Association between Post-Diagnosis Particulate Matter Exposure Among 5-Year Cancer Survivors and Cardiovascular Disease Risk in Three Metropolitan Areas from South Korea

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Abstract: Cancer survivors are at an increased risk for cardiovascular disease (CVD). However, the association between particulate matter (PM) and CVD risk among cancer survivors (alive >5 years since diagnosis) is unclear. We investigated the risk of CVD among 40,899 cancer survivors within the Korean National Health Insurance Service database. Exposure to PM was determined by assessing yearly average PM levels obtained from the Air Korea database from 2008 to 2011. PMs with sizes <2.5 (PM2.5), <10 (PM10), or 2.5-10 (PM2.5-10) μm in diameter were compared, with each PM level exposure further divided into quintiles. Patients were followed up from January 2012 to date of CVD event, death, or December 2017, whichever came earliest. Adjusted hazard ratios (aHRs) and 95% confidence intervals (CIs) for CVD were calculated using Cox proportional hazards regression by PM exposure levels. Compared with cancer survivors in the lowest quintile of PM2.5 exposure, those within the highest quintile had a greater risk for CVD (aHR 1.31, 95% CI 1.07-1.59). Conversely, increasing PM10 and PM2.5-10 levels were not associated with increased CVD risk (p for trend 0.078 and 0.361, respectively). Cancer survivors who reduce PM2.5 exposure may reduce their risk of developing CVD.

Keywords: cardiovascular disease; particulate matter; cancer survivor; metropolitan area

1. Introduction

With the increasing incidence and the improved survival of patients with cancer, the number of cancer survivors is rapidly increasing in Korea. In 2016, 229,180 Koreans were newly diagnosed with cancer, while 78,194 died from cancer. The recent 5-year relative survival rate for cancer patients in Korea was 70.6% which was similar to that reported in the US of 67.1% [1,2]. Nearly a million people in Korea have survived for more than 5 years after their cancer diagnosis compared to over 11 million in the US [3,4]. Mortality from primary cancers accounts for up to 38.0% of all-cause mortality among cancer patients, while cardiovascular diseases (CVD) accounts for 11.3% [5]. Cardiovascular risk among these patients is as a consequence of the cardiotoxicity of treatment, biological mechanisms
such as inflammation and oxidative stress, and shared lifestyle risk factors for both the cancer and CVD [6]. The increased risk for CVD has been shown among various cancers, including cancers of the breast, lung, prostate, pancreatic, hematologic, esophageal, kidney, and ovary [7,8].

Particulate matter (PM), defined as material suspended in the air in the form of minute solid particles or liquid droplets, is related to both cancers and CVD. It is currently classified as a Group 1 carcinogen by the International Agency for Research on Cancer (IARC) [9,10]. It has a dose-response relationship with carcinogenesis in various cancer types [9-11]. PM is also associated with subclinical atherosclerosis [12] and increased morbidity and mortality from CVD [13,14]. The size of PM has an influence on the pathophysiology of relative diseases and is generally classified into PM10 (<10 µm diameter); which is further subdivided into PM2.5-10 (2.5-10 µm), PM2.5 (<2.5 µm), and ultrafine (<0.1 µm) [15]. PM less than 10 µm can deposit in airways and lungs, while the finest particles can reach the alveoli, enter the bloodstream, and cause systemic inflammation. The harmful effects of PM among people with preexisting diseases such as CVD, respiratory diseases, and diabetes is well known [16] however, there is scarce evidence among cancer survivors.

Exposure to PM affects short-term survival of patients with lung and liver cancers [17,18] as well as long-term survival of patients with breast cancer [19]. Although former investigators speculated that PM affects cancer progression and leads to poor outcomes, cardiovascular morbidity and mortality due to PM has not yet been studied among cancer survivors.

Using data from the Korean National Health Insurance Service (NHIS), we assessed the effects of PM concentration and diameter on the incidence of CVD among patients with 5-year survival following their cancer diagnosis.

