cancer of bladder and other urinary organs 4.3 127.1 4 2.66 Malignant melanoma of skin 150.4 41.0 385.2 Cancer of eye 1 0.12 818.6 20.5 4,561.3 Cancer of central nervous system 6 4.76 126.0 46.2 274.2 0 0.53 Cancer of thyroid and other endocrine glands 0.0 0.0 693.1 0 0.52 0.0 0.0 706.1 Cancer of bone 19 18.18 104.5 62.9 Cancer of all lymphatic, hemopoietic tissue 163.2 Lymphosarcoma and reticulosarcoma 1 2.07 48.2 1.2 268.7 1 1.10 90.8 2.3 505.9 Hodgkins disease Leukemia and aleukemia 5 7.19 69.5 22.6 162.2 7.82 Cancer of all other lymphopoietic tissue 12 153.5 79.3 268.2 14 14.87 94.1 51.5 157.9 All other malignant neoplasms 0 2.04 0.0 0.0 180.8 Benign neoplasms 10 13.82 72.4 34.7 Diabetes mellitus 133.1 Cerebrovascular disease 38 49.72 76.4 54.1 104.9 All heart disease 287 322.42 89.0* 79.0 99.9 Rheumatic heart disease 5 4.47 111.7 36.3 260.7 280.81 Ischemic heart disease 247 88.0* 77.3 99.6 8.4** Chronic endocardial disease; other myocardial insufficiency 1 11.87 0.2 46.9 5 6.42 77.9 25.3 Hypertension with heart disease 181.8 All other heart disease 29 44.19 65.6* 44.0 94.3 2.24 Hypertension w/o heart disease 3 133.B 27.6 390.9 68.45 55 80.3 60.5 104.6 Nonmalignant respiratory disease Influenza and pneumonia 15 24.96 60.1* 33.6 99.1 Bronchitis, emphysema, asthma 12 14.01 85.6 44.2 149.6 3 2.25 133.1 27.5 389.0 Bronchitis 9 10.84 83.0 38.0 157.6 Emphysema 0 0.96 0.0 0.0 386.1 30.85 90.8 Other nonmalignant respiratory disease 28 60.3 131.2 Ulcer of stomach & duodenum 3.31 0.0 0.0 111.5 Cirrhosis of liver 19 16.86 112.7 67.8 176.0 5 4.24 117.9 38.3 Nephritis and nephrosis 275.1 58 63.69 All external causes of death 91.1 69.1 117.7 38.13 Accidents 37 97.0 68.3 133.8 193.9 22 17.18 128.1 80.3 Motor vehicle accidents All other accidents 15 21.10 71.1 39.8 117.2 Suicides 16 15.41 103.8 59.3 168.6

5

9.35

53.5

17.4

124.8

Homicides and other external causes

Number of employees = 2,965; person-years = 55,984. OBS, observed; EXP, expected; SMR, standardized mortality ratio.

^{*} Significant at 5% level.

^{**} Significant at 1% level.

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TABLE 4	
Cause-Specific Mortality of Male Employees at the Torra	ance Refinery by Length of Employment ^a

