Temporal Variation in Air Pollution Concentrations and Preterm Birth—A Population Based Epidemiological Study

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Abstract: There is growing evidence of adverse birth outcomes due to exposure to air pollution during gestation. However, recent negative studies are also reported. The aim of this study was to assess the effect of ozone and vehicle exhaust exposure (NO₂) on the length of the gestational period and risk of preterm delivery. We used data from the Swedish Medical Birth Registry on all vaginally delivered singleton births in the Greater Stockholm area who were conceived during 1987–1995 (n = 115,588). Daily average levels of NO₂ (from three measuring stations) and ozone (two stations) were used to estimate trimester and last week of gestation average exposures. Linear regression models were used to assess the association between the two air pollutants and three exposure windows, while logistic regression models were used when analyzing associations with preterm delivery (<37 weeks gestation). Five percent were born preterm. The median gestational period was 40 weeks. Higher levels of ozone during the first trimester were associated with shorter gestation as well as with an elevated risk of preterm delivery, the odds ratio from the most complex model was 1.06 (95% CI: 1.00–1.13) per 10 μg/m³ increase in the mean daily 8-h maximum concentration. Higher levels of ozone during the second trimester were associated with shorter gestation but the elevated risk of preterm delivery was not statistically significant. Higher levels of ozone and NO₂ during the last week of gestation were associated with a shorter duration of gestation and NO₂ also with preterm delivery. There were no significant associations between first and second trimester
NO₂ exposure estimates and studied outcomes. The effect of first trimester ozone exposure, known to cause oxidative stress, was smallest among women who conceived during autumn when vitamin D status, important for fetal health, in Scandinavian women is the highest.

**Keywords:** pregnancy; preterm birth; vehicle emissions; ozone; nitrogen dioxide

1. Background

There is an ever growing body of evidence that ambient air pollution levels during gestation have adverse effects on birth outcomes [1-12]. Maternal exposure to carbon monoxide (CO), or motor vehicle exhaust, late in the pregnancy was associated with preterm birth in Vancouver, Canada and in California [4-6]. Late pregnancy exposure to fine and respirable particulate matter (PM₂.₅, PM₁₀) has also been associated with an increased risk of preterm delivery in California [5,6]. Early pregnancy exposure to ozone (O₃) and PM₁₀ increased the risk of preterm delivery in Brisbane, Australia [13]. Increased concentration of several pollutants during the first and third trimester were associated with preterm birth in the Republic of Korea [9]. A study from the Czech Republic, where sulphur dioxide levels (SO₂) were rather high, found SO₂ to be associated with preterm birth regardless of period of exposure [12]. Additionally, exposure to particles during early gestation as well as exposure to NOₓ during early or late gestation was associated with preterm birth. It has been suggested that there may be some seasonality in the effects from exposure to air pollution during different exposure windows of gestation on the duration of gestation [10].

However, there is no unambiguous answer as to which specific elements of pollution or during which period of pregnancy, has an impact on pregnancy outcomes. Sram *et al.* conclude that the evidence is strong enough to infer causality of air pollution on low birth weight, LBW (birth weight < 2.5 kg), while the evidence of association between air pollution and preterm birth is deemed insufficient to infer causality [2]. More recently, Bosetti *et al.* reviewed a variety of studies and still found that the studies on maternal exposure to respirable particles during pregnancy do not provide conclusive evidence of an association with the risk of preterm birth and low birth weight [14]. Other authors conclude that for traffic related pollution, the association with an increased risk of preterm birth seems to be consistent [7]. However, several recent studies were negative: in the Dutch PIAMA birth cohort there was no significant association between preterm birth and NO₂ during first trimester, last month or entire pregnancy [15]. Neither was there any significant association between NO₂ and preterm birth in a study from California [3]. From Sydney an unexplained protective effect of NO₂ has been reported [10] and in Valencia the dose-response relationship was V-shaped [16]. The few studies of O₃ have not found robust associations [3,4,10,13]. Preterm birth is a risk factor not only in early life, but is also associated with respiratory and cardiovascular disease later in life [17,18]. Thus more and large studies are needed, especially from Europe where most of the reported studies of air pollution exposure have been small.

Studies of maternal air pollution exposure and preterm birth/LBW build on different types of exposure variables; concentrations at monitoring stations, modelled concentrations or proximity to
sources (roads). The contrasts in exposure variables may be completely spatial or completely temporal or both, but more spatial than temporal [7,8,10].

The aim of this study was to assess the association between exposure to O\textsubscript{3} and duration of gestational and preterm delivery as well as the association with vehicle exhaust, using NO\textsubscript{2} as indicator, during different windows of gestation in the Greater Stockholm area.

