

Article

Seasonal Variations in the Daily Mortality Associated with Exposure to Particles, Nitrogen Dioxide, and Ozone in Stockholm, Sweden, from 2000 to 2016

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Abstract: Urban air pollutant emissions and concentrations vary throughout the year due to various factors, e.g., meteorological conditions and human activities. In this study, seasonal variations in daily mortality associated with increases in the concentrations of PM₁₀ (particulate matter), PM_{2.5–10} (coarse particles), BC (black carbon), NO₂ (nitrogen dioxide), and O₃ (ozone) were calculated for Stockholm during the period from 2000 to 2016. The excess risks in daily mortality are presented in single and multi-pollutant models during the whole year and divided into four different seasons, i.e., winter (December–February), spring (March–May), summer (June–August), and autumn (September–November). The excess risks in the single-pollutant models associated with an interquartile range (IQR) increase for a lag 02 during the whole year were 0.8% (95% CI: 0.1–1.4) for PM₁₀, 1.1% (95% CI: 0.4–1.8) for PM_{2.5–10}, 0.5% (95% CI: –0.5–1.5) for BC, –1.5% (95% CI: –0.5––2.5) for NO₂, and 1.9% (95% CI: 1.0–2.9) for O₃. When divided into different seasons, the excess risks for PM₁₀ and PM_{2.5–10} showed a clear pattern, with the strongest associations during spring and autumn, but with weaker associations during summer and winter, indicating increased risks associated with road dust particles during these seasons. For BC, which represents combustion-generated particles, the pattern was not very clear, but the strongest positive excess risks were found during autumn. The excess risks for NO₂ were negative during all seasons, and in several cases even statistically significantly negative, indicating that NO₂ in itself was not harmful at the concentrations prevailing during the measurement period (mean values < 20 µg m^{–3}). For O₃, the excess risks were statistically significantly positive during “all year” in both the single and the multi-pollutant models. The excess risks for O₃ in the single-pollutant models were also statistically significantly positive during all seasons.

Keywords: PM₁₀; PM_{2.5–10}; BC; NO₂; O₃; excess health risks; multi-pollutant models; seasons



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1. Introduction

While air pollutant emissions occur throughout the whole year, differences in meteorological conditions and human activities cause large seasonal variations in most pollutants. Kukkonen et al. (2000) [1] showed that meteorological variables in northern Europe vary widely with the seasons and, consequently, those variations are reflected in urban air quality. Apart from anthropogenic and natural emissions, among the main factors influencing seasonal variations in air quality are occurrence of temperature inversions, wind speed, precipitation, and solar radiation [2]. Particularly in the Nordic countries, where studded winter tires are used, suspension of coarse particles from road surfaces is highly dependent

on surface moisture, which shows very large variations during the year, and with a major impact on the seasonal variations in the concentrations of both PM₁₀ and PM_{2.5} [3].

The seasonal variations in air quality give rise to differences in health effects. In a study from the Netherlands, the associations between daily mortality and short-term variations in a number of air pollutants were analyzed during the period from 1986 to 1994, and the results were, among other things, divided into summer and winter seasons [4]. They found significantly (95% CI) increased relative risks for all pollutants (PM₁₀, BS (black smoke), O₃, SO₂, NO₂, CO, SO₄⁻², and NO₃⁻) during both summer and winter, except for O₃ and SO₄⁻² during the winter. Moreover, the relative risks for total mortality associated with exposure to these air pollutants were in all cases larger during the summer compared to the winter [4].

Seasonal variations regarding short-term health effects associated with exposure to PM₁₀ have been analyzed in a number of studies. When increased mortality associated with an increase in PM₁₀ was analyzed in 100 cities in the U.S. in the period from 1987 to 2000, a significant association was found for the summer period, but no significant associations were found for the other three seasons. The seasonal pattern was also more pronounced in the northeast region of the U.S., while there were relatively small seasonal variations in the southern regions [5]. In Flanders, Belgium, where daily mortality associated with PM₁₀ was calculated during the period from 1997 to 2003, and where temperature and seasons were included as potential effect modifiers, the strongest associations were also found during the summer [6]. Similarly, a significant effect of PM₁₀ on mortality increase during the summer was found in Tallinn, Estonia, during the period from 2004 to 2011 [7]; however, during the winter, a significant decrease in mortality associated with an increase in PM₁₀ appeared. In a study from China, where the association between daily mortality and an increase in PM₁₀ was analyzed in 17 Chinese cities during different seasons, significant associations were found for summer and winter, but not for spring and autumn [8]. In another study from Wuhan, China, conducted from 2001 to 2004, the strongest associations for PM₁₀ were found during winter [9]. In a study from Korea, where the associations between PM₁₀ concentrations and increases in mortality and hospital admissions were analyzed in Seoul from 2000 to 2006, the effects on both mortality and morbidity increased during the summer [10]. Finally, in a study from Utah, U.S., the association between daily mortality and exposure to PM₁₀ was examined for the period 1985 to 1992. The largest contribution to excess mortality was for individuals 75+ years old dying in a hospital, and the strongest effect was shown during the spring [11].