	<10	Years	10-29	Years	30+ Years		
Cause of Death	OBS	SMR	OBS	SMR	OBS	SMR	
All causes	128	72.7**	425	84.2**	141	86.8	
All malignant neoplasms	38	96.5	87	75.6**	32	76.6	
Cancer of buccal cavity and pharynx	0	0.0	3	102.7	0	0.0	
Cancer of digestive organs and peritoneum	8	87.4	19	65.5	7	68.9	
Cancer of esophagus	0	0.0	2	70.1	1	97.	
Cancer of stomach	1	76.8	4	90.5	0	0.	
Cancer of large intestine	3	93.6	5	47.0	3	76.	
Cancer of rectum	0	0.0	0	0.0	2	241.	
Cancer of biliary passages and liver	1	111.2	0	0.0	0	0.	
Cancer of pancreas	3	160.2	8	137.2	1	48.	
Cancer of all other digestive organs	0	0.0	0	0.0	0	0.	
Cancer of respiratory system	13	90.2	29	71.4	10	65.	
Cancer of larynx	0	0.0	0	0.0	0	0.	
Cancer of bronchus, trachea, lung	12	87.3	29	75.0	10	68.	
Cancer of breast	0	0.0	1	694.2	0	0.	
Cancer of prostate	2	83.5	10	83.8	6	129	
Cancer of kidney	1	95.9	2	74.2	0	0	
Cancer of bladder and other urinary organs	1	122.8	1	27.9	0	0	
Malignant melanoma of skin	1	115.6	3	224.1	0	0	
Cancer of central nervous system	2	142.1	3	117.5	1	124	
Cancer of lymphatic and hemopoietic tissues	6	141.9	11	106.3	2	55	
Lymphosarcoma and reticulosarcoma	1	244.4	0	0.0	0	0	
Hodgkin's disease	1	258.1	0	0.0	0	0	
Leukemia	1	62.3	4	96.6	0	0.	
Cancer of other lymphopoietic tissue	3	164.1	7	163.5	2	117	
Benign neoplasms	0	0.0	0	0.0	ō	0	
Diabetes mellitus	3	103.0	5	61.7	2	71.	
Cerebrovascular disease	3	42.9	22	67.5	13	128	
All heart disease	38	70.1*	192	94.3	57	88.	
Ischemic heart disease	25	54.9**	171	95.6	51	90.	
Nonmalignant respiratory disease	8	76.8	34	79.4	13	85.	
Influenza and pneumonia	0	0.0*	12	74.5	3	58.	
Bronchitis, emphysema, asthma	1	51.3	6	65.6	5	171.	
Other nonmalignant respiratory disease	7	141.9	16	86.8	5	66.	
Cirrhosis of liver	1	19.6	16	169.6	2	86.	
All external causes of death	25	82.7	26	93.0	7	127.	
Accidents	15	87.9	19	108.6	3	83.	
Motor vehicle accidents	9	101.6	11	155.9	2	157.	
All other accidents	6	72.6	8	76.2	1	42.	
Suicides	6	86.8	6	86.3	4	258.	
Homicides and other external causes	4	66.7	1	33.3	0	0.	
No. of employees	2,452		1,393		283		
Person-years	28,430		23,073		4,482		

^a OBS, observed; SMR, standardized mortality ratio.

Table 9. No significant increase was detected for any categories of lymphohemopoietic cancers for any latency group.

Because of the use of asbestos at the refinery in the past, mortality from asbestos-related diseases (asbestosis, lung cancer, and mesothelioma) among cohort members was examined. Although the OCMAP program provided an analysis for lung cancer, it provided no separate analysis for asbestosis or malignant mesothelioma of the peritoneum or pleura. Analyses for asbestosis (pulmonary fibrosis) and malignant mesothelioma of the peritoneum or pleura were performed separately.