2. Population and Methods

All vaginally delivered singleton births by mothers in the Greater Stockholm area who conceived during 1988–1995 were included in the study (n = 115,588). The number of inhabitants in this area was approximately 1.6 million and the area was approximately 3,500 km\textsuperscript{2}, with approximately 40% of the population residing in the City of Stockholm, an area of approximately 200 km\textsuperscript{2}. The information was extracted from the Swedish Medical Birth Registry and included date of birth, birth weight, length at birth, smoking habits of the mother, parity, gestational age and sex of the child. Birth date and information on gestational age were used to determine date of conception and the period for the first (gestational week 1–13) and second (gestational week 14–26) of each trimester of pregnancy. Since the preterm born have a shorter third trimester, we chose not to calculate exposure for this period of varying duration. We instead calculated exposure during the last week at risk of preterm birth. Gestational age was calculated using information on date of last menstruation and from ultrasound examinations.

The air pollution data was provided by the City of Stockholm Environment and Health Administration. We used citywide daily average levels to calculate trimester averages and averages over the last week of pregnancy for NO\textsubscript{2} (daily 1-hour maxima) and O\textsubscript{3} (daily 8-hour maxima). O\textsubscript{3} was measured at one urban background station (Södermalm above roof) and the regional background station (Norr Malma), while NO\textsubscript{2} was measured at three different urban background stations (above roof) within a 1.5 km radius (Södermalm, Hornsgatan, Sveavägen). The daily citywide average was calculated from measured concentrations, or when one station was missing, after imputing values from a linear regression against the other stations as for the APHEA-2 Study [19].

Logistic regression was used to examine if the pollutants influenced the risk for preterm birth (<37 weeks gestation). Both crude models and adjusted models were fitted. In order to improve the quality of the models, they were adjusted for: smoking habits (non-smoker, moderate smoker (1–9 cigarettes/day), heavy smoker (10+ cigarettes/day) and unknown smoking habits), parity (0, 1, 2, 3 or more), and sex. We adjusted for time-varying variables; temperature (°C) and relative humidity (%), as they may confound the study of pollution effects. Additionally we adjusted for seasonal trend by fitting a cyclic regression spline on day of year for conception. Finally, we included a long term trend by fitting a cubic regression spline for conception year. We examined the effects of exposure to O\textsubscript{3} and NO\textsubscript{2} during the first and second trimesters and during the last week at risk of preterm birth. The exposure effects for the different time frames were modelled both individually and simultaneously. We also studied interaction-effects between air pollution and season of conception in the multi-pollutant fully adjusted models. The seasons were defined as; spring (March to May), summer (June to August), autumn (September to November) and winter (December to February).

The effect of the pollution exposures on duration of gestation (as a continuous outcome, in weeks) was assessed using multiple linear regression models, where adjustment was made for the same set of
potential confounders mentioned above. The different exposure windows we studied were; the first and second trimesters of pregnancy, and the last week of pregnancy.

We estimated 95% confidence intervals (CI) for all association parameters (slopes and odds ratios). All analyses were made using R statistical software package (2.13.0) [20].

3. Results

About one half of the mothers were primiparous and about a third gave birth to their second child. 5.3% of the births were preterm. The distribution of duration of gestation was skewed towards the lower end of the spectrum (Figure 1). The variability in birth weight was greater among gestations with longer durations (Figure 2).

![Figure 1. Histogram of duration of gestation.](image)

The average pollution levels and meteorology were relatively similar for different time frames of pregnancy (Table 1). There were moderate negative correlations between O₃ and NO₂ (Table 2), and positive correlations between O₃ and ambient temperature. O₃ and relative humidity were strongly negatively correlated. NO₂ and ambient temperature had a moderate negative correlation. NO₂ and relative humidity was positively correlated. Both O₃ and NO₂ had a slight positive correlation between first and second trimester averages. The correlation between the first trimester and the last week of gestation averages was negative for both pollutants. The correlation between the second trimester O₃ and the last week of gestation O₃ averages was negative. There was no correlation between the second trimester NO₂ and the last week of gestation NO₂ averages.
Figure 2. Duration of gestation plotted against birth weight.

Table 1. Descriptive statistics of air pollutants and meteorological variables.

| Time frame       | Pollutant | Mean $a$ | SD  | Median | IQR $b$ |
|------------------|-----------|----------|-----|--------|---------|
| First trimester  | O$_3$     | 57.1     | 13.3| 55.3   | 20.4    |
| Second trimester | O$_3$     | 56.7     | 13.7| 54.7   | 20.9    |
|                  | Last week $c$ | O$_3$ | 59.0 | 17.5 | 58.4 | 25.3    |
|                  | Last week $d$ | O$_3$ | 58.2 | 17.2 | 57.5 | 25.1    |
| First trimester  | NO$_2$    | 38.5     | 5.4 | 38.3   | 6.0     |
| Second trimester | NO$_2$    | 38.9     | 5.4 | 38.6   | 5.8     |
|                  | Last week $c$ | NO$_2$ | 38.3 | 8.6  | 37.6 | 11.0    |
|                  | Last week $d$ | NO$_2$ | 38.2 | 8.5  | 37.7 | 10.9    |
| First trimester  | Temperature | 7.9   | 6.3 | 7.4    | 12.2    |
| Second trimester | Temperature | 7.3   | 6.2 | 6.3    | 11.4    |
|                  | Last week $c$ | Temperature | 7.7 | 7.2 | 7.1 | 12.4    |
|                  | Last week $d$ | Temperature | 7.4 | 7.2 | 6.4 | 11.9    |
| First trimester  | Humidity   | 69.9     | 9.3 | 70.9   | 15.2    |
| Second trimester | Humidity   | 70.5     | 9.1 | 71.8   | 14.7    |
|                  | Last week $c$ | Humidity | 70.0 | 11.9 | 71.1 | 18.2    |
|                  | Last week $d$ | Humidity | 69.9 | 11.7 | 71.1 | 18.0    |