2. Experimental Section

2.1. Study population

The Korean NHIS provides mandatory health insurance for all citizens [20] covering various health services including outpatient and inpatient department visits, pharmaceutical prescriptions, surgical procedures, and procedural and laboratory examinations. All citizens aged ≥40 years are eligible for a biannual health screening exam that includes a self-reported questionnaire on their past medical history, lifestyle behaviors, anthropometric measurements including height and weight, blood pressure, and blood and urine laboratory examinations [21]. The NHIS provides some of the data from these services for research purposes and numerous epidemiological studies have used the data previously. Its validity is described in detail elsewhere [20-22].

A total of 46,334 patients from three metropolitan areas in South Korea, newly diagnosed with cancer in 2006, survived until 2011. We excluded 544 patients who were aged <20 years at diagnosis, 2,252 patients who had missing values for covariates, and 2,639 patients who were diagnosed with CVD prior to the index date from further analyses. The final study population consisted of 40,899 cancer patients who had survived for more than 5 years after initial diagnosis. Starting from the index date of 1 January 2012, all subjects were followed-up until the date of a CVD event, death, or 31 December 2017, whichever came earliest.

2.2. Ethical considerations

This study was approved by the Seoul National University Institutional Review Board (IRB number: E-1905-148-1035). The requirement for informed consent was waived by the review board as the NHIS database is anonymized with use of its data guided by strict confidentiality guidelines.

2.3. Key variables

Information on PM concentration and size was obtained from the Air Korea database which includes data on yearly average PM2.5 and PM10 levels for the administrative district areas in South Korea. The Air Korea database collects daily PM levels using more than 300 atmospheric monitoring sites distributed across the entire country. PM2.5 levels were available for three metropolitan cities (Seoul, Busan, and Incheon) starting from 2008 allowing us to determine the average PM level
exposure from 2008-2011 for people residing in these three metropolitan cities which contain 50 administrative districts and reside over a third of the entire population of South Korea. PM2.5-10 levels were calculated by subtracting the PM10 levels from the PM2.5 levels. The average PM2.5, PM10, and PM2.5-10 levels were linked with the residential district codes for each participant, after which the participants were divided into quintiles of PM levels.

Cancer survivors were defined as participants newly diagnosed with cancer in 2006 who survived until 2011 (at least 5 years). Cancer was defined as having a cancer diagnosis code (ICD-10: C00-C99) and the critical condition code for cancer as was defined in a previous study [23]. The primary outcome, CVD, was defined as any hospitalization for coronary heart disease (CHD) or stroke for ≥2 days. Upon admission, the attending physicians are required to record a primary diagnosis for each patient according to the International Classification of Diseases, Tenth Revision codes. The ICD-10 codes for CHD (I20-I25) and stroke (I60-I69) were derived from the American Heart Association guidelines [24].

We considered the following covariates in the multivariate analysis: age (continuous, years); sex (categorical: men and women); household income (categorical: 1st, 2nd, 3rd, and 4th quintiles); area of residence (categorical: Seoul, Busan, and Incheon); and Charlson comorbidity index (continuous). Household income was derived from the insurance premium. The algorithm for calculating Charlson comorbidity index values from health claims data was adopted from a previous study [25].

2.4. Statistical analysis

Differences in the distribution of covariates according to quintiles of PM2.5 or PM10 levels were determined using the Chi-squared test for categorical variables and the analysis of variance for continuous variables. The adjusted hazard ratios (aHRs) and 95% confidence intervals (CIs) for CVD, CHD, and stroke risk were calculated using the multivariate Cox proportional hazards regression. Subjects within the lowest levels of PM2.5, PM10, and PM2.5-10 were used as the reference group. The association of PM2.5 levels on CVD risk were determined according to the cancer types which included lung, breast, liver, stomach, colorectal, kidney, bladder, prostate, and smoking-related and obesity-related cancers. Smoking-related cancers included malignancies from the head and neck, esophagus, stomach, colorectum, liver, pancreas larynx, trachea, bronchus and lung, bladder, kidney, and acute myeloid leukemia [26]. Obesity-related cancers included breast, cervical, endometrial, uterine, colon, liver, gallbladder, pancreas, kidney, and esophageal cancers [27].