The term asbestosis refers to pulmonary fibrosis caused by exposure to asbestos, or pulmonary fibrosis in persons with a documented asbestos-exposure history. The ICD code (eighth revision) for asbestosis is 515.2, and is, therefore, part of the broad category "other nonmalignant respiratory disease" (ICD-8 460 to

^{*} Significant at the 5% level.

^{**} Significant at the 1% level.

TABLE 5
Cause-Specific Mortality of Male Employees at the Torrance Refinery by Time Since First Employment

	<20 `	rears	20-39	Years	40+ Years	
Cause of Death	OBS	SMR	OBS	SMR	OBS	SMI
All causes	84	59.5**	322	83.3**	288	91.2
All malignant neoplasms	20	78.4	77	79.1*	60	81.
Cancer of buccal cavity and pharynx	0	0.0	3	108.2	0	0.
Cancer of digestive organs and peritoneum	2	32.3	19	77.7	13	73.
Cancer of esophagus	0	0.0	2	76.1	1	62.
Cancer of stomach	1	93.6	3	81.0	1	42.
Cancer of large intestine	1	53.8	5	58.1	5	68.
Cancer of rectum	0	0.0	1	44.7	1	74.
Cancer of biliary passages and liver	0	0.0	1	52.2	0	0.
Cancer of pancreas	0	0.0	7	137.5	5	145.
Cancer of all other digestive organs	0	0.0	0	0.0	0	0.
Cancer of respiratory system	6	68.9	27	71.8	19	79.
Cancer of larynx	0	0.0	0	0.0	0	0.
Cancer of bronchus, trachea, lung	5	61.0	27	75.4	19	82.
Cancer of breast	0	0.0	1	831.1	0	0.
Cancer of prostate	1	140.4	4	57.7	13	114
Cancer of kidney	2	286.9	1	41.1	0	0
Cancer of bladder and other urinary organs	٥	0.0	i	38.2	1	37
Malignant melanoma of skin	0	0.0	4	327.5	0	0
Cancer of central nervous system	0	0.0	6	245.4	0	0
Cancer of lymphatic and hemopoletic tissues	8	248.4*	4	48.1	7	105
Lymphosarcoma and reticulosarcoma	0	0.0	1	82.8	0	0.
Hodgkin's disease	1	216.1	0	0.0	0	
Leukemia	2	166.3	0	0.0	3	108
Cancer of other lymphopoletic tissue	5	458.6*	3	88.2	4	120
Benign neoplasms	0	0.0	0	0.0	0	0
Diabetes mellitus	1	46.5	3	48.4	6	109
Cerebrovascular disease	2	40.1	18	81.2	18	79
All heart disease	26	65.0*	139	87.8	122	98
Ischemic heart disease	22	65.5*	119	83.8	106	100
Nonmalignant respiratory disease	2	35.2	21	76.9	32	90
Influenza and pneumonia	0	0.0	5	59.3	10	71
Bronchitis, emphysema, asthma	0	0.0	5	67.2	7	136
Other nonmalignant respiratory disease	2	109.1	11	93.5	15	86
Cirrhosis of liver	0	0.0*	18	195.5*	15	40
All external causes of death	25	73.9	22	193.5	11	129
Accidents	17	89.2		105.0	6	101
Motor vehicle accidents	12	119.9	6	110.7	4	228
All other accidents	5	54.9	8	102.8	2	47
Suicides	5	67.5	6	102.6	5	225
Homicides and other external causes	3	43.7	2	93.0	0	0
Number of employees	2,795		1,599		674	
Person-years	31,161		19,640		5,182	