$^a$ Units are $\mu g/m^3$ for the air pollutants, $^\circ C$ for temperature and $\%$ for relative humidity. $^b$ IQR is equal to the difference between the 75th and 25th percentiles. $^c$ Last week of gestation. $^d$ Last week at risk of preterm birth.
The proportion of preterm births was higher among smoking mothers (Table 3). The proportion of preterm births was higher in the subset of women who did not have any information on smoking than in the other subsets. The proportion of preterm births was higher among primiparous women than for women giving birth for the second time. Among women giving birth to their third or subsequent child the proportion of preterm births was higher than among primiparous women or women giving birth to their second child. There was only a minor difference between boys and girls in the proportion of preterm birth.

There was no clear seasonality in how preterm births were distributed over the year, there was however a slightly elevated proportion of preterm births among the mothers who conceived during Spring or Autumn (Table 3). The proportion of preterm births was somewhat lower during 1989 and 1990, the two years with the lowest first trimester O₃ averages, compared to the other years, and the proportion of preterm births was slightly elevated in 1994, when first trimester O₃ levels were the highest in this study.

### Table 2. Correlation between O₃, NO₂ and meteorological averages.

|                         | First trimester | Second trimester | Last week a | Last week b | First trimester | Second trimester | Last week a | Last week b |
|-------------------------|-----------------|------------------|-------------|-------------|-----------------|------------------|-------------|-------------|
| Second trimester O₃     | 0.13            |                  |             |             |                 |                  |             |             |
| Last week a O₃          | -0.34           | -0.45            |             |             |                 |                  |             |             |
| Last week b O₃          | -0.50           | -0.32            | 0.71        |             |                 |                  |             |             |
| First trimester NO₂     | -0.43           | 0.16             | -0.03       | 0.11        |                 |                  |             |             |
| Second trimester NO₂    | -0.33           | -0.39            | 0.30        | 0.30        | 0.25            |                  |             |             |
| Last week a NO₂         | 0.29            | 0.11             | -0.26       | -0.34       | -0.09           | 0.03             |             |             |
| Last week b NO₂         | 0.29            | 0.00             | -0.18       | -0.26       | -0.07           | 0.03             | 0.48        |             |
| First trimester temperature | 0.62          | -0.61            | 0.06        | -0.18       | -0.53           | 0.09             | 0.20        | 0.28        |
| Second trimester temperature | 0.69          | 0.56             | -0.73       | -0.77       | -0.10           | -0.50            | 0.37        | 0.33        |
| Last week a temperature | -0.79           | -0.08            | 0.54        | 0.61        | 0.44            | 0.36             | -0.40       | -0.39       |
| Last week b temperature | -0.78           | 0.17             | 0.27        | 0.54        | 0.49            | 0.27             | -0.32       | -0.40       |
| First trimester relative humidity | -0.82          | 0.24             | 0.26        | 0.46        | 0.42            | 0.19             | -0.20       | -0.25       |
| Second trimester relative humidity | -0.30          | -0.79            | 0.69        | 0.61        | -0.11           | 0.40             | -0.28       | -0.19       |
| Last week a relative humidity | 0.55           | 0.41             | -0.71       | -0.62       | -0.30           | -0.36            | 0.22        | 0.20        |
| Last week b relative humidity | 0.65           | 0.23             | -0.49       | -0.70       | -0.41           | -0.30            | 0.28        | 0.23        |

a Last week of gestation. b Last week at risk of preterm birth.
There was a clear association between O₃ levels during the first trimester and preterm delivery, both in the unadjusted and adjusted models (Table 4). The association was stronger in the multiple pollutant models than in the crude model. The Odds Ratio, OR, was equal to 1.08 per a 10 µg/m³ increase in O₃ in the individual exposure window model, CI = (1.03, 1.13). In the multiple exposure window model the OR was 1.06 per 10 µg/m³ increase in O₃, CI = (1.00, 1.13), and the crude OR was 1.03 per 10 µg/m³ increase in O₃, CI = (1.01, 1.05). Higher second trimester O₃ was positively associated with preterm birth, although the 95% confidence intervals included the null in all models. We did not observe any clear association between elevated O₃ levels towards the end of gestation and preterm birth. First and second trimester NO₂ levels were non-significant with OR < 1. Higher levels of NO₂ during the last week at risk of preterm birth was positively associated with preterm birth in all models.
Table 4. Crude and adjusted ORs for preterm delivery per 10 µg/m³ increase in pollutant concentration.