Seasonal variations regarding short-term effects on hospitalization for cardiovascular diseases associated with PM_{2.5} were analyzed in New York State during the period 1991 to 2006. The strongest effects, associated with a 10 µg m⁻³ increase in PM_{2.5}, were found during the winter. Temperature modified the PM_{2.5} effects on cardiovascular diseases, and these effects were found on low temperature days [12]. In a study in Tallinn, PM_{2.5} increased mortality during the summer, but no effects appeared during the winter [7].

For particles in the coarse fraction (PM_{2.5-10}), the difference in short-term effects on daily mortality between two annual periods was analyzed in Stockholm during the period from 2000 to 2008. The associations between PM_{2.5-10} and daily mortality were stronger during November to May in comparison with the rest of the year, which can be explained by the high levels of road dust that occurs because of the use of studded tires in Stockholm during this time of the year [13].

The seasonal variations in short-term effects associated with NO₂ were analyzed in a study from China. The daily mortality associated with NO₂ in the city of Shenzhen in southeastern China during the period from 2013 to 2017 was analyzed during the cold season (November–April) and during the warm season (May–October). Significant excess risks for cardiovascular mortality associated with an increase in NO₂ were found during the cold season at a 2-day lag and at a 6-day lag. However, no significant excess risks were found during the warm season [14].

The short-term health effects associated with O₃ and their seasonal variations have been studied in U.S., France, and China. When the short-term mortality effects associated with O₃ were analyzed in 20 communities in U.S., and where ten communities represented the northern part and ten represented the southern part, the seasonal variations in the effects estimates exhibited different results in the northern and southern parts. In the southern communities, an increase in O₃ entailed increases in mortality during autumn and winter, while there were negative excess risks during spring and summer. In the northern communities, an increase in mortality was found during spring, summer and autumn, while there was a negative excess risk during the winter. In this study, latitude and seasonal average temperature were identified as effect modifiers [15]. In a study in nine French urban areas during the period from 1998 to 2006, the association between daily mortality and the daily max-8 h O₃ concentrations was analyzed by season and by temperature strata. The strongest mortality effects were found during the summer and for the highest temperature strata [16]. In a study from China, the association between O₃ and daily mortality was analyzed in the city of Zhengzhou during the period from 2013 to 2015. Significant excess risks associated with an increase in 10 µg m⁻³ 24-h average O₃ concentrations at a 1-day lag were found during the cold season, but not during the warm season [17].

Based on the relatively few studies referenced above, it seems like the strongest excess risks of mortality for PM₁₀ (seven studies) and PM_{2.5} (only two studies) occur during the summer months. For NO₂ and O₃, no consistent results in terms of seasonal dependence of excess risks can be seen. However, there are very few studies: only one study for NO₂, and only three studies for O₃.

The purpose of the current study was to analyze the seasonal variations in the effect estimates of air pollution on mortality in Stockholm. In the earlier analyzes, only annual data were considered, but the seasonal variations were not taken into account [18]. A special focus was to analyze the causes of the negative excess risks associated with NO₂, and whether the seasons and other pollutants had any effect on these associations. Seasonal differences during the year can potentially affect the results in several ways. People are more likely to stay outdoors and have windows opened during the warm seasons, which can affect the degree of exposure. Annual variations in meteorology and air pollution sources can also be of importance. In Stockholm, the chemical composition of PM₁₀ and PM_{2.5-10} varies throughout the year with a significantly higher proportion of mechanically generated road dust particles during early spring [3]. Analyzing the associations between mortality and short-term exposure to the above-mentioned air pollutants during different seasons in Stockholm is, therefore, of great interest.

2. Materials and Methods

This study includes residents of Stockholm, with a population that increased from 0.8 to 0.9 million during the period from 2000 to 2016. Population data were obtained from the Swedish Central Bureau of Statistics. Cause of mortality data were obtained from the National Cause of Death Register. Natural cause of mortality is defined on the basis of the underlying cause of death, and these data include the daily number of deaths from non-external causes (ICD-10: A00–R99) occurring among the registered population.

Air pollution exposure was estimated from a central measuring station on the roof-top of a 20 m high building in the central part of Stockholm. The monitoring station was part of the city's regulatory air pollution control network, and equipped with reference (or equivalent) instruments for regulated pollutants according to the EU air quality directive. These air pollutants included PM₁₀ (particles with an aerodynamic diameter smaller than or equal to 10 µm), PM_{2.5} (particles with an aerodynamic diameter smaller than or equal to 2.5 µm), NO₂ (nitrogen dioxide), and O₃ (ozone) (Table 1). The O₃ measurements were based on daily maximum 8-h mean values. In addition, the monitoring included measurements of unregulated black carbon (BC), and particles in the coarse fraction (PM_{2.5-10}) estimated by subtracting PM_{2.5} from PM₁₀. The period from 2000 to 2016 was divided into

winter (December–February), spring (March–May), summer (June–August), and autumn (September–November) seasons.

Table 1. Measurement methods and instruments used to measure the pollutants.