a OBS, observed; SMR, standardized mortality ratio.

519) in the OCMAP analysis. In the Torrance cohort, no death was attributed to asbestosis. However, because certifying physicians might not be aware of the decedents' asbestos exposure (if any), such deaths could have been coded simply as pulmonary fibrosis or pneumoconiosis. Therefore, the category "pulmonary

fibrosis" was analyzed. For pulmonary fibrosis (ICD-8 515 to 517), there were 3 observed deaths among all male cohort members, compared with 4.24 expected. The corresponding SMR for pulmonary fibrosis among male employees was 70.8 (95% CI, 14.6 to 207.0). One of the three male employees who died from

pulmonary fibrosis had worked in both maintenance and process jobs, and the remaining two in administrative or technical jobs.

As stated above, lung cancer results were provided by the routine OCMAP program. The results for the overall cohort are presented in Table 2 (SMR, 76.2; 95% CI, 56.9 to

^{*} Significant at the 5% level.

^{**} Significant at the 1% level.

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TABLE 6 Cause-Specific Mortality of Male Maintenance Employees at the Torrance Refinerya

95% Confidence Limits

9				95% Confi	dence Limits
Cause of Death	OBS	EXP	SMR	Lower	Upper
All causes of death	301	313.04	96.2	85.6	107.7
All malignant neoplasms	67	73.62	91.0	70.5	115.6
Cancer of buccal cavity and pharynx	1	1.89	53.0	1.3	295.2
Cancer of digestive organs and peritoneum	18	18.06	99.6	59.1	157.5
Cancer of esophagus	3	1.93	155.5	32.1	454.4
Cancer of stomach	2	2.68	74.7	9.0	270.0
Cancer of large intestine	6	6.55	91.6	33.6	199.5
Cancer of rectum	1	1.54	64.9	1.6	361.6
Cancer of biliary passages and liver	0	1.57	0.0	0.0	235.7
Cancer of pancreas	6	3.67	163.7	60.1	356.3
Cancer of all other digestive organs	0	0.51	0.0	0.0	725.5
Cancer of respiratory system	22	26.74	82.3	51.6	124.6
Cancer of larynx	0	0.96	0.0	0.0	382.7
Cancer of bronchus, trachea, lung	22	25.50	86.3	54.1	130.6
Cancer of all other respiratory	0	0.27	0.0	0.0	1,347.7
Cancer of breast	0	0.09	0.0	0.0	4112.6
Cancer of prostate	7	6.59	106.3	42.7	219.0
Cancer of testes and other male genital organs	0	0.23	0.0	0.0	1,616.9
Cancer of kidney	1	1.79	55.7	1.4	310.6
Cancer of bladder and other urinary organs	1	2.03	49.3	1.2	274.5
Malignant melanoma of skin	3	1.06	282.9	58.4	826.8
Cancer of eye	1	0.05	2,172.8	54.3	12,107.0
Cancer of central nervous system	2	1.89	106.1	12.8	383.1
Cancer of thyroid and other endocrine glands	0	0.21	0.0	0.0	1,793.7
Cancer of bone	0	0.20	0.0	0.0	1,844.4
Cancer of all lymphatic, hemopoietic tissue	5	6.87	72.8	23.6	169.8
Lymphosarcoma and reticulosarcoma	0	0.79	0.0	0.0	465.8°
Hodgkins disease	0 ;	-	0.0	0.0	819.9
Leukemia and aleukemia	2	2.70	74.2	9.0	268.0
Cancer of all other lymphopoietic tissue	3	2.93	102.2	21.1	298.7
All other malignant neoplasms	6	5.62	106.7	39.2	232.3
Benign neoplasms	0	0.76	0.0	0.0	482.7
Diabetes mellitus	3	5.16	58.2	12.0	170.0
Cerebrovascular disease	15	17.76	84.5	47.3	139.3
All heart disease	125	117.84	106.1	88.3	126.4
Rheumatic heart disease	0	1.70	0.0	0.0	216.5
Ischemic heart disease	105	102.54	102.4	83.8	124.0
Chronic endocardial disease; other myocardial insufficiency	1	4.08	24.5	0.6	136.5
Hypertension with heart disease	3	2.41	124.5	25.7	363.8
All other heart disease	16	16.30	98.2	56.1	159.4
Hypertension w/o heart disease	2	0.82	243.4	29.5	879.3
Nonmalignant respiratory disease	18	24.22	74.3	44.0	117.5
	5	8.66	57.7	18.7	134.8
Influenza and pneumonia	4	5.06	79.1	21.6	202.5
Bronchitis, emphysema, asthma	0	0.80	0.0	0.0	458.6
Bronchitis	4	3.90	102.7	28.0	262.8
Emphysema	0	0.36	0.0	0.0	1,012.7
Asthma	9	10.97	82.1	37.5	155.8
Other nonmalignant respiratory disease	0	1.21	0.0	0.0	303.9
Ulcer of stomach and duodenum		6.84	190.1*	101.2	325.1
Cirrhosis of liver	13		0.0		239.2
Nephritis and nephrosis	0	1.54		0.0	
All external causes of death	34	26.34	129.1	89.4	180.4
Accidents	17	15.67	108.5	63.2	173.7
Motor vehicle accidents	12	7.24	165.7	85.6	289.4
All other accidents	5	8.49	58.9	19.1	137.5
Suicides	13	6.23	208.6*	111.1	356.7
Homicides and other external causes	4	4.10	97.6	26.6	249.9

^a Number of employees = 1,075; person-years = 23,419. OBS, observed; EXP, expected; SMR, standardized mortality ratio.

^{*} Significant at 5% level.

^{**} Significant at 1% level.

TABLE 7
Cause-Specific Mortality of Male Process Employees at the Torrance Refinery^a