| Time frame   | Pollutant | Crude OR (95% CI) | ORs and (95% CI) from multiple pollutant model a. Exposure windows studied individually | ORs and (95% CI) from multiple pollutant model a. Exposure windows studied simultaneously |
|--------------|-----------|-------------------|---------------------------------------------------------------------------------------|---------------------------------------------------------------------------------------|
| First trimester | O₃       | 1.03 (1.01, 1.05) | 1.08 (1.03, 1.13)                                                                     | 1.06 (1.00, 1.13)                                                                     |
| Second trimester | O₃       | 1.01 (0.99, 1.03) | 1.02 (0.97, 1.07)                                                                     | 1.05 (0.98, 1.12)                                                                     |
| Last week     | O₃       | 0.99 (0.98, 1.01) | 1.01 (0.98, 1.04)                                                                     | 1.02 (0.98, 1.05)                                                                     |
| First trimester | NO₂      | 0.96 (0.92, 1.01) | 0.95 (0.88, 1.03)                                                                     | 0.95 (0.87, 1.04)                                                                     |
| Second trimester | NO₂      | 0.95 (0.91, 1.00) | 0.96 (0.88, 1.04)                                                                     | 0.93 (0.85, 1.02)                                                                     |
| Last week     | NO₂      | 1.05 (1.02, 1.08) | 1.07 (1.03, 1.11)                                                                     | 1.06 (1.02, 1.11)                                                                     |

a Adjusted for maternal smoking, parity, sex of the child, temperature, relative humidity, seasonal variation and long term trend.

Table 5. Crude and adjusted slope estimates for duration of gestation (in weeks) per 10 µg/m³ increase in pollutant concentration.

| Time frame   | Pollutant | Crude slope estimate (95% CI) | Slope estimates and (95% CI) from multiple pollutant model a. Exposure windows studied individually | Slope estimates and (95% CI) from multiple pollutant model a. Exposure windows studied simultaneously |
|--------------|-----------|-------------------------------|---------------------------------------------------------------------------------------------------|---------------------------------------------------------------------------------------------------|
| First trimester | O₃       | −0.02 (−0.03, −0.01)          | −0.06 (−0.08, −0.04)                                                                                 | −0.05 (−0.08, −0.02)                                                                                 |
| Second trimester | O₃       | −0.01 (−0.02, −0.01)          | −0.03 (−0.05, −0.01)                                                                                 | −0.07 (−0.10, −0.04)                                                                                 |
| Last week     | O₃       | 0.01 (0.00, 0.01)             | −0.02 (−0.03, −0.01)                                                                                 | −0.02 (−0.03, −0.01)                                                                                 |
| First trimester | NO₂      | 0.04 (0.02, 0.06)             | 0.04 (0.00, 0.07)                                                                                    | 0.01 (−0.03, 0.05)                                                                                  |
| Second trimester | NO₂      | 0.05 (0.03, 0.07)             | 0.02 (−0.01, 0.06)                                                                                  | 0.04 (−0.01, 0.08)                                                                                  |
| Last week     | NO₂      | −0.03, (−0.04, −0.02)         | −0.04 (−0.05, −0.02)                                                                                  | −0.04 (−0.06, −0.02)                                                                                  |

a Adjusted for maternal smoking, parity, sex of the child, temperature, relative humidity, seasonal variation and long term trend.

The associations were very similar in the analyses of gestational period. In the fully adjusted model high O₃ levels were associated with a shorter duration of gestation, regardless of window of exposure (Table 5). The strongest association, in the multiple pollutant—multiple exposure windows model, was observed for second trimester O₃ and duration of gestation. A 10 µg/m³ increase in O₃ was associated with a 0.07 weeks reduction of the duration of gestation, CI = (−0.10, −0.04). Higher first and second trimester NO₂ levels had a tendency to be positively associated with gestation, but the 95% confidence intervals for the slope estimates excluded the null only for the crude models. Exposure to high levels of O₃ and NO₂ during the last week of gestation was associated with a shorter duration of gestation.