Pollutant	Method	Instrument
PM ₁₀	Gravimetric	TEOM 1400A, Thermo Fisher Scientific, Waltham, MA, USA
PM _{2.5}	Gravimetric	TEOM 1400A, Thermo Fisher Scientific, Waltham, MA, USA
PM _{2.5–10}	Gravimetric (Subtracting PM _{2.5} from PM ₁₀)	TEOM 1400A, Thermo Fisher Scientific, Waltham, MA, USA
BC	Transmission of lightthrough a filter	Aethalometers 8100, AE31, AE33, Magee Scientific Corporation, Berkeley, CA, USA
NO ₂	Chemiluminescence	AC32M, Environnement S.A., Poissy, France
O ₃	UV absorption	O342M, Environnement S.A., Poissy, France

Temperature data were collected from the urban meteorological station Observatoire-lunden. In this study, daily maximum temperature was used as exposure variable.

The associations between different air pollutants and daily mortality were modelled using a quasi-Poisson regression model with a logistic link function. The concept “quasi-Poisson” refers to a model that adjusts for overdispersed data, and a logistic link function defines the relationship of the dependent variables to the mean of the Poisson distributed independent variables. The modeling procedure was replicated from a previous study [18] in order to ensure comparability of the results. The model estimated the effect of an interquartile range (IQR) increase in air pollutants on daily mortality for lag 02 (average concentration during the same and the previous two days). The IQR values were calculated based on data for the whole year. Adjustments for other time-varying factors were made by assuming a linear additive effect on a logarithmic scale:

$$\text{Log}(Y_i) = \text{Intercept} + AP_i + W_i + \text{DOW}_i + \text{long-term trend} \quad (1)$$

where Y_i represents the daily number of deaths from non-external causes, AP_i represents the concentration of a specific or a combination of air pollutants on day i , W_i represents variables controlling for the weather on day i using smooth spline functions for the maximum temperature and snowfall, DOW_i represents the day of the week, and the long-time trend is a smooth function varying over time to capture any long-term and seasonal patterns in mortality. The effects of air pollution were estimated by using a seasonal factor resulting in individual dose-responses for each season while keeping the other variable estimates constant for the whole period. Snowfall was included since it is a risk factor for daily mortality, as described in Auger et al. (2017) [19]. The smooth function describing the long-term time trend was a penalized regression spline restricted to 5 d.f. (degrees of freedom) per year. All pollutants were modelled by assuming a linear relationship with daily mortality. Air pollutants were first modelled in single-pollutant models, and traffic-related pollutants were included in multi-pollutant models together with O₃ and PM_{2.5–10}. Temperature effects were adjusted by using two different smooth functions corresponding to the different lag-windows of 0–2 and 3–10. The model allowed for the use of 4 d.f. for each function. All analyses were conducted using R statistical software version 3.6.0 (R Foundation for Statistical Computing, Vienna, Austria).

3. Results

3.1. Descriptive Data

In Table 2, summary statistics of the daily data regarding mortality, maximum temperature, and the measured air pollutants during the period from 2000 to 2016 are presented as mean values and IQR values for the whole year, and divided into four different seasons. PM₁₀, PM_{2.5-10} and O₃ show the highest concentrations during spring and the lowest during autumn/winter. For NO₂ and BC, the highest values are seen during winter and the lowest during summer. The seasonal variations reflect the different importance of emissions as discussed by Olstrup et al. (2019) [18].

Table 2. Summary statistics of the daily data from 2000 to 2016 during whole year and in four different seasons.

Variable	Whole Year	Winter	Spring	Summer	Autumn
			Mean (IQR)		
Mortality (cases per day)	18.5 (7.0)	20.2 (6.0)	18.7 (6.0)	17.0 (6.0)	17.9 (6.0)
Maximum temperature (°C)	11.4 (15.0)	1.4 (5.6)	10.9 (9.8)	22.2 (5.0)	11.0 (8.3)
PM ₁₀ (µg m ⁻³)	14.5 (8.7)	12.2 (7.7)	20.5 (13.7)	12.9 (5.9)	12.2 (6.7)
PM _{2.5-10} (µg m ⁻³)	8.0 (5.5)	5.5 (3.7)	13.1 (10.8)	6.8 (3.8)	6.3 (3.8)
BC (µg m ⁻³)	0.6 (0.5)	0.7 (0.6)	0.6 (0.4)	0.5 (0.5)	0.7 (0.5)
NO ₂ (µg m ⁻³)	14.4 (9.9)	17.6 (11.5)	14.0 (9.4)	10.8 (7.4)	15.2 (9.6)
O ₃ (µg m ⁻³) *	51.2 (25.2)	41.6 (20.1)	66.1 (17.4)	58.0 (17.5)	38.8 (18.0)

* O₃ measurements were based on daily maximum 8-h mean values.

The number of days with valid data is presented for each variable in Table A1 in Appendix A. Pearson correlation coefficients between the different air pollutants during all year and divided into different seasons are presented in Tables A2–A6 in Appendix A.

