Exposure to traffic and lung function in adults: a general population cohort study

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ABSTRACT

Objectives: To investigate the association between living near dense traffic and lung function in a cohort of adults from a single urban region.

Design: Cross-sectional results from a cohort study.

Setting: The adult-onset asthma and exhaled nitric oxide (ADONIX) cohort, sampled during 2001–2008 in Gothenburg, Sweden. Exposure was assessed as the distance from participants’ residential address to the nearest road with dense traffic (>10 000 vehicles per day) or very dense traffic (>30 000 vehicles per day). The exposure categories were: low (>500 m; reference), medium (75–500 m) or high (<75 m).

Participants: The source population was a population-based cohort of adults (n=6153). The study population included 5441 participants of European descent with good quality spirometry and information about all outcomes and covariates.

Outcome measures: Forced vital capacity (FVC) and forced expiratory volume in 1 s (FEV1) were measured at a clinical examination. The association with exposure was examined using linear regression adjusting for age, gender, body mass index, smoking status and education in all participants and stratified by sex, smoking status and respiratory health status.

Results: We identified a significant dose–response trend between exposure category and FEV1 (p=0.03) and borderline significant trend for FVC (p=0.06) after adjusting for covariates. High exposure was associated with lower FEV1 (−1.0%, 95% CI −2.5% to 0.5%) and lower FVC (−0.9%, 95% CI −2.2% to 0.4%). The effect appeared to be stronger in women. In highly exposed individuals with current asthma or chronic obstructive pulmonary disease, FVC was lower (−4.5%, 95% CI −8.8% to −0.1%).

Conclusions: High traffic exposure at the residential address was associated with lower than predicted FEV1 and FVC lung function compared with living further away in a large general population cohort. There were particular effects on women and individuals with obstructive disease.

BACKGROUND

Vehicular traffic is a major contributor to air pollution in urban areas, and the emissions contain a plethora of substances, including nitrogen oxides and fine particles, which have acute negative effects on respiratory health.1 2 There is less evidence for the long-term effects of air pollution on respiratory health, including effects on lung function.3 Some studies have shown traffic-related air pollution associated with attenuated lung development in children, specifically forced expiratory volume in 1 s (FEV1) between ages 10 and 18,4 and modelled traffic exposure during the first year of life was associated with lower FEV1 at age 8.5

In adults, there are some studies showing that exposure to traffic pollution is related to reductions in lung function.6–10 However, none of the previous studies concern adults from a random population sample living within a single urban region.

In the earlier studies of long-term effects, crude exposure assessment with low spatial resolution contributed to exposure misclassification. Long-term individual exposure is influenced by a number of factors, including regional and nearby pollution sources,11 and is modified by individual factors, for example, occupational and lifestyle exposures.4

Geographic metrics are straightforward methods for classifying exposure as they increase the spatial resolution and reduce exposure misclassification compared to emission models or ground level measurements.
The levels of pollutants from vehicle exhaust decay to near-background level as the distance to a road increases beyond some hundreds of metres from the site of generation. When studying a mixture of substances such as vehicle emissions, a general exposure indicator such as the distance to a large road may be more useful than models of specific pollutants, as this approach has been used in a number of studies.\textsuperscript{13–16} The aim of this study was to investigate the association between residential proximity to dense and very dense traffic and lung function in adults within a single coherent urban region, using cross-sectional data from a large population-based cohort.

**MATERIALS AND METHODS**

The study population was based on the adult-onset asthma and exhaled nitric oxide (ADONIX) cohort, which consists of a random sample of adults (25–75 years old) in Gothenburg. In total, 14,554 participants were invited to a clinical examination with spirometry, anthropometric measurements and a clinical examination. Information about age, respiratory health, education, asthma status, smoking history and level of education was collected as described by Olin et al.\textsuperscript{17} Spirometry was performed with a dry wedge spirometer (Vitalograph; Buckingham, UK) except in 164 participants who were measured using EasyOne, and percentages of predicted values of lung function variables (FEV\textsubscript{1}, FVC and FEV\textsubscript{1}/FVC ratio) were calculated.\textsuperscript{18}

**Variable definitions**

Age was reported in whole years, and body mass index (BMI) as kg/m\textsuperscript{2}. Smoking was coded into three categories: never, former and current smokers, where former smokers were those who had stopped smoking at least a year ago. Obstructive lung disease was defined as reporting asthma in the previous 12 months, or fulfilling criteria for chronic obstructive pulmonary disease (COPD) as assessed from spirometry without a reversibility test according to the Global Initiative for Chronic Obstructive Lung Disease criteria\textsuperscript{19} in a regular spirometry test without reversibility testing. Education level was reported in one of six categories (primary, upper primary, vocational, high school, university equivalent or other) and used as an indicator of socioeconomic status.