The observed association between first trimester O₃ and preterm birth was modified by season, the ORs for preterm delivery ranged from 1.07, CI = (0.98, 1.16), during autumn to 1.13, CI = (1.05, 1.21), during spring per 10 µg/m³ increase in O₃ during the first trimester (Table 6). The weakest association
for first trimester exposure to O₃ was for those pregnancies where the conception day happened during autumn, and this was the only season-specific O₃ effect where the confidence interval included the null.

| Season  | Crude OR (95% CI) | Adjusted OR (95% CI) a | Crude slope estimates (95% CI) | Adjusted slope estimates (95% CI) a |
|---------|-------------------|-------------------------|--------------------------------|-----------------------------------|
| Spring  | 1.10 (1.02, 1.18) | 1.13 (1.05, 1.21)       | −0.06 (−0.09, −0.03)           | −0.07 (−0.10, −0.04)              |
| Summer  | 1.05 (0.99, 1.10) | 1.13 (1.02, 1.26)       | −0.03 (−0.06, 0.00)            | −0.04 (−0.07, −0.02)              |
| Autumn  | 1.05 (0.96, 1.14) | 1.07 (0.98, 1.16)       | −0.06 (−0.09, −0.02)           | −0.06 (−0.09, −0.03)              |
| Winter  | 1.15 (1.08, 1.21) | 1.12 (1.04, 1.20)       | −0.07 (−0.10, −0.04)           | −0.06 (−0.09, −0.03)              |

a Multiple pollutant model adjusted for maternal smoking, parity, sex of the child, first trimester temperature, first trimester relative humidity, first trimester NO₂, a seasonal spline and long term cubic regression spline.

High exposure to O₃ during the first trimester was associated with a shorter duration of gestation, regardless of season of conception, although the association was weakest during summer (Table 5).

4. Discussion

We found an association between first trimester O₃ exposure and increased risk of preterm birth, which was consistent with descriptive data for the different years. We have studied the effects of temporal fluctuations in air pollution exposure between pregnancies. This approach gives us the ability to detect effects better for exposures that vary more strongly in time than in space. That is, this study is well suited to detect effects of O₃, but not so well suited to detect the full effects from air pollutants with an additional within city variation, i.e., NO₂. At this latitude, O₃ levels are mainly dependent on the incoming air masses, and the local formation is very limited [21]. At the same time, the regional background concentrations of pollutants such as NO₂ are low, and the fluctuations in concentrations are highly dependent on the dispersion of emissions from local traffic in the city. This means that periods of stagnation will increase exhaust levels everywhere in the city, though more in some streets and districts than in others. As such, there will have been some misclassification of NO₂ exposures, which as non-differential typically would tend to produce attention in slopes and odds ratios. We use a few stations only for a quite large area. However, air pollution data from the same monitoring stations have for this population shown significant associations in time-series studies of daily mortality and hospitalisations in APHEA-2 and other studies, indicating their validity [22].

The Medical Birth Registry has existed in Sweden since 1973, is almost complete (99%) and of very high quality [23]. Because of this it is very unlikely that missing data has had any impact on the results of this study. One quality issue we observed with the Medical Birth Registry was the lack of information on smoking habits. In 7.4% of the births in this study we did not have information on smoking status. Possible explanations to these missing records are; unwillingness to respond to this question during the first antenatal care visit, failure to report information from antenatal care by the birthing centre or antenatal care started in a later stage of pregnancy. This is probably more common among women with social problems. There is of course the issue of migration. Exposure misclassification will occur for women who moved to the study area during gestation. However, although such movements may bias associations towards the null, we do not believe these migration
patterns could confound the associations driven by studying temporal fluctuations in air pollution concentrations. The only criterion for a child to be entered into the study was to be a vaginally delivered singleton birth during the study period; this means that we have no restriction as to how many times one woman is represented. A consequence of this is that the observations are not independent, and there may be some individual factors that strongly predicts duration of gestation, for example some genetic predisposition. However, the direction and magnitude of the associations did not change when restricting the models to primiparous women only (Appendix A).

A typical problem in air pollution studies is potential confounding, and in studies of birth outcome socioeconomic status, lifestyle, country of origin etc. may differ between areas with more or less air pollution. As we only study the influence of temporal variability in the air pollutants, we primarily need to worry about potential confounding for variables that vary in time in a similar way [11,24]. We did not observe any confounding effect on the results for NO2. We observed some confounding from the meteorological factors and time of conception on the results for first trimester O3, inflating the ORs from 1.06 per IQR increase in the unadjusted model to 1.16 per IQR increase in the adjusted model. Season of conception, temperature and relative humidity could be good proxies for sunlight exposure, which is correlated with ground level O3 formation. Sunlight is associated with maternal vitamin D status. Maternal vitamin D levels are important for fetal health, and vitamin D deficiency during pregnancy has been associated with preeclampsia [25]. Inflammation in early pregnancy is associated with an increased risk of preterm delivery [26]. Vitamin D deficiency is associated with increased circulating inflammatory proteins, and vitamin D status seems to modulate C-reactive protein levels [27]. In a recent study exposures to PM and O3, but not NO2, were associated with increased C-reactive protein concentrations in early pregnancy [28]. Vitamin D status has a clear seasonal variation in Scandinavian women, and is best in the autumn [29]. In this study the effect of first trimester O3 levels was smallest among women who conceived during the autumn.