3.2. Calculated Excess Risks

Figures 1–6 show the calculated excess risks for daily mortality associated with an IQR increase in the measured pollutants in Stockholm during the period from 2000 to 2016. The measured air pollutants are presented in Table 2. The figures are divided into different sections where the whole year is presented furthest to the left, and then from left to right winter (December–February), spring (March–May), summer (June–August), and autumn (September–November). Figure 1 presents a single-pollutant model, and Figures 2–6 present multi-pollutant models where adjustments for other pollutants have been made. The excess risks for daily mortality associated with exposure to the different air pollutants were based on lag02, which means a lagging effect of the air pollution exposure during the same and the previous two days.

The excess risks associated with exposure to PM₁₀ and PM_{2.5-10} (Figures 2 and 3) showed a clear pattern with the strongest associations during spring and autumn, but with weaker associations during summer and winter. For BC (Figure 4), which represents combustion-generated particles, the pattern was not very clear, but the strongest positive excess risks were found during autumn. The excess risks associated with exposure to NO₂, presented in Figure 5, were significantly (95% CI) negative based on “all year”, and especially strong negative associations were shown during the autumn after adjustments for PM_{2.5-10} and BC. For O₃ (Figure 6), the excess risks were significantly (95% CI) positive during “all year” in both the single and the multi-pollutant models. The single-pollutant models were also significantly positive during all seasons, and the highest excess risk appeared during the autumn.

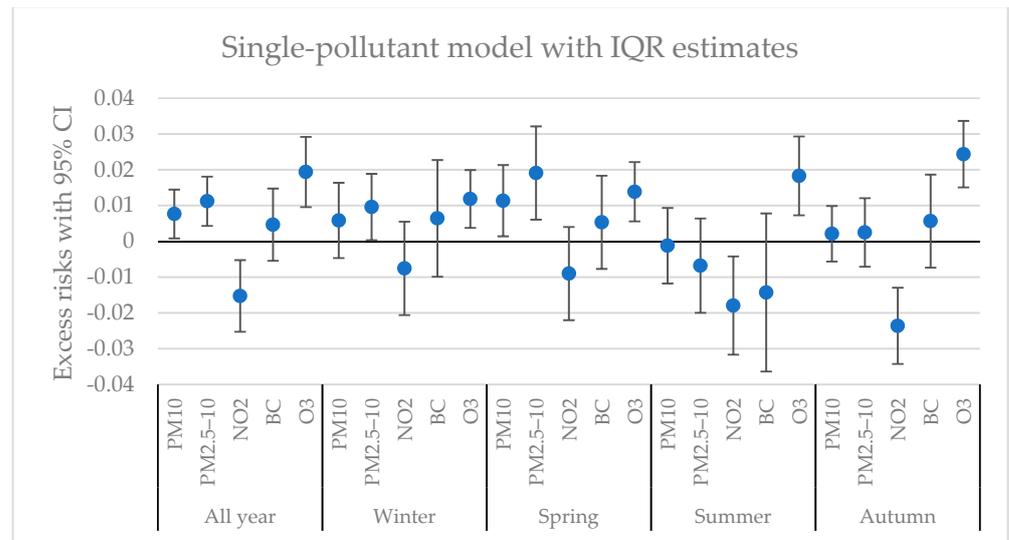


Figure 1. Estimated excess risks in daily mortality with 95% CI (lag02) for an IQR increase in the concentrations of PM₁₀, PM_{2.5-10}, NO₂, BC and O₃ in single-pollutant models divided into all year and four different seasons.

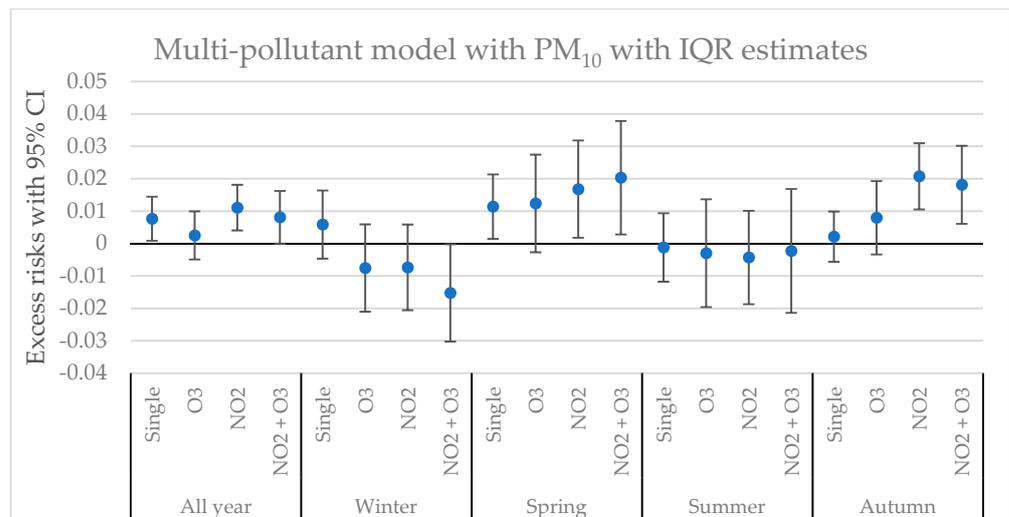


Figure 2. Estimated excess risks in daily mortality with 95% CI (lag02) for an IQR increase in the concentrations of PM₁₀. The calculations are divided into all year and four different seasons. The excess risks are presented as single-pollutant estimates and multi-pollutant estimates with O₃, NO₂ and NO₂ + O₃ included in the calculations.