**Residential traffic exposure**

Modelled values of NO\textsubscript{2} and NO\textsubscript{x} were only available for Residential traffic exposure other) and used as an indicator of socioeconomic status.

A primary, vocational, high school, university equivalent or reported in one of six categories (primary, upper tertile for chronic obstructive pulmonary disease (COPD) as assessed from spirometry without reversibility testing. Education level was reported in one of six categories (primary, upper primary, vocational, high school, university equivalent or other) and used as an indicator of socioeconomic status.

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**Residential traffic exposure**

Modelled values of NO\textsubscript{2} and NO\textsubscript{x} were only available for some of the study participants (60% and 77%); instead, the distance from the participants’ home address to the nearest densely trafficked road was used as a measure of exposure to traffic pollution (the distance to road with 10,000 cars and modelled 2006 values of NO\textsubscript{2} and NO\textsubscript{x} have a correlation coefficient of −0.57 and −0.55 respectively). All participants’ residential addresses at the time of examination were geocoded, and the distance from each address to the nearest road with more than 10,000 vehicles per day (‘dense traffic’) and more than 30,000 vehicles per day (‘very dense traffic’) as an annual average for the year 2004 was calculated by the Gothenburg municipality office.

The distances to the nearest road were coded into three categories: closer than 75 m (high exposure), between 75 and 500 m (medium exposure), and 500 m or more (low exposure). The cut-offs were chosen based on prior data regarding the dispersion of air pollutants after they are emitted from the source,\textsuperscript{12} that is, an almost exponential increase in the proximity of a road, as well as the distribution of the data and available literature.\textsuperscript{20–23} The study population was then classified into these three exposure categories in combination with the density of road traffic (either 10,000 or 30,000 vehicles per day). Analyses were performed separately for the two levels of traffic density.

Stratified analyses were performed for subgroups based on gender, smoking status and respiratory health status (healthy vs self-reported current asthma or COPD assessed from spirometry).

**Statistical methods**

After excluding participants with missing data, we explored the association between categories of exposure and FVC and FEV\textsubscript{1}, with linear regression using the low exposure category as reference. Initially, the outcomes were regressed on exposure and each potential confounding variable individually. Variables which were statistically significantly associated with at least one of the outcomes and also changed the estimated effect by 10% were included in the final models. For the main analysis, both unadjusted results and results adjusted for age, gender, BMI, education and smoking status are presented. The data were analysed stratified by gender, smoking status and respiratory health status. The percentage of lorry traffic was added to the model in a sensitivity analysis.

The results are reported as change in percent predicted FEV\textsubscript{1} or FVC, with a 95% CI. We report p for trend, which was obtained through transforming the exposure categories to linear variables with values 0, 1 and 2 for each category, and including them in the same models instead of the category indicator variables. Results were considered statistically significant at the 0.05 level. R V.2.14.1 and STATA were used for all analyses.

**RESULTS**

In total, 6686 participants were clinically examined once during 2001–2003 or 2005–2008, as previously described.\textsuperscript{18} The participation rate for the whole cohort was 46%. Of the total 6686 individuals in the cohort, 5441 participants (81%) lived within the greater Gothenburg area, had data available on all covariates, were of European descent and could be included in the analysis. The skewness of the distribution of % predicted FVC was −0.123, rather symmetric. The skewness of % predicted FEV\textsubscript{1} was −0.483, within the range of being approximately symmetric.
In all, 363 participants were highly exposed to dense traffic (<75 m), 2668 had medium exposure (75–500 m) and 2410 had low exposure (>500 m). Demographic characteristics were similar among the participants regardless of exposure category, except for university education which was more common in the high exposure group (table 1).

The crude effect estimate suggested a 1% reduction in predicted FEV1 and FVC among people living within 75 m of a densely trafficked road in comparison to those living more than 500 m away from a similar traffic flow (table 2). The associations followed a dose–response relationship with a near-significant trend for FEV1. In the adjusted analysis, the associations were strengthened with the medium exposure category estimate for FEV1 becoming statistically significant, and the dose–response trend significant for FEV1 and near-significant for FVC (table 2).