One of the limitations in this study is the lack of adjustment for socioeconomic factors. There are no good proxies for socioeconomic status in the Medical Birth Registry. Socioeconomic status varies more strongly in space than in time, at least in a shorter time span. It might be plausible that there is confounding from temporal variations in socioeconomic status on the associations between air pollution and preterm delivery. Most of this confounding should be accounted for by adjusting for time, as both the variation of air pollution and the variation of socioeconomic status are functions of time. Although the risk of confounding from socioeconomic status is limited, the overall performance of the models would likely improve by adjusting for socioeconomic status.

Our finding of an early pregnancy effect of O3 is similar, albeit smaller, than the findings of Hansen et al. [13]. Liu et al. found an OR of 1.08 for a 10 ppb increase in first month O3 in their unadjusted model, but this effect was reversed, OR = 0.98 per 10 ppb increase, when adjusting for possible confounders; maternal age, parity, infant sex, birth weight and season of birth [4]. The first finding is similar to the first trimester crude effect in this study, but the results diverge in the adjusted models. The difference in results may be due to difference in how seasons were defined, Liu et al. adjusted for season of birth (which may be modified by preterm birth) while we adjusted for season of conception. No associations between early gestation exposure to O3 were found in the two studies from the southern California [3,5]. The inconsistent findings could be caused by differences in susceptibility (e.g., different anti-oxidant status in different populations), differences in exposure patterns (e.g., high
prevalence of air conditioning reduces O₃ exposure) and differences in the pollution mix (e.g., negative correlation with primary combustion components).

There could be two explanations to the overall null effect of early and mid-gestation NO₂ concentrations. One explanation is that there is no effect; the other is that we fail to catch important variability in exhaust exposure during the first and second trimester as we were unable to take into account spatial variation in exposure. However, in the Californian studies there was spatial resolution in the exposure data, still they found no or weak indications of an adverse effect of early gestation exposure to higher ambient NO₂ levels [3,5]. The observation of an adverse association of exposure to higher NO₂ levels during the last period of gestation has been observed in several studies [3-5,9,13]. There have also been observations of a null effect of exposure to higher NO₂ levels during the last part of pregnancy [10]. The correlations between NO₂ and other pollutants, such as particles, may differ between the different study locations.

All season-specific estimates of exposure to higher O₃ levels during the first trimester were associated with an elevated risk of preterm delivery and a shorter duration of gestation, although the association with preterm birth was slightly weaker for conceptions in autumn than for conceptions during any other season. This strengthens our belief that there seems to be a negative effect of early gestation exposure to O₃ on the duration of the gestational period. The season-specific findings are not in accordance with the findings of Jalaludin et al., who found statistically significant positive associations with O₃ levels during the first trimester for women who conceived during spring or summer [10]. The association with O₃ levels we found is robust in that we observe the effect regardless of whether we study duration of gestation or preterm delivery.

The pathway by which we believe O₃ may cause preterm delivery is through oxidative stress. Oxidative stress caused by exposure to air pollution may lead to inflammatory response in the respiratory system [30]. Expression of proinflammatory cytokines could disrupt trophoblast invasion during the formation of the placenta [13,31]. This could lead to preeclampsia, which is an important predictor of preterm birth. Alternatively systemic oxidative stress may affect the embryo at the earliest phase of development, which may cause intra-uterine growth retardation or preterm birth [32]. Exposure to air pollution in a later stage of gestation may cause disturbances in cord blood flow or increased susceptibility for maternal infections, which may cause premature rupture of membranes leading to an earlier birth [5].

We have shown that although the air pollution level is moderate in Sweden there is an association between exposure to air pollution during pregnancy and reduction of gestational age. Our results support the observation that there is some seasonality in the association with air pollution. Further studies are needed to find out more about these associations and possible mechanisms.

How important are these effects of moderate air pollution from a public health point of view? A comparison with smoking shows that the impact of air pollution must be taken seriously. 6.5% of the mothers (7,520) were heavy smokers with a prevalence of preterm birth of 6.76% compared to 4.84% preterm among non-smoking mothers. This corresponds to 144 preterm births attributed to heavy smoking. The estimated attributable fraction preterm births associated with heavy smoking was 2.1%. 10.6% of the mothers were moderate smokers with a prevalence of preterm birth of 5.36%, and 64 preterm births attributed to moderate smoking. The adjusted ORs (heavy smoking 1.37, moderate smoking 1.12) would give similar numbers. The upper quartile of first trimester O₃ exposure has a mean value of 27.7 µg/m³ over the lowest quartile, which corresponds to an OR of 1.18. The fraction
of preterm births attributed to the upper quartile of first trimester O₃ exposure was 7.8% (corresponding to 129 preterm births) for this exposure category. The upper quartile of last week NO₂ exposure has a mean value of 16.9 µg/m³ over the lowest quartile, which corresponds to an OR of 1.11. The estimated fraction of preterm births attributable to the highest quartile of late pregnancy NO₂ exposure was 5.2% (corresponding to 85 preterm births) for this exposure category.