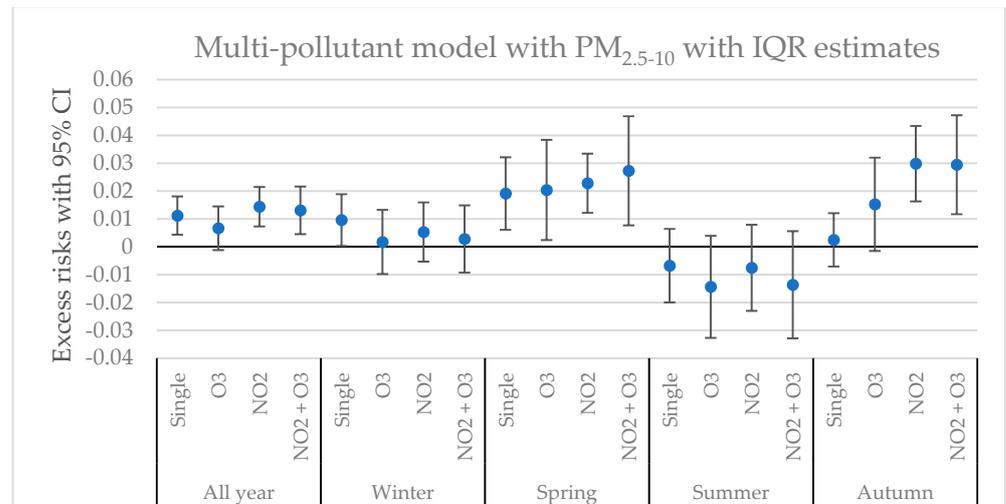


Figure 3. Estimated excess risks in daily mortality with 95% CI (lag02) for an IQR increase in the concentrations of PM_{2.5-10}. The calculations are divided into all year and four different seasons. The excess risks are presented as single-pollutant estimates and multi-pollutant estimates with O₃, NO₂ and NO₂ + O₃ included in the calculations.

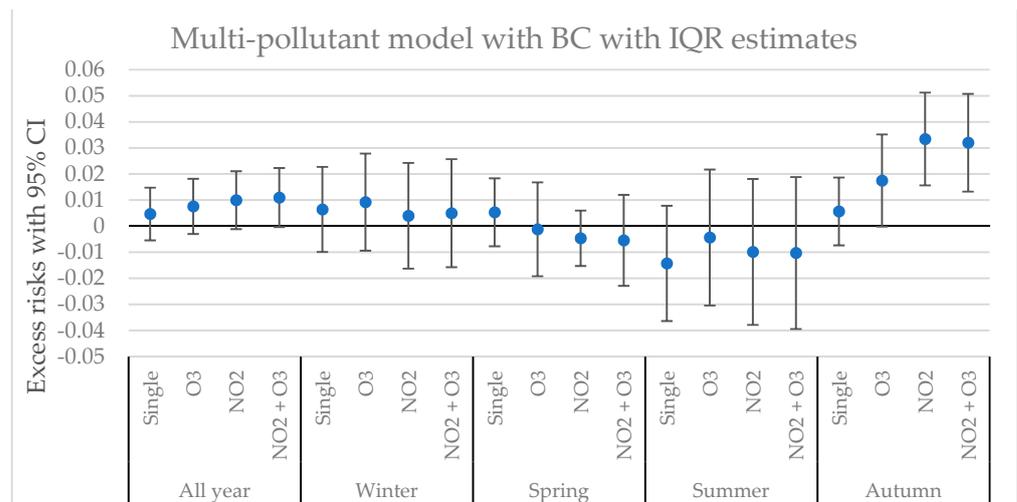


Figure 4. Estimated excess risks in daily mortality with 95% CI (lag02) for an IQR increase in the concentrations of BC. The calculations are divided into all year and four different seasons. The excess risks are presented as single-pollutant estimates and multi-pollutant estimates with O₃, NO₂ and NO₂ + O₃ included in the calculations.