Among the covariates, current smoking, BMI, age and male gender were significantly associated with a lower percentage of predicted values for FEV1 and FVC. Former smoking was significantly associated with a lower FEV1. University education, however, was associated with a higher FEV1.

The estimated reduction in FEV1 associated with living close to a road with >30 000 vehicles per day was somewhat greater than for those living close to a road with >10 000 vehicles per day, and the dose–response trend was slightly stronger (table 3).

Adding the percentage of lorry (heavy) traffic to the model did not improve the fit, and neither was it a significant predictor (results not shown).

Stratifying the analysis by gender, we found that associations between exposure and predicted FEV1 and FVC were stronger in women, although this analysis had less statistical power. For FVC, there was a significant negative association for the medium exposure category and a significant dose–response trend; for FEV1, the dose–response trend did not reach statistical significance (table 4).

Stratifying the data by respiratory health status, we found that high exposure in individuals with obstructive lung disease was associated with significantly lower FVC (−4.5%, 95% CI −8.8% to −0.1%). FEV1 was also lower, but did not reach statistical significance (−2.2% (95% CI −7.6% to 3.2%) (figure 1 and online supplementary table S1). In current smokers with high exposure, FEV1 was lower (−2.8%, 95% CI −6.2% to 0.6%), but the result was not significant (figure 1 and online supplementary table S2).

In the sensitivity analysis, including the percentage of traffic larger than cars (highest percentage was 8) into the model did not improve the fit, and nor was it a significant predictor of lung function.

**DISCUSSION**

In this study of adults from a single metropolitan area, we found an association between living close to densely trafficked roads and lung function, with a dose–response relationship for FEV1 and near-significant trend for FVC. The associations were stronger in women and among individuals with obstructive pulmonary disease. The estimated reduction in FEV1 for those living close to a road with >30 000 vehicles per day was somewhat greater than for those living close to a road with >10 000 vehicles per day, and the dose–response trend was slightly stronger (table 3).
trafficked roads and having a lower than predicted FEV₁ and FVC. The associations were stronger after adjusting for covariates, and they followed a dose–response trend, indicating that high exposure to traffic was more detrimental than medium exposure. The estimated effects of living close to a road with very dense traffic flows (more than 30,000 vehicles per day) were generally higher, but less consistent, indicating that a dose response could be present, but that the relatively few people exposed to high levels of very dense traffic compromised the statistical power. Though we cannot exclude bias due to unmeasured factors, or the role of chance, the observed association was most likely due to traffic exposure from large roads near the residence. There were more current smokers in the high and medium exposure areas, but there were also more people with a university education. The adjusted analyses had more significant associations and yielded higher estimates and a higher explanatory power ($R^2$ was <0.001 in the unadjusted models, and 0.064 in the adjusted models).

The effect sizes found in the current study are on par with those found in other studies. For example, Kan et al. found that women living closer than 150 m from a road with 10,000 vehicles had, on average, a 15.7 mL lower FEV₁ and 24.2 mL lower FVC than those living further away. Converting our results from percentage predicted to mL, women in our study with medium or high exposure had, on average, a 29.7 and 43.8 mL lower FEV₁, and a 35.1 and 63.7 mL lower FVC, respectively. Forbes et al. reported that a 10 µg/m³ increase in NO₂ in the area was associated with a 22 mL reduction in mean FEV₁ (0.7% of the population mean). In our study, 10 µg/m³ corresponds roughly to the difference between high and low exposure in our data, making the result similar to ours (table 2). In a cross-sectional study of the nationwide National Health and Nutrition Examination Survey (NHANES, data collected between 1971 and 1975), decreases in FEV₁ and FVC were found with increasing levels of modelled total suspended particulates at the participants’ residence. In a cross-sectional study of the Swiss Study on Air Pollution and Lung Diseases in Adults (SAPALDIA), community-level concentration of all pollution types was inversely associated with FVC, whereas a decrease in FEV₁ was only associated with traffic pollutants. In a follow-up of the same cohort, declining particulate matter (PM₁₀) exposure over 12 years was associated with less attenuated age-related deterioration in mean FEV₁ compared to having more constant exposure. Other studies have found indications of susceptible groups when looking at modelled levels of ozone, and exposure to high levels of PM₁₀ in relation to FEV₁, showing only significant associations among men with a family history of respiratory disease. The association between exposure and