Heavy smokers had a reduction in duration of gestation estimated to approximately 0.19 [95% CI = (−0.23, −0.14)] weeks, while moderate smokers had no reduction. The upper quartile of first trimester O₃ exposure has a mean value of 27.7 µg/m³ over the lowest quartile, which corresponds to 0.13 [95% CI = (−0.21, −0.05)] weeks shorter gestation (using the adjusted slope). However, this is a mean value, but the real effect is likely not evenly distributed. The average reduction in duration of gestation for women exposed to levels within the highest quartile of last week NO₂ compared to the lowest quartile was 0.068 [95% CI = (−0.096, −0.040)] weeks.

5. Conclusions

The results found in this study indicate that there is an early (first trimester) effect of O₃ in accordance with suggested pathways for the early exposure window. The effect of O₃ was smallest among women who conceived during the autumn when the vitamin D levels in Scandinavian women should be the highest. We observed an adverse effect of exposure to elevated NO₂ levels in late gestation on preterm delivery. However, no spatial variation in NO₂ was modelled. There remains a need to study these associations further with subject specific exposure assessments. However, the total size of these effects supports improvements in air quality also in a moderately polluted region.

Conflict of Interest

The authors declare no conflict of interest.

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References

1. Maisonet, M.; Correa, A.; Misra, D.; Jaakkola, J.J.K. A review of the literature on the effects of ambient air pollution on fetal growth. Environ. Res. 2004, 95, 106-115.
2. Srám, R.J.; Binková, B.; Dejmek, J.; Bobak, M. Ambient air pollution and pregnancy outcomes: A review of the literature. Environ. Health Perspect. 2005, 113, 375-382.
3. Ritz, B.; Wilhelm, M.; Hoggatt, K.J.; Ghosh, J.K.C. Ambient air pollution and preterm birth in the environment and pregnancy outcomes study at the University of California, Los Angeles. Am. J. Epidemiol. 2007, 166, 1045-1052.
4. Liu, S.; Krewski, D.; Shi, Y.; Chen, Y.; Burnett, R.T. Association between gaseous ambient air pollutants and adverse pregnancy outcomes in Vancouver, Canada. *Environ. Health Perspect.* **2003**, *111*, 1773-1778.

5. Ritz, B.; Yu, F.; Chapa, G.; Fruin, S. Effect of air pollution on preterm birth among children born in Southern California between 1989 and 1993. *Epidemiology* **2000**, *11*, 502-511.

6. Wilhelm, M.; Ritz, B. Local variations in CO and particulate air pollution and adverse birth outcomes in Los Angeles County, California, USA. *Environ. Health Perspect.* **2005**, *113*, 1212-1221.

7. Wu, J.; Ren, C.; Delfino, R.J.; Chung, J.; Wilhelm, M.; Ritz, B. Association between local traffic-generated air pollution and preeclampsia and preterm delivery in the south coast air basin of California. *Environ. Health Perspect.* **2009**, *117*, 1773-1779.

8. Landgren, O. Environmental pollution and delivery outcome in Southern Sweden: A study with central registries. *Acta Paediatr.* **1996**, *85*, 1361-1364.

9. Leem, J.-H.; Kaplan, B.M.; Shim, Y.K.; Pohl, H.R.; Gotway, C.A.; Bullard, S.M.; Rogers, J.F.; Smith, M.M.; Tylenda, C.A. Exposures to air pollutants during pregnancy and preterm delivery. *Environ. Health Perspect.* **2006**, *114*, 905-910.

10. Jalaludin, B.; Mannes, T.; Morgan, G.; Lincoln, D.; Sheppeard, V.; Corbett, S. Impact of ambient air pollution on gestational age is modified by season in Sydney, Australia. *Environ. Health* **2007**, *6*, doi:10.1186/1476-069X-6-16.

11. Slama, R.; Darrow, L.; Parker, J.; Woodruff, T.J.; Strickland, M.; Nieuwenhuijsen, M.; Glinianaia, S.; Hoggatt, K.J.; Kannan, S.; Hurley, F.; *et al.* Meeting report: Atmospheric pollution and human reproduction. *Environ. Health Perspect.* **2008**, *116*, 791-798.

12. Bobak, M. Outdoor air pollution, low birth weight, and prematurity. *Environ. Health Perspect.* **2000**, *108*, 173-176.

13. Hansen, C.; Neller, A.; Williams, G.; Simpson, R. Maternal exposure to low levels of ambient air pollution and preterm birth in Brisbane, Australia. *BJOG* **2006**, *113*, 935-941.

14. Bosetti, C.; Nieuwenhuijsen, M.J.; Gallus, S.; Cipriani, S.; La Vecchia, C.; Parazzini, F. Ambient particulate matter and preterm birth or birth weight: A review of the literature. *Arch. Toxicol.* **2010**, *84*, 447-460.