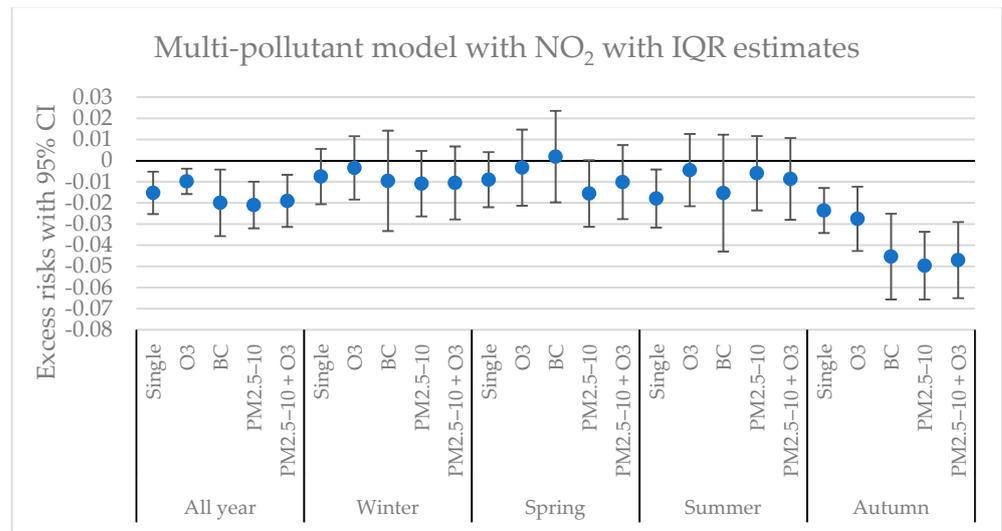


Figure 5. Estimated excess risks in daily mortality with 95% CI (lag 02) for an IQR increase in the concentrations of NO₂. The calculations are divided into all year and four different seasons. The excess risks are presented as single-pollutant estimates and multi-pollutant estimates with O₃, BC, PM_{2.5-10} and PM_{2.5-10} + O₃ included in the calculations.

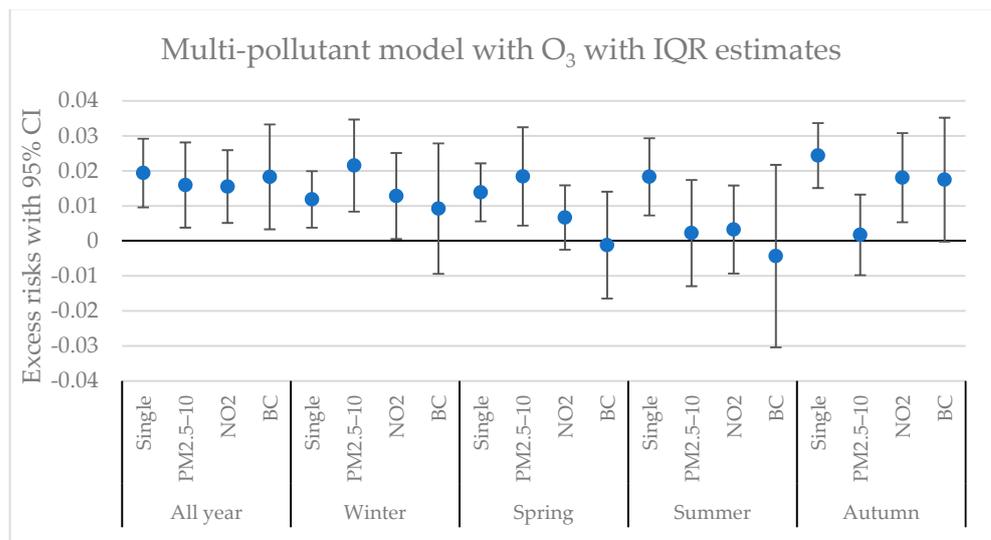


Figure 6. Estimated change in daily mortality with 95% CI (lag02) for an IQR increase in the concentrations of O₃. The calculations are divided into all year and four different seasons. The excess risks are divided into single-pollutant estimates and multi-pollutant estimates with PM_{2.5-10}, NO₂ and BC included in the calculations.

4. Discussion

Calculated Excess Risks and Possible Explanations for Their Seasonal Variations

The main objective of this study was to analyze if the daily mortality associated with exposure to the different air pollutants were different when different seasons were analyzed separately, and if the results were significantly different compared to when the analyses were based on the whole year without regard to the seasons. An important issue was whether behavioral factors regarding the amount of outdoor stay, which varies depending on the season, could have affected the results.

The excess risks associated with PM_{2.5-10} and PM₁₀ exhibited a broadly similar pattern with larger excess risks in daily mortality during springtime compared to the other seasons. The seasonal pattern of the excess risks for PM₁₀ and PM_{2.5-10} can possibly be explained by

the seasonal variation in the chemical composition throughout the year. During springtime, up to 90% of the mass of PM₁₀ in Stockholm originates from road abrasion that occurs when the road surfaces become drier [3]. The large proportion of PM₁₀ during springtime is road dust that is mainly represented in the coarse fraction (PM_{2.5–10}), and this is mainly caused by the use of studded tires during the winter months [20]. A large number of detrimental effects on the respiratory and the cardiovascular system linked to the components of road dust particles have been showed in a literature review from 2018 [21].