Statistical significance was determined as having a p-value <0.05 in a two-sided test. All data collection and statistical analyses were conducted using SAS Enterprise Guide 7.1 (SAS Institute, Cary, NC, USA).

3. Results

3.1. Patient characteristics

Table 1 shows the characteristics of the study population. The number of subjects residing in the 1st, 2nd, 3rd, 4th, and 5th quintiles of PM2.5 levels were 8,206, 8,100, 7,837, 9,087, and 7,669, respectively. The range of PM2.5 levels in order of increasing quintile groups were 19.8-23.9, 24.1-25.3, 25.4-26.4, 26.5-27.8, and 28.2-33.8 μg/m³. Participants residing in areas with the highest PM2.5 levels were younger, had higher household income, and had less comorbidities (all p<0.001). The number of participants within the 1st, 2nd, 3rd, 4th, and 5th quintiles of PM10 levels were 8,891, 7,210, 7,984, 8,519, and 8,295, respectively. The range of PM10 levels in order of increasing quintile groups were 35.5-49.1, 49.7-50.4, 50.4-52.6, 52.7-54.1, and 54.4-61.9 μg/m³. Compared to those residing in the lowest levels of PM10, cancer survivors residing in the highest levels of PM10 were younger, had higher household income, and had less comorbidities (all p<0.001).

| Table 1. Descriptive characteristics of the study population. |
|---------------------------------------------------------------|
| Particulate matter, quintiles                                  |
| 1st (lowest)        | 2nd | 3rd | 4th | 5th (highest) | p value |
|---------------------|-----|-----|-----|---------------|---------|
|                     |  |    |    |               |         |
The association of PM2.5 levels with CVD risk is shown in Table 2. Compared to participants within the lowest quintile of PM2.5, those in the highest PM2.5 quintile group were at higher risk for CVD (aHR 1.31, 95% CI 1.07-1.59). Similarly, participants residing in the highest PM2.5 areas were at higher risk for CHD (aHR 1.47, 95% CI 1.10-1.95) compared to those within the lowest quintile. Moreover, there was a risk-increasing trend for CVD and CHD (p for trend 0.011) according to higher groups of PM2.5 quintiles. The risk for CVD according to PM10 quintile levels are shown in Table 3. Compared to those within the lowest quintile of PM10, cancer survivors with the highest exposure to PM10 did not have a higher risk for CVD (aHR 1.04, 95% CI 0.89-1.21), CHD (aHR 1.09, 95% CI 0.87-1.36), or stroke (aHR 0.99, 95% CI 0.80-1.22). Table 4 shows the association of PM2.5-10 levels with CVD risk among cancer survivors. Compared to subjects within the lowest quintile of PM2.5-10, participants residing in the highest quintile of PM2.5-10 did not have higher risk for CVD (aHR 1.04, 95% CI 0.90-1.20), CHD (aHR 1.02, 95% CI 0.83-1.25), and stroke (aHR 1.06, 95% CI 0.87-1.29). Finally, the risk of CVD among survivors from lung, breast, liver, stomach, colorectal, kidney, bladder, prostate, smoking-related, and obesity-related cancers according to PM2.5 levels is shown in Table 5. There was a tendency towards increased CVD risk for higher PM2.5 levels, although the results were not significant due to the reduced number of participants and statistical power.
Table 2. Hazard ratios for cardiovascular disease according to post-diagnosis PM$_{2.5}$ levels among 5-year cancer survivors.