15. Gehring, U.; Wijga, A.H.; Fischer, P.; de Jongste, J.C.; Kerkhof, M.; Koppelman, G.H.; Smit, H.A.; Brunekreef, B. Traffic-related air pollution, preterm birth and term birth weight in the PIAMA birth cohort study. *Environ. Res.* **2011**, *111*, 125-135.

16. Llop, S.; Ballester, F.; Estarlich, M.; Esplugues, A.; Rebagliato, M.; Iniguez, C. Preterm birth and exposure to air pollutants during pregnancy. *Environ. Res.* **2010**, *110*, 778-785.

17. Kwinta, P.; Pietrzyk, J.J. Preterm birth and respiratory disease in later life. *Expert. Rev. Respir. Med.* **2010**, *4*, 593-604.

18. Skilton, M.R.; Viikari, J.S.; Juonala, M.; Laitinen, T.; Lehtimaki, T.; Taattonen, L.; Kahonen, M.; Celermajer, D.S.; Raitakari, O.T. Fetal growth and preterm birth influence cardiovascular risk factors and arterial health in young adults: The cardiovascular risk in young Finns study. *Arterioscler Thromb Vasc. Biol.* **2011**, *31*, 2975-2981.
19. Katsouyanni, K.; Schwartz, J.; Spix, C.; Touloumi, G.; Zmirou, D.; Zanobetti, A.; Wojtyniak, B.; Vonk, J.M.; Tobias, A.; Ponka, A.; et al. Short term effects of air pollution on health: A European approach using epidemiologic time series data: The APHEA protocol. J. Epidemiol. Community Health 1996, 50(Suppl 1), S12-S18.

20. R Development Core Team. R: A Language and Environment for Statistical Computing; R Foundation for Statistical Computing: Vienna, Austria, 2011. Available online: http://www.r-project.org (accessed on 18 April 2011).

21. Andersson-Sköld, Y.; Janhäll, S. Simulations of ozone formation from different emission sources in Sweden. Water Air Soil Pollut. 1995, 85, 2045-2050.

22. Gryparis, A.; Forsberg, B.; Katsouyanni, K.; Analitis, A.; Touloumi, G.; Schwartz, J.; Samoli, E.; Medina, S.; Anderson, H.R.; Niciu, E.M.; et al. Acute effects of ozone on mortality from the “air pollution and health: A European approach” project. Am. J. Respir. Crit. Care Med. 2004, 170, 1080-1087.

23. Cnattingius, S.; Ericson, A.; Gunnarskog, J.; Kallen, B. A quality study of a medical birth registry. Scand. J. Soc. Med. 1990, 18, 143-148.

24. Strickland, M.J.; Klein, M.; Darrow, L.A.; Flanders, W.D.; Correa, A.; Marcus, M.; Tolbert, P.E. The issue of confounding in epidemiological studies of ambient air pollution and pregnancy outcomes. J. Epidemiol. Community Health 2009, 63, 500-504.

25. Shin, J.S.; Choi, M.Y.; Longtine, M.S.; Nelson, D.M. Vitamin D effects on pregnancy and the placenta. Placenta 2010, 31, 1027-1034.

26. Catov, J.M.; Bodnar, L.M.; Ness, R.B.; Barron, S.J.; Roberts, J.M. Inflammation and dyslipidemia related to risk of spontaneous preterm birth. Am. J. Epidemiol. 2007, 166, 1312-1319.

27. Ngo, D.T.; Sverdlov, A.L.; McNeil, J.J.; Horowitz, J.D. Does vitamin D modulate asymmetric dimethylarginine and C-reactive protein concentrations? Am. J. Med. 2010, 123, 335-341.

28. Lee, P.C.; Talbott, E.O.; Roberts, J.M.; Catov, J.M.; Sharma, R.K.; Ritz, B. Particulate air pollution exposure and C-reactive protein during early pregnancy. Epidemiology 2011, 22, 524-531.

29. Moosgaard, B.; Vestergaard, P.; Heickendorff, L.; Melsen, F.; Christiansen, P.; Mosekilde, L. Vitamin D status, seasonal variations, parathyroid adenoma weight and bone mineral density in primary hyperparathyroidism. Clin. Endocrinol. (Oxf.) 2005, 63, 506-513.

30. Donaldson, K.; Stone, V.; Seaton, A.; MacNee, W. Ambient particle inhalation and the cardiovascular system: Potential mechanisms. Environ. Health Perspect. 2001, 109(Suppl 4), 523-527.

31. Gitto, E.; Reiter, R.J.; Karbownik, M.; Tan, D.-X.; Gitto, P.; Barberi, S.; Barberi, I. Causes of oxidative stress in the pre- and perinatal period. Biol. Neonate 2002, 81, 146-157.

32. Kannan, S.; Misra, D.P.; Dvorch, J.T.; Krishnakumar, A. Exposures to airborne particulate matter and adverse perinatal outcomes: A biologically plausible mechanistic framework for exploring potential effect modification by nutrition. Environ. Health Perspect. 2006, 114, 1636-1642.

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