The excess risks associated with BC were statistically significant only in two out of the studied 20 cases. In terms of percentage of days with valid data, BC differed from the other pollutants by having a data coverage of just over 50% in comparison with NO₂, O₃, and PM₁₀ with a data coverage of 96–100%, and PM_{2.5–10} with a data coverage of 83–88% (Table A1, Appendix A). A relatively smaller amount of data might have made the excess risk estimates less accurate. However, the data coverage for BC was greater during the latter part of the measurement period. Additionally, in a previous study, where BC measurements were compared at different measuring sites in Stockholm, the spatiotemporal variability was not found to be very high, and different urban sites were poorly correlated even for daily averages ($R < 0.70$) [22]. Consequently, this means that the use of measurement data for BC from one single measurement station may have led to exposure misclassification, leading to even greater uncertainties regarding the calculations of the excess risks.

The reasons for the negative excess risks associated with NO₂ are not entirely clear. The hypothesis in the previous study [18] was that seasonal variations in exposure could be an explanation, and that the exposure was greater during the warm season. In that case, an expected result would be that the excess risks during the summer would be greater compared to the winter, but no indications of higher excess risks during the warm season could be seen. However, a lack of positive associations between daily mortality and exposure to NO₂ has been shown for Stockholm in previous studies as well [23,24]. It may be possible that NO₂ in itself, at these low concentrations, has no significant effects on short-term mortality, but that oxidants in the air might have a crucial importance.

The lack of positive excess risks associated with NO₂ may also be due to concentrations that were below the threshold for adverse health effects. Experimental studies with humans demonstrated noticeable effects after short-term exposure to NO₂ concentrations at or above 400 $\mu\text{g m}^{-3}$, and health effects on patients with mild asthma could not be detected after short-term exposure to concentrations below 200 $\mu\text{g m}^{-3}$ [25]. Based on a review of several studies on exposure to NO₂, increased mortality was suggested above a threshold value of 20 $\mu\text{g m}^{-3}$ [26]. The average NO₂ concentrations in this study were below 20 $\mu\text{g m}^{-3}$ during all seasons. An earlier analysis including older data with higher NO₂ values (median concentration of 26.8 $\mu\text{g m}^{-3}$) in Stockholm increased the mortality risk [27], whereas the current study with lower concentrations did not. However, in the former study, the control for temperature effects on daily mortality included only a short lag. In this study, the control for low temperature effects included a lag-window of 3–10 days. There is a possibility that the temperature adjustments capture some effects of exhaust gases. Another factor that could contribute to the lack of positive associations with daily mortality is exposure misclassification as discussed above in the case of BC.

Effect modifications by oxidants regarding the health effects associated with PM_{2.5} were shown in a multi-city case-crossover study in Canada where the relationships between PM_{2.5} and daily mortality were greater in connection with elevated oxidant gas concentrations [28]. The importance of reactive oxygen species (ROS) has also been addressed in a cohort study from Canada where the association between ROS in lung fluid and exposures to PM_{2.5} containing Fe and Cu were estimated using land use regression models. Long-term exposure to Fe and Cu in PM_{2.5} and their combined impact on ROS were consistently associated with increased cardiovascular mortality. Interestingly, adjustments for ROS did not significantly change the effects on cardiovascular diseases and mortality associated with exposure to PM_{2.5}, but the effects associated with exposure to NO₂ decreased significantly after adjustments for ROS, indicating the importance of non-exhaust particulate air

pollution [29]. The results in this study, with a lack of positive excess risks associated with NO_2 , but with significantly positive excess risk for $\text{PM}_{2.5-10}$ during spring time when there is a large amount of road dust, indicate that ROS could have been a contributing factor. However, in the current study, there was no information on ROS in Stockholm.

Among all the air pollutants that were analyzed in this study, the excess risks associated with O_3 were the most robust in terms of health impact, with significantly positive excess risks in 12 out of the 20 cases. Based on the “all year” calculations, the excess risks associated with O_3 were fairly similar in both the single and the multi-pollutant models. However, the results for the different seasons were inconclusive. However, relatively higher excess risks for O_3 during summer and autumn compared to winter and spring were shown in the single-pollutant models. Higher excess risks for O_3 -related mortality during the warm season were also shown in Stockholm in a previous study by Bedada et al. (2016) [23]. Since there are few indoor sources of ozone, an increased exposure during the warmer season can at least partially explain the higher excess risks in the single-pollutant models during the warm seasons.

5. Conclusions

The main objective of this study was to analyze seasonal variations in the excess risks for daily mortality associated with an interquartile range increase in PM_{10} , $\text{PM}_{2.5-10}$, BC, NO_2 , and O_3 in Stockholm during the period from 2000 to 2016. Both single and multi-pollutant models were used in the analysis.

The current study showed a broadly similar pattern for excess risks associated with $\text{PM}_{2.5-10}$ and PM_{10} with larger risk increases in daily mortality during springtime. The seasonal pattern of the excess risks for PM_{10} and $\text{PM}_{2.5-10}$ can possibly be explained by the variation in the chemical composition throughout the year, with a larger amount of road dust present during springtime. The excess risks associated with BC were, in most cases, statistically insignificant, which may be due to the fact that the amount of data on which the calculations were based was smaller in comparison with the other air pollutants.