### Table 2

Lung function change in those exposed to dense traffic (>10,000 vehicles per day) versus reference (>500 m)

| Outcome | Traffic exposure | n    | Unadjusted models | p for trend | Adjusted* models | p for trend |
|---------|------------------|------|-------------------|------------|------------------|------------|
| FEV₁    | >500 m           | 2441 | –                 | –          | –                | –          |
|         | 75–500 m         | 2688 | –0.66 (–1.41 to 0.09) | 0.065      | –0.80 (–1.53 to –0.07) | 0.027 |
|         | <75 m            | 367  | –0.93 (–2.44 to 0.57) | 0.101      | –2.47 to 0.45     | 0.027 |
| FVC     | >500 m           | 2441 | –                 | –          | –                | –          |
|         | 75–500 m         | 2688 | –0.37 (–1.05 to 0.32) | 0.167      | –0.57 (–1.23 to 0.10) | 0.058 |
|         | <75 m            | 367  | –0.80 (–2.16 to 0.58) | 0.224      | –2.24 to 0.42     | 0.058 |

*Adjusted for age, gender, body mass index, education and smoking status.

FEV₁, forced expiratory volume in 1 s; FVC, forced vital capacity.

### Table 3

Lung function change in those exposed to dense traffic (>30,000 vehicles per day) versus reference (>500 m)

| Outcome | Traffic exposure | n    | Unadjusted model | p for trend | Adjusted* model | p for trend |
|---------|------------------|------|------------------|------------|------------------|------------|
| FEV₁    | >500 m           | 4382 | –                 | –          | –                | –          |
|         | 75–500 m         | 1020 | –0.87 (–1.80 to 0.06) | 0.057      | –1.06 (–1.97 to –0.16) | 0.012 |
|         | <75 m            | 39   | –1.31 (–5.61 to 2.99) | 0.223      | –6.40 to 1.94    | 0.012 |
| FVC     | >500 m           | 4382 | –                 | –          | –                | –          |
|         | 75–500 m         | 1020 | –0.86 (–1.705 to –0.01) | 0.111      | –1.11 (–1.94 to –0.39) | 0.018 |
|         | <75 m            | 39   | 1.23 (–2.69 to 5.14) | 0.33       | –3.47 to 4.12    | 0.018 |

*Adjusted for age, gender, BMI, education and smoking status.

FEV₁, forced expiratory volume in 1 s; FVC, forced vital capacity.
decreases in FEV1 and FVC averages were stronger in women than in men. A possible explanation for this observation is that the exposure assessment based on residential address could be more accurate in women, as women tend to have lower labour market participation and are therefore more often at home. However, several studies suggest women to be more susceptible to adverse respiratory health effects (mortality, symptoms, lower FEV1 or FVC) from exposure to air pollution, or other exposures, although other studies found that men were more susceptible.

Other studies have shown associations between traffic proximity and respiratory health. Living near a large road was associated with increased risk of developing asthma and chronic obstructive pulmonary disease (COPD). However, the specific mechanisms through which traffic exposure affects lung function are not fully understood.

Table 4: Lung function change in women and men exposed to dense traffic (>10,000 vehicles per day) versus reference (>500 m)

| Outcome | Traffic exposure | Women | p for trend | n | Men | p for trend |
|---------|------------------|-------|-------------|---|-----|-------------|
| FEV1 | >500 m | 1257 | − | 1153 | − |
| 75–500 m | 1451 | −0.82 (−1.83 to 0.19) | 0.82 (−1.79 to −0.31) |
| <75 m | 198 | −1.21 (−3.28 to 0.89) | 165 | −0.85 (−2.98 to 1.28) | 0.169 |
| FVC | >500 m | 1257 | − | 1153 | − |
| 75–500 m | 1451 | −0.97 (−1.90 to −0.04) | 1217 | −0.09 (−1.03 to 0.86) | 0.169 |
| <75 m | 198 | −1.76 (−3.61 to 0.09) | 0.014 | 165 | 0.07 (−1.84 to 1.99) | 0.950 |

Estimates are adjusted for age, BMI, education and smoking status. FEV1, forced expiratory volume in 1 s; FVC, forced vital capacity.