95% Confidence Limits Cause of Death OBS **EXP** SMR Lower Upper All causes of death 219 269.45 81.3** 70.9 92.8 42 62.44 67.3** All malignant neoplasms 48.5 90.9 Cancer of buccal cavity and pharynx 0 1.59 0.0 0.0 232.3 13 15.24 85.3 Cancer of digestive organs and peritoneum 45.4 145.8 Cancer of esophagus 1 1.61 62.1 1.6 346.3 Cancer of stomach 3 2.22 135.3 27.9 395 4 Cancer of large intestine 3 5.60 53.6 11.1 156.6 Cancer of rectum 1 1.29 77.5 1.9 432.1 0 1.35 0.0 Cancer of biliary passages and liver 0.0 273.6 Cancer of pancreas 5 3.08 162.3 52.7 378.8 0 Cancer of all other digestive organs 0.42 0.0 0.0 872.0 10 22.48 44.5** 21.3 Cancer of respiratory system 81.8 0 0.81 0.0 Cancer of larynx 0.0 455.3 46.7** Cancer of bronchus, trachea, lung 10 21.43 22.4 85.8 Cancer of all other respiratory 0 0.23 0.0 0.0 1,592.5 0 0.08 Cancer of breast 0.0 0.0 4,883.0 3 5.82 Cancer of prostate 51.5 10.6 150.6 0 0.19 Cancer of testes and other male genital organs 0.0 0.0 1,933.0 Cancer of kidney 2 1.51 132.7 16.1 479.5 Cancer of bladder and other urinary organs 0 1.75 0.0 0.0 211.1 0.89 112.5 2.8 Malignant melanoma of skin 1 627.0 Cancer of eye 0 0.04 0.0 0.0 9,620.5 Cancer of central nervous system 2 1.58 126.4 15.3 456.5 0 0.17 Cancer of thyroid and other endocrine glands 0.0 0.0 2.142.1 0 0.17 0.0 Cancer of bone 0.0 2,166.8 7 5.88 Cancer of all lymphatic, hemopoietic tissue 119.0 47.8 245.1 Lymphosarcoma and reticulosarcoma 0 0.65 0.0 0.0 564 4 0.37 267.8 Hodakins disease 6.7 1,492.1 2 2.32 10.5 Leukemia and aleukemia 86.4 312.1 Cancer of all other lymphopoietic tissue 4 2 54 157.3 42.9 402.7 All other malignant neoplasms 4 4.78 83.6 22.8 214.2 0 0.66 Benign neoplasms 0.0 0.0 562.7 Diabetes mellitus 0 4.41 0.0* 0.0 83.6 Cerebrovascular disease 14 15.03 93.2 50.9 156.3 92 100.22 91.8 74.0 All heart disease 112.6 Rheumatic heart disease 1.42 70.6 1.8 393.5 1 Ischemic heart disease 78 86.86 89.8 71.0 112.1 Chronic endocardial disease; other myocardial insufficiency 0 3.73 0.0* 0.0 98.9 Hypertension with heart disease 0 2.07 0.0 0.0 178.3 All other heart disease 13 14.26 91.2 48.5 155.9 0 0.69 0.0 0.0 532.1 Hypertension w/o heart disease 18 Nonmalignant respiratory disease 21.34 84.4 50.0 133.3 4 7.67 52.1 14.2 133.5 Influenza and pneumonia 7 4.29 163.2 65.6 Bronchitis, emphysema, asthma 336.2 2 0.68 292.7 **Bronchitis** 35.4 1,057.5 5 Emphysema 3.30 151.6 49.2 353.7 0 0.31 0.0 0.0 1,178.8 Asthma Other nonmalignant respiratory disease 7 9.80 71.4 28.7 147.1 0 Ulcer of stomach and duodenum 1.01 0.0 0.0 364.6 8 5.71 140.1 60.5 276.1 Cirrhosis of liver 3 1.33 Nephritis & nephrosis 226.0 46.6 660.4 25 All external causes of death 23.70 105.5 68.3 155.7 Accidents 17 13.89 122.4 71.3 195.9 Motor vehicle accidents 12 6.48 185.3 95.8 323.7 All other accidents 5 7.47 67.0 21.7 156.2 Suicides 6 5.60 107.2 39 4 233.4 Homicides and other external causes 3.98 50.3 6.1 181.6

a Number of employees = 1,069; person-years = 20,815. OBS, observed; EXP, expected; SMR, standardized mortality ratio.

^{*} Significant at 5% level.

[&]quot; Significant at 1% level.

TABLE 8

Mortality from Lymphohemopoietic Cancers of Male Employees at the Torrance Refinery^a

Cause of Death	OBS	EXP	SMR	95% CI
Total cohort				
Acute lymphoid leukemia	0	0.36	0.0	0.0-1,035.2
Chronic lymphoid leukemia	1	1.34	74.4	1.9-414.3
Acute myeloid leukemia	1	2.20	45.4	1.1-252.8
Chronic myeloid leukemia	2	1.02	195.6	23.7-706.6
Multiple myeloma	4	3.28	122.1	33.3-312.5
Non-Hodgkin's lymphoma	9	7.42	121.4	55.5-230.4
Maintenance				
Acute lymphoid leukemia	0	0.14	0.0	0.0-2,721.8
Chronic lymphoid leukemia	0	0.48	0.0	0.0-769.7
Acute myeloid leukemia	0	0.83	0.0	0.0-443.0
Chronic myeloid leukemia	1	0.39	253.6	6.3-1,412.8
Multiple myeloma	0	1.22	0.0	0.0-302.4
Non-Hodgkin's lymphoma	2	2.81	71.2	8.6-257.2
Process				
Acute lymphoid leukemia	0	0.12	0.0	0.0-3,042.5
Chronic lymphoid leukemia	1	0.42	240.6	6.0-1,340.6
Acute myeloid leukemia	1	0.72	139.4	3.5-776.5
Chronic myeloid leukemia	0	0.34	0.0	0.0-1,083.8
Multiple myeloma	1	1.03	96.8	2.4-539.2
Non-Hodgkin's lymphoma	4	2.41	166.1	45.3-425.4

^a OBS, observed; EXP, expected; SMR, standardized mortality ratio; CI, confidence interval.