The calculated excess risks for NO_2 were negative in almost all cases throughout the whole year. Differences in exposure during the year, which previously was hypothesized as an explanation for the negative excess risks, are unlikely. One possible reason for the negative excess risks associated with NO_2 is that the concentrations (on average $14.4 \mu\text{g m}^{-3}$) in this study were too low to cause harmful health effects.

The excess risks associated with O_3 were most robust in terms of the number of statistically significantly positive relationships, indicating that O_3 and its oxidative potential were particularly important in terms of daily mortality associated with air pollution exposure. Higher excess risks were shown during summer and autumn, indicating a higher degree of exposure during the warm seasons.

From a policy point of view, there were clear indications that the health effects associated with exposure to $\text{PM}_{2.5-10}$ and PM_{10} were most evident during the spring. Additional action strategies to reduce the emissions of road dust particles are, therefore, needed.

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Appendix A

Table A1. Numbers and percentage of days with valid data for the measured variables during the period from 2000 to 2016 divided into whole year and four different seasons. Mortality is indicated as cases per day, maximum temperature in °C, and air pollutants in $\mu\text{g m}^{-3}$.

Variable	Whole Year	Winter	Spring	Summer	Autumn
	#Days (% Valid Data)				
Mortality	6210 (100%)	1535 (100%)	1564 (100%)	1564 (100%)	1548 (100%)
Maximum temperature	6210 (100%)	1535 (100%)	1564 (100%)	1564 (100%)	1548 (100%)
PM ₁₀	5999 (97%)	1474 (96%)	1533 (98%)	1500 (96%)	1492 (96%)
PM _{2.5-10}	5352 (86%)	1298 (85%)	1378 (88%)	1356 (87%)	1277 (83%)
BC *	3316 (53%)	764 (50%)	906 (58%)	791 (51%)	855 (55%)
NO ₂	6101 (98%)	1504 (98%)	1533 (98%)	1527 (98%)	1537 (99%)
O ₃	6133 (99%)	1531 (100%)	1553 (99%)	1532 (98%)	1517 (98%)

* No data available for BC before April 2006.

Table A2. Correlation matrix with R-values (Pearson correlation coefficients) between the different air pollutants measured in Stockholm during the whole year during the period from 2000 to 2016.

Pollutant	PM ₁₀	PM _{2.5-10}	BC	NO ₂	O ₃
PM ₁₀	1	0.81	0.49	0.25	0.31
PM _{2.5-10}	0.81	1	0.13	0.12	0.35
BC	0.49	0.13	1	0.49	−0.25
NO ₂	0.25	0.12	0.49	1	−0.46
O ₃	0.31	0.35	−0.25	−0.46	1

Table A3. Correlation matrix with R-values (Pearson correlation coefficients) between the different air pollutants measured in Stockholm during wintertime (December, January and February) during the period from 2000 to 2016.

Pollutant	PM ₁₀	PM _{2.5-10}	BC	NO ₂	O ₃
PM ₁₀	1	0.67	0.56	0.24	−0.02
PM _{2.5-10}	0.67	1	0.05	0.11	0.13
BC	0.56	0.05	1	0.48	−0.45
NO ₂	0.24	0.11	0.48	1	−0.62
O ₃	−0.02	0.13	−0.45	−0.62	1

Table A4. Correlation matrix with R-values (Pearson correlation coefficients) between the different air pollutants measured in Stockholm during springtime (March, April and May) during the period from 2000 to 2016.

Pollutant	PM ₁₀	PM _{2.5-10}	BC	NO ₂	O ₃
PM ₁₀	1	0.83	0.66	0.46	0.18
PM _{2.5-10}	0.83	1	0.31	0.33	0.14
BC	0.66	0.31	1	0.54	−0.06
NO ₂	0.46	0.33	0.54	1	−0.40
O ₃	0.18	0.14	−0.06	−0.40	1

Table A5. Correlation matrix with R-values (Pearson correlation coefficients) between the different air pollutants measured in Stockholm during summertime (June, July and August) during the period from 2000 to 2016.

Pollutant	PM ₁₀	PM _{2.5–10}	BC	NO ₂	O ₃
PM ₁₀	1	0.71	0.55	0.40	0.44
PM _{2.5–10}	0.71	1	0.19	0.16	0.25
BC	0.55	0.19	1	0.55	0.11
NO ₂	0.40	0.16	0.55	1	−0.16
O ₃	0.44	0.25	0.11	−0.16	1

Table A6. Correlation matrix with R-values (Pearson correlation coefficients) between the different air pollutants measured in Stockholm during autumn time (September, October and November) during the period from 2000 to 2016.

Pollutant	PM ₁₀	PM _{2.5–10}	BC	NO ₂	O ₃
PM ₁₀	1	0.69	0.69	0.19	0.06
PM _{2.5–10}	0.69	1	0.32	0.14	0.12
BC	0.69	0.32	1	0.37	−0.30
NO ₂	0.19	0.14	0.37	1	−0.55
O ₃	0.06	0.12	−0.30	−0.55	1

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