|                          | PM$_{2.5}$, quintiles |                  |                  |                  |                  | $p$ for trend |
|--------------------------|------------------------|------------------|------------------|------------------|------------------|--------------|
|                          | 1$^\text{st}$ (lowest) | 2$^\text{nd}$    | 3$^\text{rd}$    | 4$^\text{th}$    | 5$^\text{th}$    |              |
| Cardiovascular disease   |                        |                  |                  |                  |                  |              |
| Events                   | 359                    | 329              | 325              | 349              | 365              |              |
| Person-years             | 45,317                 | 44,937           | 43,626           | 50,065           | 42,207           |              |
| aHR (95% CI)             | 1.00 (reference)       | 1.02 (0.86-1.19) | 1.06 (0.90-1.24) | 1.11 (0.96-1.27) | 1.31 (1.07-1.59) | 0.011        |
| Coronary heart disease   |                        |                  |                  |                  |                  |              |
| Events                   | 158                    | 133              | 133              | 202              | 166              |              |
| Person-years             | 45,317                 | 44,937           | 43,626           | 50,065           | 42,207           |              |
| aHR (95% CI)             | 1.00 (reference)       | 1.06 (0.83-1.36) | 1.11 (0.86-1.42) | 1.19 (0.96-1.46) | 1.47 (1.10-1.95) | 0.011        |
| Stroke                   |                        |                  |                  |                  |                  |              |
| Events                   | 201                    | 196              | 192              | 237              | 199              |              |
| Person-years             | 45,317                 | 44,937           | 43,626           | 50,065           | 42,207           |              |
| aHR (95% CI)             | 1.00 (reference)       | 0.98 (0.79-1.21) | 1.02 (0.83-1.26) | 1.06 (0.87-1.26) | 1.18 (0.91-1.55) | 0.257        |

Particulate matter levels determined by 4-year average levels during 2008-2011.
Adjusted hazard ratio calculated by Cox proportional hazards regression analysis after adjustments for age, sex, household income, area of residence, and Charlson comorbidity index.
Acronyms: PM, particulate matter; aHR, adjusted hazard ratio; CI, confidence interval.

Table 3. Hazard ratios for cardiovascular disease according to post-diagnosis PM$_{10}$ levels among 5-year cancer survivors.

|                          | PM$_{10}$, quintiles |                  |                  |                  |                  | $p$ for trend |
|--------------------------|-----------------------|------------------|------------------|------------------|------------------|--------------|
|                          | 1$^\text{st}$ (lowest) | 2$^\text{nd}$    | 3$^\text{rd}$    | 4$^\text{th}$    | 5$^\text{th}$    |              |
| Cardiovascular disease   |                        |                  |                  |                  |                  |              |
| Events                   | 394                   | 267              | 384              | 387              | 385              |              |
| Person-years             | 49,015                | 40,138           | 44,160           | 47,328           | 45,510           |              |
| aHR (95% CI)             | 1.00 (reference)      | 0.98 (0.83-1.15) | 1.18 (1.02-1.36) | 1.21 (1.05-1.41) | 1.04 (0.89-1.21) | 0.078        |
| Coronary heart disease   |                        |                  |                  |                  |                  |              |
| Events                   | 181                   | 105              | 162              | 157              | 185              |              |
| Person-years             | 49,015                | 40,138           | 44,160           | 47,328           | 45,510           |              |
| aHR (95% CI)             | 1.00 (reference)      | 0.88 (0.69-1.13) | 1.14 (0.92-1.42) | 1.18 (0.94-1.48) | 1.09 (0.87-1.36) | 0.111        |
| Stroke                   |                        |                  |                  |                  |                  |              |
| Events                   | 213                   | 162              | 222              | 228              | 200              |              |
| Person-years             | 49,015                | 40,138           | 44,160           | 47,328           | 45,510           |              |
| aHR (95% CI)             | 1.00 (reference)      | 1.05 (0.85-1.30) | 1.20 (0.99-1.46) | 1.24 (1.02-1.51) | 0.99 (0.80-1.22) | 0.348        |

Particulate matter levels determined by 4-year average levels during 2008-2011.
Adjusted hazard ratio calculated by Cox proportional hazards regression analysis after adjustments for age, sex, household income, area of residence, and Charlson comorbidity index.
Acronyms: PM, particulate matter; aHR, adjusted hazard ratio; CI, confidence interval.
Table 4. Hazard ratios for cardiovascular disease according to post-diagnosis PM$_{2.5-10}$ levels among 5-year cancer survivors.