TABLE 9
Mortality From Lymphohemopoietic Cancers of Male Employees at the Torrance Refinery by Length of Employment and Time Since Hire^a

Cause of Death	OBS	SMR	95% CI	OBS	SMR	95% CI	OBS	SMR	95% CI
Length of employment		<10 Years 10-29 Years		30+ Years					
Years									
Acute lymphoid leukemia	0	0.0	0.0-3,332.3	0	0.0	0.0-1,989.3	0	0.0	0.0-6,125.3
Chronic lymphoid leukemia	0	0.0	0.0-1,869.5	1	120.5	3.0-671.6	0	0.0	0.0-1,160.3
Acute myeloid leukemia	0	0.0	0.0-706.9	1	80.6	2.0-449.4	0	0.0	0.0 834.1
Chronic myeloid leukemia	1	344.2	8.6-1,918.0	1	179.3	4.5-999.3	0	0.0	0.0-2,115.9
Multiple myeloma	1	166.6	4.2-928.1	1	51.1	1.3-284.7	2	277.9	33.6-1,003.9
Non-Hodgkin's lymphoma	4	225.9	61.1-578.5	5	119.4	38.8-278.7	0	0.0	0.0-253.0
Years									
Time since hire		<20 '	Years		20-39	Years		40+	Years
Acute lymphoid leukemia	0	0.0	0.0-3,613.3	0	0.0	0.0-2,892.9	0	0.0	0.0-2,910.3
Chronic lymphoid leukemia	0	0.0	0.0 - 4,546.3	0	0.0	0.0-628.1	1	147.8	3.7-823.8
Acute myeloid leukemia	0	0.0	0.0-958.5	0	0.0	0.0-357.7	1	126.9	3.2-707.2
Chronic myeloid leukemia	2	729.5	93.1-2,780.0	0	0.0	0.0-861.3	0	0.0	0.0-1,103.6
Multiple myeloma	1	271.3	6.8-1,511.6	2	125.3	15.2-452.7	1	76.2	1.9-424.6
Non-Hodgkin's lymphoma	4	298.3	81.3-763.9	2	57.9	7.0-209.1	3	114.5	23.6-334.6

a OBS, observed; SMR, standardized mortality ratio; Cl, confidence interval.

100.0). The lung cancer SMR for all male employees was 76.0 (95% CI, 56.6 to 99.9). Exposure to asbestos has been linked to malignant peritoneal and pleural mesothelioma in epidemiologic studies. Malignant peritoneal mesothelioma is part of "malignant neoplasm of the perito-

neum and retroperitoneal tissue" (ICD-8 158), and malignant pleural mesothelioma is part of "malignant neoplasm of the pleura" (ICD-8 163). The most comprehensive mortality rates for malignant peritoneal and pleural mesothelioma for the United States are those compiled by

the National Cancer Institute in the SEER program. 8.9 In the SEER program, cases were selected having the International Classification of Diseases for Oncology morphology code 905 (mesothelial neoplasms) and topography codes 158 (peritoneum) or 163 (pleura). In addition, only

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malignant cases were included. In the present study, diagnostic information was based entirely on death certificates and no morphologic information was available. In the Torrance cohort, there was one death coded as ICD-8 163 (malignant neoplasm of the pleura). Based on the SEER rates, approximately 1.08 deaths from malignant mesothelioma of the pleura were expected. The corresponding SMR was 92.6 (95% CI, 2.4 to 516.0).

Discussion

First, we will compare some major results in the updated study with those previously reported for the Torrance cohort.³ We will then discuss several specific findings in the updated study that are of particular interest in epidemiologic studies of refinery workers.

Extended enrollment and follow-up in this update on Torrance refinery workers added 1337 workers (+67.2%), 27,281 person-years (+81.8%), and 297 additional deaths (+72.8%). One of the reasons for the sizable increases was that only white men were included in the previous study, whereas no restriction on gender or race was imposed on cohort eligibility in the update. In the previous analysis, 20.5% of cohort members were deceased, compared with 21.2% in this update. Updated results for the total cohort were consistent with earlier reports, which demonstrated a favorable overall mortality experience of the Torrance refinery workers. According to national rates, the updated SMR of 81.9 for all causes showed a significant deficit of 18.1%, which was similar to the deficit previously observed (SMR, 81). As in the previous study, the favorable overall mortality experience of these refinery workers might be attributed to the "healthy worker effect," whereby working populations exhibit decreased overall mortality because of initial selection into the workplace and maintenance of this healthier status through benefits derived from employment.

In terms of specific causes, for the cohort as a whole or the male subcohort, no significant mortality increase was observed for any cause of death. The earlier reports showed nonsignificant deficits for cancer of the digestive system and cancer of the respiratory system. In the updated study, similar deficits were observed, and because of the larger numbers of deaths, the deficits for both cancer of the digestive system and cancer of the respiratory system were statistically significant. In the previous reports, a significant deficit of ischemic heart disease was reported. In the current update, a similar significant deficit was observed for ischemic heart disease. In addition, a significant deficit was also reported for the category "all other heart disease."