Figure 1: Lung function outcomes and exposure to dense traffic (>10,000 vehicles per day) adjusted for age, gender, BMI, and education (also adjusted for smoking) stratified by respiratory health status and smoking status.
asthma in a prospective cohort study. In a multicentre study, living near dense traffic was associated with a lower FEV₁ and FVC in women, while in men only age-adjusted means of FEV₁ were lower to near significance. In a multicentre study, women living near dense traffic had a lower mean FEV₁ and FVC and increased risk of COPD. However, other studies found no associations between living near a road with dense traffic and FEV₁ or other measures of respiratory morbidity. In individuals with obstructive lung disease, high exposure to dense traffic was associated with airway restriction, with a significantly lower FVC than for the low-exposed individuals in the same group. Interestingly, the effect estimate for medium exposure was very near the reference (figure 1). However, these results barely reached statistical significance, possibly due to the low number of participants in the highest exposed groups. The lower than predicted FVC indicates a restrictive effect on pulmonary function, which is more associated with obesity, systemic and metabolic morbidities and mortality, which is generally increased in COPD. In our analyses, we found that age was a negative predictor of lung function, although using a predicted value should have accounted for this. Other studies have found air pollution effects in elderly participants and in the subgroup analysis we found that the association between age and lung function was only significant in those with obstructive lung disease. This finding supports that the observed association between age and age-adjusted lung function could be explained by age-related accumulation of comorbidities, which could also be associated with exposure to traffic pollution. A general deterioration in health could also increase susceptibility to exposure to air pollution, which could contribute to accelerated lung function decline. The composition of vehicles has been found to influence the impact of air pollution on respiratory health in children, and lorry traffic within 50 m has been shown to predict indoor air pollution, but estimates for this variable were near the null.

Our study has some limitations; we used distance to the nearest road to estimate traffic exposure at the current address, as modelled values of NO₂ and NOₓ were only available for 73% of the study population. However, distance to road also has advantages as an exposure indicator for the complex traffic-related air pollution mix, and there is good correlation (Pearson) between the modelled values and the absolute distance to the nearest road with (correlation coefficient −0.51 for NO₂ and −0.47 for NOₓ; p values > 0.001). The cut-offs at 75 and 500 m from roads were selected based on previous studies about distribution of pollutants near large roads, but the literature holds a large span of options for cut-offs. The crude measure of distance to road is significantly associated with lung function in the adjusted models (not shown), but the resulting coefficients are very small. We had no information about how long the participants had lived at their current address. Older people tend to have lived longer in the same place, and exposure assessment based on the residence may also improve as people age and leave the workforce, as people spend more time in their homes. However, misclassifications of exposure are most likely to bias the results towards the null. The most relevant traffic data available (from 2004) predate the index date of some participants by as much as 3 years. However, comparing 2004 traffic data with 1997 traffic data, only 2.3% of participants changed exposure category. From 2004 to 2010, 3.1% of participants changed exposure category, a precision we consider acceptable and which is unlikely to bias our results appreciably. Education was used as a measure of socioeconomic status, but unfortunately no information about income or occupation was available, which could have improved the measure. Education is usually a better marker of socioeconomic status in men than in women.

It is a strength of the study that data were collected at one centre, and the fact that the population came from a large cohort sampled from the same area increases the likelihood that the observed effects were due to the traffic exposure rather than regional confounding. However, the low participation rate is a concern. A non-participation analysis of the data gathered in 2001–2003 showed that women, the elderly and the university educated were more likely to participate. If non-participation was selective in such a way that those who live close to the main traffic areas and also suffer from a respiratory illness were less likely to participate, our effect estimates could be exaggerated. However, the social stratification of the cohort is expected and is particular to this area, where the high exposed study group had the highest proportion of highly educated people, opposite to many other urban areas. Most results adhere to a dose–response pattern showing increasing effects with increasing exposure which further indicates that differences were related by the exposure rather than bias from exposure to other unmeasured confounders.

In conclusion, in this study of a large cohort in a single metropolitan area, high residential exposure to dense traffic was associated with reductions in percentage of the predicted FEV₁ and FVC. In those with obstructive lung diseases, the association of FVC with high exposure was particularly strong. This is potentially an important finding, but it should be verified in other studies before being incorporated into official advice for this patient group.

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Contributors HKC was involved in statistical analyses, interpretation of the results and drafting of the article; FN and A-CO was involved in development of the research questions, designed and coordinated the study, interpretation of results and preparation of the manuscript; KT and AL was involved in development of the research questions, interpretation of results and preparation of the manuscript; LM was involved in development of the research questions, interpretation of results and preparation of the manuscript; all authors read and approved the final version of the manuscript.

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Data sharing statement Additional data from the ADONIX study exist and are held by the communicating author.

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