| PM$_{2.5-10}$, quintiles | 1$^\text{st}$ (lowest) | 2$^\text{nd}$ | 3$^\text{rd}$ | 4$^\text{th}$ | 5$^\text{th}$ (highest) | p for trend |
|--------------------------|-----------------------|---------|---------|---------|----------------------|------------|
| Cardiovascular disease   |                       |         |         |         |                      |            |
| Events                   | 419                   | 326     | 286     | 423     | 363                  |            |
| Person-years             | 46,989                | 43,996  | 42,215  | 50,648  | 42,304               |            |
| aHR (95% CI)             | 1.00 (reference)      | 1.03 (0.88-1.20) | 0.94 (0.80-1.11) | 1.10 (0.95-1.27) | 1.04 (0.90-1.20) | 0.361      |
| Coronary heart disease   |                       |         |         |         |                      |            |
| Events                   | 203                   | 128     | 116     | 178     | 167                  |            |
| Person-years             | 46,989                | 43,996  | 42,215  | 50,648  | 42,304               |            |
| aHR (95% CI)             | 1.00 (reference)      | 0.92 (0.72-1.17) | 0.87 (0.68-1.11) | 1.03 (0.82-1.28) | 1.02 (0.83-1.25) | 0.576      |
| Stroke                   |                       |         |         |         |                      |            |
| Events                   | 216                   | 198     | 170     | 245     | 196                  |            |
| Person-years             | 46,989                | 43,996  | 42,215  | 50,648  | 42,304               |            |
| aHR (95% CI)             | 1.00 (reference)      | 1.11 (0.90-1.37) | 1.00 (0.80-1.24) | 1.16 (0.95-1.41) | 1.06 (0.87-1.29) | 0.468      |

Particulate matter levels determined by 4-year average levels during 2008-2011.
PM$_{2.5-10}$: particulate matter with sizes between 2.5 and 10 μm.
Adjusted hazard ratio calculated by Cox proportional hazards regression analysis after adjustments for age, sex, household income, area of residence, and Charlson comorbidity index.
Acronyms: PM, particulate matter; aHR, adjusted hazard ratio; CI, confidence interval.
Table 5. Association of post-diagnosis PM₂.₅ levels with cardiovascular disease risk among 5-year cancer survivors according to cancer types.