Analysis by length of employment also indicated that there was no work-related mortality pattern for male employees. No significant increase of mortality was found for any cause for any length of employment group, and no upward trend was detected for any cause of death. Analysis by latency (time since hire) showed that there was a significant mortality excess from the miscellaneous category "cancer of other lymphopoietic tissue" for the group with less than 20 years of latency (5 observed vs 1.09 expected; SMR, 458.6). Of the five deaths in this category, one was attributed to multiple myeloma (ICD-8 203), and the other four involved "other lymphomas" (ICD-8 202). As discussed earlier, a proper analysis of NHL should include "lymphosarcoma and reticulosarcoma" (ICD-8 200) and "other lymphomas" (ICD-8 202). In the group with less than 20 years of latency, no death was observed for "lymphosarcoma and reticulosarcoma" (ICD-8 200). Thus, according to the disease grouping provided in the OCMAP analysis, NHL was reported in two places: no death was attributed to "lymphosarcoma and reticulosarcoma" (ICD-8 200) and four deaths involved "other lymphomas" (ICD-8 202). When NHL (ICD-8 200 and 202) was analyzed as a group (Table 9), the SMR was 298.3 (95% CI, 81.3 to 763.9) and not significant. The more important observation was that there was no upward trend by length of employment for NHL.

The finding of no association between NHL and employment at the Torrance refinery is consistent with the conclusion reached in a recent review of NHL in petroleum workers. 14 The review was based on more than 308,000 petroleum workers from the United States, the United Kingdom, Canada, Australia, Italy, and Finland. None of the individual studies showed a significant increase of NHL. For all refinery studies combined, there were 408 observed NHL deaths, comparable with the 412.20 expected (SMR, 99; 95% CI, 90 to 109). Specifically for US refinery studies, there were 317 observed deaths, slightly fewer than the 330.79 expected (SMR, 96; 95% CI, 86 to 107). There is no evidence that employment at petroleum refineries is associated with an increased risk of NHL.

For male employees with 20 to 39 years of latency, mortality from cirrhosis of the liver was significantly elevated, but it was significantly low for male employees with fewer than 20 years of latency (Table 5). There was no upward trend of mortality from cirrhosis of the liver by length of employment (Table 4). Cirrhosis of the liver is associated with alcohol consumption. Unfortunately, no information on alcohol consumption was available in the study. Cirrhosis of the liver has not been linked to any petrochemicals present at refineries, and no other epidemiologic studies of petroleum refinery workers have reported an increased mortality from cirrhosis of the liver.

Because of potential differences in exposures and exposure patterns, separate analyses were performed for maintenance and process workers at the refinery. Among maintenance workers, mortality from cirrhosis of the liver was significantly elevated (13 observed vs 6.84 expected; SMR, 190.1; 95% CI, 101.2 to 325.1). As discussed above, the excess of cirrhosis of the liver was not likely to be related to employment at the refinery. There was a significant increase of mortality from suicide among maintenance workers (13 observed vs 6.23 expected; SMR, 208.6; 95% CI, 111.1 to 356.7). However, there is no reason to believe that the increase was related to exposures at the refinery. Among process workers, no significant increase of mortality from any cause of death was detected.

As stated above, the OCMAP program does not provide analyses specific to subgroups of lymphohemopoietic cancers. When analyses were performed separately for acute lymphatic leukemia, chronic lymphatic leukemia, acute myeloid leukemia, chronic myeloid leukemia, MM, and NHL, no significant increase was found for any category for the cohort as a whole or for the sub-cohorts of maintenance or process workers. However, it must be noted that the number of deaths for each category was small, with the possible exception of NHL (9 observed deaths vs 7.42 expected). Nevertheless, the data indicated that there was no increased mortality from any of the subcategories of lymphohemopoietic

Similar to other industrial facilities, asbestos was used at the Torrance refinery in the past, and detailed analyses of asbestos-related diseases were performed for male employees. Although specific exposure measurements were not available, data from the industry as a whole indicated that asbestos concentrations at refineries were generally extremely low. I Among Torrance refinery employees, no death was attributed to "asbestosis" as the underlying cause of death. The term asbestosis refers to pulmonary fibrosis in persons with documented asbestos exposure history. Because certifying physicians might not be

aware of the decedents' asbestos exposure, such deaths could have been coded simply as pulmonary fibrosis or pneumoconiosis. In terms of pulmonary fibrosis, 3 deaths were reported, compared with 4.24 expected (SMR, 70.8; 95% CI, 14.6 to 207.0); one of these decedents had worked in both process and maintenance jobs, and the other two in administrative or technical jobs.