| Cancer Type          | PM₂.₅, quintiles | p for trend |
|----------------------|------------------|-------------|
|                      | 1st (lowest)     | 2nd          | 3rd          | 4th          | 5th (highest) |
| Lung cancer          |                  |              |              |              |              |
| Events               | 12               | 9            | 10           | 9            | 12           |
| Person-years         | 841              | 799          | 846          | 946          | 783          |
| aHR (95% CI)         | 1.00 (reference) | 0.71 (0.29-1.73) | 0.83 (0.34-1.98) | 0.79 (0.33-1.89) | 2.46 (0.93-6.52) | 0.389 |
| Breast cancer        |                  |              |              |              |              |
| Events               | 34               | 30           | 41           | 36           | 48           |
| Person-years         | 8,013            | 8,016        | 7,250        | 8,591        | 7,461        |
| aHR (95% CI)         | 1.00 (reference) | 1.08 (0.63-1.86) | 1.60 (0.97-2.65) | 1.04 (0.65-1.66) | 2.25 (1.32-3.85) | 0.045 |
| Liver cancer         |                  |              |              |              |              |
| Events               | 14               | 9            | 10           | 5            | 12           |
| Person-years         | 1,006            | 962          | 996          | 1,171        | 853          |
| aHR (95% CI)         | 1.00 (reference) | 0.90 (0.35-2.31) | 1.04 (0.41-2.63) | 0.34 (0.12-0.96) | 0.84 (0.23-2.98) | 0.107 |
| Stomach cancer       |                  |              |              |              |              |
| Events               | 79               | 70           | 77           | 90           | 81           |
| Person-years         | 7,618            | 7,601        | 6,931        | 8,867        | 7,624        |
| aHR (95% CI)         | 1.00 (reference) | 0.96 (0.68-1.35) | 1.14 (0.81-1.60) | 0.99 (0.73-1.35) | 1.33 (0.88-2.02) | 0.379 |
| Colorectal cancer    |                  |              |              |              |              |
| Events               | 81               | 70           | 54           | 86           | 62           |
| Person-years         | 6,160            | 6,007        | 5,630        | 6,466        | 5,588        |
| aHR (95% CI)         | 1.00 (reference) | 0.88 (0.62-1.24) | 0.77 (0.53-1.12) | 1.00 (0.74-1.36) | 1.17 (0.74-1.86) | 0.630 |
| Kidney cancer        |                  |              |              |              |              |
| Events               | 13               | 6            | 6            | 15           | 11           |
| Person-years         | 1,154            | 1,079        | 1,070        | 1,182        | 947          |
| aHR (95% CI)         | 1.00 (reference) | 0.50 (0.18-1.38) | 0.55 (0.20-1.54) | 1.07 (0.50-2.28) | 1.65 (0.60-4.56) | 0.301 |
| Bladder cancer       |                  |              |              |              |              |
| Events               | 13               | 23           | 13           | 29           | 19           |
| Person-years         | 1,117            | 1,062        | 1,061        | 1,518        | 966          |
| aHR (95% CI)         | 1.00 (reference) | 2.86 (1.34-6.12) | 1.62 (0.70-3.75) | 1.77 (0.91-3.43) | 2.44 (1.00-5.95) | 0.210 |
| Prostate cancer      |                  |              |              |              |              |
| Events               | 20               | 17           | 21           | 28           | 15           |
| Person-years         | 1,115            | 1,073        | 1,502        | 1,276        | 737          |
| aHR (95% CI)         | 1.00 (reference) | 0.86 (0.43-1.69) | 0.83 (0.43-1.59) | 1.26 (0.70-2.26) | 1.23 (0.52-2.90) | 0.304 |
| Smoking-related cancer |                |              |              |              |              |
| Events               | 226              | 201          | 185          | 253          | 218          |
| Person-years         | 19,067           | 18,508       | 17,502       | 21,442       | 17,977       |
| aHR (95% CI)         | 1.00 (reference) | 0.99 (0.81-1.22) | 0.99 (0.80-1.22) | 1.00 (0.84-1.20) | 1.34 (1.04-1.73) | 0.201 |
| Obesity-related cancer |               |              |              |              |              |
| Events               | 166              | 137          | 133          | 174          | 161          |
| Person-years         | 21,160           | 21,073       | 18,987       | 22,623       | 19,772       |
| aHR (95% CI)         | 1.00 (reference) | 0.88 (0.69-1.13) | 0.97 (0.76-1.24) | 0.98 (0.79-1.21) | 1.33 (0.99-1.78) | 0.209 |

Particulate matter levels determined by 4-year average levels during 2008-2011.
Smoking-related cancer included head and neck, esophagus, stomach, colorectum, liver, pancreas larynx, trachea, bronchus and lung, bladder, kidney cancer, and acute myeloid leukemia.
Obesity-related cancer included breast, cervical, endometrial, uterine, colon, liver, gallbladder, pancreas, kidney, and esophageal cancer.
Adjusted hazard ratio calculated by Cox proportional hazards regression analysis after adjustments for age, sex, household income, area of residence, Charlson comorbidity index.
Acronyms: PM, particulate matter; aHR, adjusted hazard ratio; CI, confidence interval.

4. Discussion

In the three-metropolitan areas with over 40,000 cancer survivors, long-term exposure to PM was associated with increased CVD incidence. To our knowledge, this is the first study to identify an increasing effect of PM2.5 levels on CVD in long-term cancer survivors. The higher the concentration of PM2.5, the more incident CVD, confirming the concentration-response relationship. The hazard ratio for CVD increased from PM2.5 concentrations of 28.2 µg/m³.

In previous studies, PM2.5 has been reported to be the most pathogenic for CVD in the general population, with PM2.5-10 and PM10 levels having inconsistent results [15,28,29]. Our study also
showed similar results. We found that a yearly average concentration of 28.2 µg/m³, which is above the yearly PM2.5 regulation guideline in South Korea (15 µg/m³), US (12 µg/m³), European Union (25 µg/m³), and Japan (15 µg/m³) [30], increased the risk for CVD.

In the general population, an increase in the PM2.5 concentration by 10 µg/m³ was associated with an increase in the number of hospital admissions for coronary artery disease, arrhythmias, heart failure, cerebrovascular disease, and peripheral artery disease [31]. The susceptible population included children, older adults, those that were obese, had low socioeconomic status (SES), and specific genetic factors [32]. Unlike traditional Framingham risk factors, women were more susceptible to CVD when exposed to PM2.5 than men probably due to their smaller coronary arteries and micro vessels [33-36].