One death in the cohort was coded as ICD-8 163 (malignant neoplasm of the pleura), whereas 1.08 deaths were expected. This individual worked for approximately 2 years in the late 1950s at the Torrance refinery and died 4 years after termination of employment. Epidemiologic studies have reported the average latency of malignant pleural mesothelioma to be around 30 to 40 years, even among heavily exposed workers. The extremely short interval of 6 years between employment at the Torrance refinery in 1958 and death in 1964 does not support an association between the two. In addition to the malignant pleural cancer case, the words "malignant mesothelioma" were mentioned on another death certificate. The cause of death listed on the death certificate was "malignant mesothelioma, metastatic" (with no specific site) and was coded as ICD-8 199.1. According to the SEER criteria, the information provided on the death certificate was not sufficient to classify this death specifically as either malignant mesothelioma of the peritoneum or malignant mesothelioma of the pleura.

For lung cancer, an asbestosrelated malignancy, no increase was found for the overall cohort or the sub-cohorts of maintenance or process workers. In fact, for all three groups, lung cancer mortality was less than expected. Equally important is the observation that no upward trend by length of employment was detected. Thus, the data indicated no relationship between lung cancer mortality and employment at the Torrance refinery. This finding is consistent with the conclusion reached in a recent review of the data of more than 350,000 petroleum workers from the United States, the United Kingdom, Canada, Australia, Italy, and Finland.15 Lung cancer SMRs from individual studies ranged from 44 to 107. For all studies combined, there were 5695 observed lung cancer deaths, significantly fewer than the 7056.25 expected (SMR, 81; 95% CI, 79 to 90). For US refinery workers, there were 2696 observed lung cancer deaths, significantly fewer than the 3573.27 expected (SMR, 75; 95% CI, 73 to 78). Thus, there was no indication that petroleum workers were at an increased risk of lung cancer.

It should be pointed out that there are some potential limitations in this study. Most limitations are typical of a historical mortality study of industrial populations. First, although both the percentage of individuals with unknown vital status (1.1%) and the proportion of deaths with no death certificate information (1.3%) were low, it was possible, but unlikely, that some deaths from causes of interest might have been missed. We do not believe that such low percentages would have any significant impact on our results.

Second, the health endpoint in our study was mortality, which is a reasonable surrogate for incidence of many cancers. Analyses were based on the underlying cause of death listed on death certificates. As such, the investigation inherited the problems associated with death certificates (diagnostic accuracy, for example). No detailed clinical information was available on the deaths in our study. However, it must be pointed out that although detailed information derived from medical records or pathology reports may be more accurate than that based on death certificates, it would be inappropriate to use such information for comparisons with national mortality rates. In our study, we compared diagnoses based on death certificates with national statistics that were derived

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from death certificates as well. Furthermore, our analysis was adjusted for calendar time, thus at least partially controlling for changes in survival and diagnostic practices.

As in most historical cohort mortality studies, little quantitative exposure data were available during the early part of the study, and analyses by quantitative exposure indices were not possible. In the present study, we relied on duration of employment and the broad classification of jobs (process and maintenance). Although such exposure surrogates could be informative, they might lack both sensitivity and specificity.

For some analyses in the overall cohort, and in some sub-cohort analyses (particularly those stratified by job category or by length of employment), the numbers of deaths from some rare causes were small. As a result, the corresponding 95% CIs were wide, and the findings must be interpreted with caution. However, for common disease categories such as all cancers, digestive cancer, lung cancer, prostate cancer, heart disease, and nonmalignant respiratory disease, the statistical power of the study was quite adequate.

Finally, information on lifestyle or exposures from employment elsewhere was not available, although it would have been extremely helpful in interpreting some of the findings in the present study, such as alcohol consumption in relation to cirrhosis of the liver. From a methodologic point of view, the most appropriate approach toward incorporating lifestyle risk factors and exposures from employment elsewhere is the nested case-control study design.

Conclusion

This updated study confirms earlier findings of an overall favorable mortality experience of employees at the Torrance, California, refinery when compared with the US general population. In addition, significant mortality deficits were found for several causes of death. For the entire cohort, no significant increase was detected in any cause-specific mortality that could be attributed to employment at the refinery. Furthermore, analyses by job category and length of employment indicated no mortality increase from any cause of death associated with employment at the refinery.

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