In addition to those risk factors, cancer survivors may be more susceptible to the harmful effects of PM. They are often asymptomatic but several times higher prevalence of CVD compared with the Framingham Heart Study population [37]. Chemotherapy agents for cancer patients have several vascular complications including endothelial dysfunction, capillary rarefaction, vascular remodeling, oxidative stress, platelet activation, thrombosis, and vasospasm [38]. In addition, radiotherapy impairs diastolic and systolic function [39]. The time-lapse between chemotherapy and radiotherapy and incident CVD can be very long, lasting even more than 10 years [40,41].

In the present study, breast and bladder cancer survivors were the most affected by PM2.5 which is consistent with a previous study [19]. PM seems to have a greater influence on CVD-susceptible cancers of all kind. In the previous study, endometrial, breast, melanoma, prostate, and urinary bladder cancer were susceptible among cancers in 28 sites [42]. Breast cancer survivors treated with taxanes, vascular endothelial growth factor (VEGF) inhibitor, and hormonal agents experience increased thrombotic events, severe hypertension, and heart failure [38,39].

The relationship between bladder cancer and the PM remains controversial. Environmental and socioeconomic factors are known to affect bladder cancer mortality [43]. Those engaged in occupations such as taxi driver, truck driver, or street vendor are highly exposed to air pollution and have been reported to be at increased risk for bladder cancers [44]. However, there was no association reported between PM2.5 concentrations and bladder cancer incidence in 15 European cohort studies and one Spanish study [45,46]. The relationship between non-cancer mortality and PM has scarce evidence for bladder cancers s patients with bladder cancers had high to no risk for CVD in the previous studies [7,42]. In our long-term bladder cancer survivors, the concentration of 24.1-25.3 and 28.2-33.8 µg/m³ of PM2.5 was associated with CVD. Further research in other large-scale populations is needed.

In the present study, there was no association between PM concentration and stroke in cancer survivors. Both hemorrhagic and ischemic stroke risk were increased in cancer patients [47,48] even after 10 years of surviving [49] in previous studies. This phenomenon may need to be investigated further.

Our study has several limitations. First, operationally defined CVD may be incorrectly coded. Unstable angina is one of the most up coded conditions [50] and there is also the possibility that patients with multiple CVD could be down coded should the doctor not have recorded all ICD codes for multiple complex CVD. However, the operational diagnosis was validated and had been widely used in previous studies [51,52]. Second, ischemic and hemorrhagic strokes could not be differentiated due to lack of data. Third, the study population was from three metropolitan cities, so the results may not be generalizable to rural areas. Fourth, the residential areas and areas where the population spends most of their time (such as the workplace) may be different. Fifth, the individual CVD risk factors (smoking, diet, alcohol use, access to care, obesity, blood pressure, lipid profile, and family history) were not identified, although non-smokers had a greater risk for air-pollution related disease than current smokers in the previous study [53].

5. Conclusions

We found that higher levels of PM2.5 were associated with increased CVD among cancer survivors, implying that cancer survivors are a vulnerable group when exposed to PM. Among the
various cancer types, breast and bladder cancer survivors were the most affected. Future studies will be needed to evaluate the impact of PM exposure reduction among cancer survivors.

Author Contributions:

SM Park had full access to all of the data in the study and takes responsibility for the integrity of the data and the accuracy of the data analysis; Study concept and design: SM Park; Acquisition of data: S Choi, K Kim, SM Park; Analysis and interpretation of data: S Choi, KH Kim, K Kim, J Chang, SM Kim, SR Kim, Y Cho, G Lee, JS Son, SM Park; Drafting of the manuscript: S Choi, KH Kim, SM Park; Critical revision of the manuscript: S Choi, KH Kim, K Kim, J Chang, SM Kim, SR Kim, Y Cho, G Lee, JS Son, SM Park; Statistical analysis: S Choi; Administrative, technical, or material support: K Kim, SM Kim

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